

Identifying Differences in U.S. Exposures to Ubiquitous Carcinogens

Adam Theising, Tina Bardot and Ann Wolverton

Working Paper 24-13
December, 2024

Identifying differences in U.S. exposure to ubiquitous carcinogens*

Adam Theising
U.S. EPA

Tina Bardot
ORISE Fellow
U.S. EPA

Ann Wolverton
U.S. EPA

December 2024

Abstract

Cancer is one of the leading causes of death in the United States, and substantial racial-, ethnicity-, sex-, and income-based differences in cancer incidence and mortality persist despite declining overall trends. Underlying differences in exposure to carcinogenic chemicals are often cited as a likely factor contributing to persistent differences in the incidence of certain cancer types. In this exploratory analysis, we construct a novel database of actual or potential exposure to ubiquitous carcinogenic chemicals based on nationally-representative biomonitoring and environmental data to produce demographically-differentiated exposure statistics, where possible. Despite major data gaps – one or more measures of direct or indirect exposure were available for only 37% of these 622 carcinogens – our results show evidence of potential exposure disparities for at least 28 carcinogens. We also review readily-available information on exposure pathways and cancer types associated with these carcinogens to identify common pathways through which households may be exposed, and explore correlations between relative exposure levels and cancer incidence rates. We end the paper with a discussion of key data gaps and limitations that future data gathering and investigation could address.

*Corresponding author: theising.adam@epa.gov. We thank Zach Stanfield for providing key data and support with the household products analysis and Wes Austin for support with the drinking water sampling data. We are grateful to Charles Bevington (CPSC), Suzanne Condon (ATSDR), Kristi Fedinick (OSTP), Erica Kimmerling (OSTP), Ruth Lunn (NIH), Suril Mehta (CEQ), and Ben Place (CEQ) for valuable early stage discussions and feedback on ways of identifying carcinogens and exposure data collection. We also thank Wes Austin (EPA), Heather Klemick (EPA), Danelle Lobdell (EPA), Greg Miller (EPA), Rachel Schaffer (EPA), Mitch Sumner (EPA), and several reviewers from various CDC centers (NCEH, DCPC, NCHS, and NIOSH) for very helpful comments on an earlier version of the manuscript. This work was completed with the support of the Oak Ridge Institute for Science and Technology (ORISE) fellowship program. The findings, conclusions, and views expressed here are those of the authors alone and do not necessarily reflect those of the United States Environmental Protection Agency or any of the federal agencies for the acknowledged individuals. No official agency endorsement has been granted, nor should such endorsement be inferred.

1 Introduction

Chemicals are commonly used and widely occurring in many products found in the home or used in the workplace (Helm et al., 2018; Dodson et al., 2012). Dust, furniture, and air samples indicate their presence in the indoor environment, and urine and blood samples confirm widespread exposure (Mitro et al., 2016; Schildroth et al., 2022). While exposure to some of these chemicals is nearly universal, researchers have documented substantial disparities in exposure to certain chemicals by race, ethnicity, income, and sex. For instance, studies have found that Mexican American and Non-Hispanic Black women are exposed to much higher levels of parabens and phthalates via personal care products, plastics, diet, and food packaging (Chan et al., 2021; James-Todd et al., 2016; Calafat et al., 2010). Others have found that low-income households are exposed to higher concentrations of volatile organic chemicals that off-gas from less-expensive furniture and building materials in older homes (Bi et al., 2018; Dodson et al., 2017). In addition, studies have observed that those highly exposed to one type of chemical are often highly exposed to other chemicals, including through outdoor and workplace exposure to environmental contaminants (Woodruff et al., 2011; Bertin et al., 2018; Kapraun et al., 2017). Exposure to a subset of these chemicals, individually or as part of mixtures, has been linked to adverse health effects such as higher incidence of pre-term births, gestational diabetes and other metabolic disorders, hypertension, and in some cases cancer, among specific populations.

The evidence cited above is based on the chemical content of specific products, location-specific surveys and sample collections, and national-level data for specific chemicals. In this paper, we contribute to the growing literature by combining multiple national-level datasets on actual or potential exposure to carcinogenic chemicals and calculating demographically-differentiated exposure statistics for individual chemicals where such information is available. We find the strongest evidence of potential exposure disparities for 28 carcinogens; for the chemicals arsenic, 1-naphthylamine, acrylonitrile, furan, 1,4-dichlorobenzene, 2,4-dichlorophenoxyacetic acid (2,4D), and ethylene oxide, at least three unique statistical measures indicate potential exposure disparities. We also explore broader demographic trends in carcinogen exposure. Though we observe that all of the demographic subgroups we examine likely experienced relatively high exposure to specific carcinogens, several measures indicate that non-Hispanic Black Americans – especially lower-income and male individuals – had the highest cumulative carcinogen exposure across the chemicals for which data are available. We emphasize that while these findings may identify potential disparities in exposure, they do not immediately imply national disparities in cancer hazard or risk.

As a complement to the exposure analysis, we identify what is known regarding the main pathways through which exposure to these 28 carcinogens may occur. We also discuss key data gaps where additional data gathering and future study could be useful. While it is beyond the scope of this paper to identify the contribution of individual carcinogenic chemicals to overall cancer risk, we also examine whether trends in national level incidence and mortality for specific cancer types associated with these chemicals seem consistent with demographically-differentiated exposure.

We focus our analysis on observed differences in individual or household exposure to commonly used and widely occurring (i.e., ubiquitous) carcinogenic chemicals. Cancer is the second leading cause of death in the United States, after heart disease. The National Cancer Institute estimates about 2 million new cancer diagnoses and more than 600,000 cancer deaths will occur in the United States in 2024 (NCI, 2024). Additionally, the financial burden of cancer care is substantial: annual medical care costs, alone, are expected to exceed \$246 billion by 2030 in the United States (Mariotto et al., 2020). External factors such as lifestyle choice and exposure to toxic chemicals are major contributors to lifetime cancer risk (Wu et al., 2016). In

addition, while overall cancer rates have declined since the 1990s due to reduced use of tobacco products, early detection, and better treatment methods, Black individuals lag in early-stage diagnoses of cancer; Black women and people of color continue to have higher than average rates of cancer incidence; and individuals with lower socioeconomic status have higher mortality rates after being diagnosed with cancer (O’Keefe et al., 2015; Islami et al., 2024; Shaw et al., 2024).

The paper is organized as follows. Section 2 explains the methodology to identify ubiquitous carcinogens, gather exposure data from various sources, and classify carcinogens based on notable differences in exposure across demographic subgroups. Section 3 highlights our main findings: nationally-representative statistical measures of differential exposure to carcinogens by demographic (race, ethnicity, and sex) and economic categories (income level). We also summarize our findings on common exposure pathways and describe potential correlations between differences in exposure and disparities in cancer incidence by cancer type. In Section 4, we discuss the limitations of the analysis and note key data gaps. Section 5 concludes.

2 Methods

2.1 Scope of analysis

For our analysis, we combined multiple data sources that shed light on carcinogenic chemicals that are widely occurring in the environment, at home, and in the workplace. To set the scope of our analysis, we first determined the universe of carcinogenic chemicals eligible for our analysis, established criteria for classifying ubiquity, and then selected the main data sources for use in identifying national-level demographic differences in exposure.

2.1.1 Scoping chemicals on carcinogenicity

Several international, federal, and state agencies classify and maintain lists of carcinogens, and while there is substantial overlap, differences exist. In an attempt to be inclusive, we compiled a super-list of the carcinogens appearing on one or more of several lists, without favoring one institution over another, and we considered known, probable and possible carcinogens. Specifically, we included carcinogenic chemicals on the International Agency for Research on Cancer’s (IARC) Group 1 (known carcinogens), IARC Group 2A (probable carcinogens), and IARC Group 2B (possible carcinogens) lists; known and reasonably anticipated human carcinogens from the National Toxicology Program’s Report on Carcinogens (RoC); cited carcinogens from both the European Union and United Kingdom Candidate Lists for Substances of Very High Concern (SVHC); and cited carcinogens from California’s Proposition 65 list.

After compiling the carcinogenic chemicals that appear on these source lists, we removed duplicates, and then matched them to the U.S. Environmental Protection Agency’s (EPA) CompTox Chemicals Dashboard based on the Chemical Abstracts Service (CAS) registry number.¹ We then standardized all chemical names to the preferred name listed in the dashboard (Williams et al., 2017). Carcinogens without a CAS number or CompTox identification number (DTXSID) and those directly related to radiation, viruses, alcohol, and

¹A CAS number is a unique identification number assigned to every chemical substance described in the open scientific literature since 1957. The CompTox Chemicals Dashboard is a free, searchable online repository of information on the chemistry, toxicity, and exposure of thousands of chemicals. It is maintained and updated regularly by the U.S. EPA. See <https://comptox.epa.gov/dashboard/>.

tobacco were considered out of scope for this exercise and dropped from the sample.² In total, we identified 727 chemicals as within scope for carcinogenicity.

2.1.2 Scoping chemicals on ubiquity

We also limited the scope of our analysis to chemicals that can be considered ubiquitous in the human environment based on currently available data sources. Chemical ubiquity as a scientific construct remains ill-defined and ambiguous, so we explored several readily-available metrics to inform our scoping exercise. For each of the 727 carcinogens identified, we assessed whether the chemical was (1) active on EPA’s Toxic Substances Control Act (TSCA) Inventory (EPA, 2024b); (2) produced in or imported to the US per the EPA’s 2020 Chemical Data Reporting database (EPA, 2020); or (3) reported in the harmonized Multimedia Monitoring Database (Isaacs et al., 2022). We also assessed the chemical’s (4) numeric count of related patents per PubChem (Kim et al., 2018) and (5) numeric count of scientific articles in which a chemical was mentioned per PubChem (Kim et al., 2016).³

We eliminated any chemical from our sample that satisfied none of (1)-(3) **and** also fell at or below the 25th percentile for either criteria (4) or (5). Jointly, these scoping requirements restricted our focus to carcinogens that are actively used or produced at volume, specialty chemicals with substantial patenting and research activity, or legacy carcinogens which are/were environmentally monitored by governmental agencies.⁴

In total, 105 chemicals did not meet these criteria, leaving 622 ubiquitous carcinogens as candidates for our analysis. 223 chemicals (36%) were on only one carcinogen source list, 230 chemicals (33%) were on two source lists, 173 chemicals (27%) were on three source lists, and 28 (4%) were on all four source lists. See panel A of Appendix Table A1 for additional details. This variance in carcinogen classification across authoritative source lists lends some *prima facie* support for our inclusive scoping approach.

2.1.3 Scope of exposure data considered

We focused our analysis on nationally-representative data. While additional insights would likely result from reviewing alternative data sources including the academic literature, location-specific study samples, state or local governments, or community-generated activities, the primary purpose of this analysis is to screen for notable national-level differences in chemical exposure using readily-available public information.⁵ We leave a systematic review of alternative data sources for future work.

2.2 Datasets and analytical approach

We conducted four separate descriptive statistical exercises to assess the extent of differences in exposure across demographic subgroups to the chemicals on our candidate list of 622 carcinogens.

Our first analysis used direct measures of exposure – biomonitoring data – while the other three analyses relied on indirect environmental measures of possible exposure - in the ambient air, in drinking water, and

²While some chemicals without CAS Numbers are recognized as possible or probable carcinogens they were dropped from consideration for this analysis because they are more difficult to link to exposure data. A full list of candidate carcinogens and their scoping attributes are available in supplemental Appendix Table E1

³PubChem is an open chemical database maintained by the U.S. National Institutes of Health.

⁴While legacy carcinogens may not be ubiquitous in terms of active production, exposure remains an ongoing problem. See, for example, Barr et al. (2022), Sun et al. (2016), McGoldrick and Murphy (2016).

⁵While scoping the national availability of biomonitoring data, we contacted several state biomonitoring programs to see if they could supplement the chemical coverage available through NHANES. Ultimately, additional chemical coverage was limited, so we retained a focus on national biomonitoring data. A summary of findings from this outreach is available in Appendix D.

via purchase and use of consumer products. This section describes the data sources and analytical methods used. All data processing and analysis were conducted using scripts written in the R language (version 4.4.0), which are available in an online appendix.

2.2.1 Analysis of NHANES biomonitoring data

The natural starting point for assessing chemical exposure is biomonitoring data. These data provide a direct measure of the amount of a chemical found in the human body via all possible routes of exposure, including inhalation, ingestion, and dermal absorption. There are two types of biomarker samples generally available: blood/serum samples and urine samples. Blood/serum samples are most useful for understanding exposure to chemicals with longer half-lives (residing in the body for months to years) or steady-state ubiquity but can be challenging to collect. Some examples of suitable chemicals include polychlorinated biphenyls (PCBs), per- and polyfluoroalkyl substances (PFAS), and metals. Urine samples are more useful for understanding exposure to chemicals with shorter half-lives (residing in the body for hours to days) such as phenols and tobacco but results can be highly variable across time (LaKind et al., 2014; Aylward et al., 2014). While biomarker concentrations are de facto evidence of exposure intensity, several factors may complicate the relationship between true exposure levels and biomarker concentrations (Tan et al., 2012). Given the lack of alternative national-level data sources, we nonetheless relied on biomarker concentrations as a proxy of carcinogen exposure.

We used chemical-specific biomonitoring data collected via blood, serum, and urine samples from the U.S. population as part of the National Health and Nutrition and Examination Survey (NHANES) administered by the Centers for Disease Control and Prevention’s (CDC) National Center for Health Statistics. This survey has been conducted periodically, including the 1988-1994 survey, and continuously since 1999 with data acquired and released at two-year intervals (i.e. survey cycles). Though the survey cycle is on a two year schedule, the analytes evaluated for a given cycle have varied over time.

While the CDC produces disaggregated national biomonitoring summary statistics for a select set of demographic subgroups (CDC, 2024), we used the underlying NHANES microdata, which allowed us to create national exposure summary statistics further stratified by intersections of age, sex, household income, and race/ethnicity (CDC, 2020). This analysis primarily relied on a merged and harmonized version of the continuous NHANES data; we analyzed examination-level biomonitoring samples drawn from individuals between 2003-2004 and 2017-2018 (Nguyen et al., 2023). For a subset of biomonitored chemicals – namely, organochloride pesticides, dioxins/furans, PCBs, and brominated flame retardants – the most recent samples available are based on pooled measurement. In those cases, we directly acquired and relied on the original, unharmonized NHANES microdata from the 2011-2012 and 2015-2016 cycles (CDC, 2020).

We examined blood, serum, and urine biomarkers only for chemicals that were on the ubiquitous carcinogen list. We followed best practices in the use of NHANES: examination or subsample survey weights were used to account for NHANES’ sampling design (Nguyen et al., 2023; CDC, 2020), efforts were taken to ensure metabolites of carcinogens were correctly assigned to the parent chemical (Stanfield et al., 2024), and the analysis followed the CDC’s convention of not reporting geometric mean estimates for stratification subgroups with >40% of test results below the reported limit of detection (CDC, 2024).⁶ We also did not report geometric mean estimates for a carcinogen if >40% of all national test results were non-detects. For carcinogens included in multiple NHANES cycles, statistics were based on only the most recent cycle of

⁶We imputed a value of $\frac{LOD}{\sqrt{2}}$ for samples below the detection limit when reporting geometric mean values. This is a common approximation method of censored values below the detection threshold for screening work of this nature.

data.⁷ In total, 106 of the 622 candidate carcinogens (17%) were surveilled in NHANES by blood, serum or urine – directly or via a metabolite – at least once since 2003. 74 carcinogens were only monitored directly, 21 carcinogens were only monitored via a metabolite, and 11 carcinogens were monitored both directly and by a metabolite.⁸

For each carcinogen and/or metabolite with individual-level biomonitoring data, we calculated the national geometric mean and 95th percentile concentrations for several demographic subgroups of interest. Our primary results focused on exposure differences by sex, race/ethnicity, and household income, and relied on samples from all age groups. In supplemental results, we also assessed statistical decompositions by age, interactions of sex and race/ethnicity, and interactions of household income and race/ethnicity.⁹ Of the 106 surveilled carcinogens, only 39 (6% of candidate carcinogens) had a sufficient count of samples above the limit of detection for complete geometric mean estimates stratified by race/ethnicity, sex, and household income. Therefore, our primary analysis also considered subgroups’ 95th percentile concentrations, for which a broader set of 62 carcinogens (10% of candidates) can be analyzed. See Appendix Table A2 for a list of carcinogens matched to NHANES biomonitoring data and further information on the analyzable subsets.

Based on these statistics, we extended the methods of Belova et al. (2013) and Nguyen et al. (2020) and screened for exposure differences across subgroups using both geometric mean ratios and 95th percentile ratios as metrics:

$$GMR_{bcs} = \frac{GM_{bcs}}{GM_{bcr}} \quad (1)$$

$$PR_{95,bcs} = \frac{P_{95,bcs}}{P_{95,bcr}} \quad (2)$$

We defined the geometric mean ratio (*GMR*) as the ratio between the geometric mean of biomarker *b* for chemical *c* in subgroup *s* and the geometric mean of that chemical biomarker in a reference group *r*. Analogously, the 95th percentile ratio (*PMR*₉₅) is defined as the ratio between the 95th percentile of biomarker *b* for chemical *c* in subgroup *s* and the 95th percentile of that chemical biomarker in a reference group *r*. For ease of comparability across all of the different subgroups considered, we selected the entire national population as the reference group.

Turning to the 10 matched carcinogens with pooled biomonitoring data, we elected to calculate arithmetic mean concentrations by demographic subgroup. Calculation of geometric mean or percentile statistics for these carcinogens would be assumption-laden and computationally challenging due to the pooled nature of the data and the survey’s complex sampling design. Instead, we followed the CDC’s standard practice in using these pooled data and assessed arithmetic means differentiated by race/ethnicity, sex, and age (CDC, 2021a). Only 5 of the 10 carcinogens (1% of all candidate carcinogens) had a sufficient count of samples above the limit of detection for a complete set of arithmetic mean estimates stratified by race/ethnicity and sex. We used these estimates from the pooled data to calculate arithmetic mean ratios (*AMRs*) analogous to the geometric mean ratio described above, again using the entire national population as the reference group.

⁷The most recent cycle varied by chemical: the earliest cycle considered in this analysis was from 2003-2004 and the latest was from 2017-2018.

⁸These 106 chemicals are a broad mix of known, probable, and possible carcinogens; see panel B of Table A1 for a summary. NHANES selects chemicals based on evidence of exposure, known/potential health impacts, availability of analytical methods, and cost.

⁹Decompositions by sex are male or female; by household income, below two times the poverty line or above two times the poverty line; by race/ethnicity, non-Hispanic White, non-Hispanic Black, Hispanic, and non-Hispanic Other; by age, younger or older than 20 years. See Appendix C for additional details of the NHANES data construction.

2.2.2 Analysis of RSEI air data

In the absence of more complete biomonitoring coverage, we analyzed three additional data sources that provide indirect measures of exposure to specific chemicals via environmental contaminant pathways. We first examined potential exposure to individual carcinogens emitted into the air by industrial facilities. Research has shown that households residing in proximity to airborne industrial releases face increased risks of adverse health effects, including cancer, and that these neighborhoods are often home to lower-income, racial/ethnic minority, or non-native-English-speaking populations (Brender et al., 2011; Madrigal et al., 2024). We conducted an analysis of EPA’s Risk-Screening Environmental Indicators (RSEI) geographic microdata using a spatially aggregated version of these data that provide average airborne concentrations of each chemical reported to the Toxics Release Inventory (TRI) at the census block group (CBG) resolution (EPA, 2022).¹⁰

After filtering the 2022 RSEI air data to only include chemicals on the ubiquitous carcinogen list, we matched CBG-level air concentrations of individual carcinogens to CBG-level sociodemographic information from the Census Bureau’s latest available five-year (2018-2022) American Community Survey (Census, 2022). In total, 186 carcinogens from the candidate list are represented in the data. For each of these carcinogens, we separately calculated population-weighted national average airborne concentrations ($Conc_{cs}^*$) for subgroups of different races/ethnicities and by household poverty status:

$$Conc_{cs}^* = \sum_I \frac{Conc_{ic} Pop_{is}}{Pop_s} \quad (3)$$

where I indexes the national set of census block groups, $Conc_{ic}$ is the average annual concentration of chemical c in block group i , Pop_{is} is the population count of demographic subgroup s in block group i , and Pop_s is the national population count of subgroup s . To ensure mutual exclusivity, we considered the following self-identified race and ethnicity subgroups: non-Hispanic Asian, non-Hispanic American Indian, non-Hispanic Black, non-Hispanic Pacific Islander, non-Hispanic White, and Hispanic. We also considered income-based subgroupings, differentiating households earning less than twice the poverty line and households earning more than twice the poverty line. Using each subgroup’s nationally-weighted average concentration, we calculated carcinogen-specific relative ambient air concentration ratios as in Equation 1, again with the national average concentration for the carcinogen as the reference value. These ratios are nationally representative, as all census block groups were included.

2.2.3 Analysis of EPA drinking water samples

We next examined potential exposure to individual carcinogens present in public drinking water. Previous national-scale research has shown demographic subgroups face inequalities in exposure to individual chemicals present in public drinking water (Hefferon et al., 2024; Martinez-Morata et al., 2022; Nigra et al., 2020; Ravalli et al., 2022). We collected and harmonized the universe of EPA’s drinking water sampling data from community water systems (CWS) for chemicals sampled via the Six Year Review process (SYR) and Unregulated Contaminant Monitoring Rules (UCMR). Our analysis only included sampling results for chemicals on the ubiquitous carcinogen list; this exercise resulted in millions of drinking water samples for 70 matched carcinogens, with temporal coverage spanning 2006 to 2019 from SYR3 through SYR4 (EPA, 2016b;

¹⁰These data input self-reported facility-level air emissions from the TRI into a medium-complexity fate-and-transport model over a spatial 810m by 810m national grid to develop estimates of the average airborne concentration of a specific TRI-reported chemical in a given grid cell. These grid cell averages are then spatially aggregated up to the census block group geography.

EPA, 2022b) and 2008 to 2020 from UCMR2 through UCMR4 (EPA, 2024a).

Every sample in the database is attributed to a single CWS. For each CWS, we calculated an average carcinogen concentration based on all of the system’s collected samples from 2006-2020. Of the 70 matched carcinogens, only 15 had a sufficient count of samples with concentrations above the limit of detection for calculation of meaningful CWS-level averages.¹¹ We used information from Austin et al. (2024) on the demographic characteristics of residents within each CWSs’ service area boundaries – namely, the total population count, the fraction of the population identifying as a specific race or ethnicity, and the fraction of the population earning income above and below 2x the poverty line, all as of 2022. With this information, we calculated population-weighted average carcinogen concentrations in drinking water (DW_{cs}^*) for each demographic subgroup:

$$DW_{cs}^* = \sum_J \frac{DW_{jc} Pop_{js}}{Pop_s} \quad (4)$$

where J indexes the complete set of CWSs included in our sampling database, DW_{jc} is the average concentration level of carcinogen c in the drinking water of CWS j , Pop_{js} is the population count of demographic subgroup s in CWS j , and Pop_s is the national population count of subgroup s across all CWSs in our sampling database. With these average drinking water concentrations, we calculated carcinogen-specific relative concentration ratios for each racial/ethnic or household income subgroup as in Equation 1, again with the national average drinking water concentration for the carcinogen serving as the reference value. These relative concentration ratios serve as a proxy of differences in exposure via drinking water.

2.2.4 Re-analysis of Stanfield et al. household product data

Our third and final analysis of potential differences in indirect exposure examined carcinogens present in household products. Stanfield et al. (2021) linked a year’s worth of nationally-representative consumer product purchasing data from Nielsen with information on chemicals present in specific products from EPA’s Chemical and Products Database (Dionisio et al., 2018). With this information, Stanfield et al. (2021) assessed how possible chemical co-exposures varied across demographics due to real-world purchasing patterns.

We accessed relevant portions of the 2021 paper’s data, and re-purposed them to analyze the extent of differences in exposure to carcinogens via consumer products. The underlying Nielsen product purchasing data are monthly observations from 60,000 participating households for the year 2012. While there have certainly been changes to consumer markets’ product and chemical mix over the ensuing decade, but we are unaware of any other readily available data of this type that has been linked to product chemical information. We found a total of 44 carcinogens from the candidate list reflected in these household purchasing and products data. See Stanfield et al. (2021) for details of the monthly household-chemical panel data construction.

We measured the *national prevalence* of potential exposure to a given carcinogen via household products as the fraction of total household-months that the chemical appears in the purchased bundle:¹²

$$Prev_c^* = \frac{\sum_H \sum_T \mathbb{I}_{cht}}{HT} \quad (5)$$

¹¹See Appendix C for a description of data processing, including imputation methods for non-detect samples and a list of matched carcinogens. These data were generally representative of the US population that receives public drinking water from a CWS with SYR and UCMR samples. For the 15 carcinogens studied, 12 statistics represented >200 million Americans, and 1 statistic each represented >140 million, >32 million, and 90,000 Americans, respectively.

¹²For example, in a 60,000 household panel over 12 months, there were 720,000 observations. If a carcinogen showed up in the purchased good bundle for 360,000 household-months, our national prevalence measure would be 0.5.

where H indexes the complete set of households in the Nielsen data, t indexes the monthly time component of the panel data (with $T = 12$), and $\mathbb{I}_{cht}$ denotes an indicator that the household-month goods bundle contained carcinogen c . We calculated analogous disaggregated chemical purchase prevalence statistics for households with lower- or higher-income levels, members of a racial/ethnic subgroup (Hispanic, Asian, Black, White), or which include a female of childbearing age (<45 years old):

$$Prev_{cs}^* = \frac{\sum_{H_s} \sum_T \mathbb{I}_{scht}}{H_s T} \quad (6)$$

where H_s indexes the set of households that were members of subgroup s . Again following from Equation 1, we defined our measure of potential differences in exposure to a carcinogen via use of household products as the ratio of a given subgroup’s purchase prevalence and the overall national purchase prevalence.

2.3 Synthesizing analyses to identify potential exposure disparities

With the set of differentiated exposure measurements described above, we identified at least one source of exposure for analysis for 230 (37%) of the 622 identified ubiquitous carcinogens. The final task was to synthesize statistics that identify the subset of these carcinogens with the strongest evidence of potential exposure disparities. To do this, we operationalized a set of selection criteria that balance the body of evidence available with tractability, consistency and a predisposition to inclusiveness in marginal cases.

Our primary screening for potential exposure disparities identified any carcinogen that met at least one of the following criteria based on the analyses described above:

1. NHANES geometric or arithmetic mean ratio greater than 1.5 for any primary demographic subgroup¹³;
2. NHANES 95th percentile ratio greater than 1.5 for any primary demographic subgroup;
3. At least two of the measures of indirect exposure to chemicals (i.e. via ambient air, drinking water, or consumer product purchases) scored greater than 1.5 for any subgroup.

A uniform inclusion threshold value across the different criteria ensures comparability across exposure metrics and data sources. Direct evidence of differences in exposure (via biomonitoring metrics) was implicitly given heavier weight than indirect evidence. By using a value of 1.5 as the primary inclusion threshold, this approach highlighted the set of chemicals where potential disparities are most evident; a subgroup with a ratio value of 1.5 faces 50% more exposure than the nationally-representative level. In supplemental results, we also tested (i) an alternative threshold value of 1.25 and (ii) alternative NHANES-based inclusion criteria, which considered results from demographic interactions of race/ethnicity, household income, sex, and age. Throughout the results section, we note any instances where findings meaningfully differed under these alternative criteria.

2.4 Assessing potential exposure pathways

For all carcinogens identified based on these primary criteria, we also reviewed readily available evidence from several sources to inventory their major exposure pathways. Epidemiologists and the broader risk assessment community have compiled an immense body of research studying exposure to potential and known hazardous

¹³The primary demographic subgroups considered for the NHANES analyses include {Hispanic, non-Hispanic Black, non-Hispanic White, non-Hispanic Other, household income below 2x poverty line, household income above 2x poverty line, female sex, and male sex}.

agents. Generally speaking, exposure assessment focuses on measuring the duration, frequency, and source of exposures, and is ultimately integrated with a separate hazard and dose-response assessment to characterize risks to groups and the general population (EPA, 2019). In regulatory or policy work, exposure assessment is often highly targeted, looking at potential exposure risk for specific population subgroups (e.g., agriculture workers, industrial workers, children, consumers) or via certain products and product categories that are known to contain hazardous agents. As such, the format of exposure assessments – and exposure research more broadly – is varied, their publication locations are diffuse, and unfortunately, there is not a “one-stop-shop” for researchers aiming to compile the entirety of the available literature and secondary data relating exposure pathways to a given chemical agent.

We therefore took a tractable but ultimately limited approach to assessing potential exposure pathways for the carcinogens with notable differences in exposure. We did this by reviewing readily available evidence from several authoritative sources and categorizing the major exposure pathways that could be contributing to exposure disparities. These sources include CompTox, PubChem, Haz-Map, and NTP’s 15th Report on Carcinogens (Williams et al. (2017); Kim et al. (2023); Brown (2024); NTP, 2021)¹⁴. Using all available information from these sources, we compiled common exposure pathways and associated product categories for each chemical. We additionally derived summaries of cancers associated with each carcinogen, separately noting when the association was based on animal or epidemiological (human) evidence.

We reviewed and presented exposure and use information in two separate but related ways. To reflect common ways of considering sources of vulnerability to chemicals, we explored **product categories** and **routes of exposure**. The former focused on classifications of products used in the home or workplace, reflecting frequently-used objects and materials that might contain carcinogens that pose a hazard. The latter focused on the activities and environments in which a chemical is used or otherwise present. We separately explored routes for general population and occupational exposures.¹⁵ These characterizations of exposure overlapped, but each provided an angle for considering the “where” and “how” of chemical exposure.

To classify product categories for each carcinogen, we began by adapting CompTox’s Chemical and Products Database (CPDat) categories, which were constructed using the methods described in Isaacs et al. (2022) for use in risk assessment. We focused on CPDat’s highest-aggregation-level of product use categories in households and in industry, and expanded the set to include other major product types found across the remaining three search sources. Most carcinogens for which we found exposure disparities are used in multiple products types. For example, as a preview of our findings, in Figure A5 we observed that nearly all of our identified carcinogens with likely exposure differences are in multiple product categories, and more than half of them are associated with 5 or more.

We then delineated exposure pathways by their capacity for general population and occupational exposure, noting the activities and places in which exposure to carcinogens is possible (e.g. on furniture in the home, through art or office-related materials, in agricultural settings). We stopped short of exploring specific mechanisms or estimating the likelihood of exposure via a particular pathway, lacking enough data to consistently and confidently suggest the intensity of use or extent of influence a given classification has on vulnerability.

Finally, using the same set of databases described above plus the IARC Monographs (IARC, 2024),

¹⁴For more information, hyperlinks here: CompTox, PubChem, ROC15, Haz-Map.

¹⁵We included both general population and occupational exposure to differentiate chemicals that are strictly industrial from those that may pose risk through everyday activities. This was in part to identify chemicals which have a “double incidence”, where workers may be exposed at home and at work to the same chemical, and in part to develop a better understanding of the types of chemicals to which different subpopulations are at greater risk based on their activities and occupation

we identified the forms of cancer and exposure types (e.g. inhalation, dermal, etc.) associated with each carcinogen. Because there is variability in the completeness and update frequency of each source database, we were inclusive of all listed exposure types and forms of cancer. For carcinogens with more thorough exposure and toxicological information available in the source databases – especially those with noted associations to particular cancer types in RoC and IARC reports – we emphasized the most recent available information on carcinogenicity and exposure/use. In many cases, however, the available information is limited or decades old.

3 Results

We begin by summarizing results on potential exposure disparities. Section 3.1 provides an overview of our findings based on the biomonitoring data, while section 3.2 summarizes our findings based on each of the indirect exposure data sources. For each individual analysis, we highlight (1) which carcinogens had the strongest evidence of potential exposure disparities, and (2) instances where media-specific evidence suggested differences in cumulative exposure to multiple carcinogens may exist. Next, in section 3.3 we discuss our cross-media synthesis of the exposure results and highlight carcinogens for which evidence was strongest. In section 3.4, we leverage additional information to better understand the main exposure pathways for each of the identified carcinogens in sections 3.1 and 3.2. Section 3.5 describes the extent to which cancer types associated with these carcinogenic chemicals also demonstrated disparities in national cancer incidence by race, ethnicity, and sex.

As a final point before presenting results, we emphasize that the following simply calls attention to **exhibited differences in exposure**; the results that follow **do not** immediately imply or explain national disparities in cancer hazard or risk. We assessed relative differences in exposure levels – not absolute exposure levels – and therefore cannot speak to potential risk. More formal risk assessment work is outside the scope of this analysis, which can best be characterized as an initial evaluation indicating areas where in-depth investigation could be beneficial.

3.1 Direct measures of exposure: NHANES biomonitoring data

3.1.1 Geometric mean ratios

We start by discussing the geometric mean (GM) ratio results. We had a sufficient fraction of detected biomarker samples to calculate GM ratios for 42 carcinogen metabolites and parent chemicals (representing 39 carcinogens in total). Of those, 13 total carcinogens – measured as 5 parent chemicals and 7 metabolites – returned a GM ratio of 1.5 or greater for at least one primary demographic subgroup. Figure 1 provides an overview of these results.

Panel (a) presents the distributions of GM ratios across all of the detected chemicals. These distributions are shown separately for each primary demographic subgroup. A few findings are visually striking. We found that the distribution of GM ratios for lower income individuals was skewed slightly higher than the distribution for higher income individuals. Similarly, the distribution of GM ratios for males was centered at a higher median value than it was for females. And while the GM distributions for Hispanic, non-Hispanic White, and non-Hispanic Other subgroups were all centered at median values close to 1, the GM ratio distribution for the non-Hispanic Black subgroup had a median value of greater than 1.25 and a considerable right tail. This last result evidences higher than national average exposure to a substantial number of

detectable chemicals for the average non-Hispanic Black American.

Panel (b) lists the subset of individual chemicals for which at least one primary demographic subgroup had a GM ratio of 1.5 or greater. We highlight some of the findings here. Lower income, Hispanic, and non-Hispanic Black subgroups showed high exposure to 1,4-dichlorobenzene (as measured by its metabolite 2,5-dichlorophenol) relative to the national average. Of particular note for 1,4-dichlorobenzene – a common ingredient in mothballs, deodorant cakes, and insecticides – the GM ratio for non-Hispanic Black individuals was 6.15, congruent with exposure levels more than 600% above the national average. GM ratios for the legacy pesticide nitrofen (as measured by its metabolite 2,4-dichlorophenol) showed potentially elevated exposure levels for Hispanic and Black individuals.¹⁶ The tobacco-specific carcinogen NNK (as measured by its metabolite NNAL) was found to have elevated GM ratios for lower-income and non-Hispanic Black subgroups.¹⁷ And finally, to re-emphasize a key finding, non-Hispanic Black individuals showed elevated exposure to 11 of the 13 carcinogens for which we observed *any* GM ratio of greater than 1.5, including ubiquitous carcinogens like acrylonitrile, naphthalene, acrolein, and carbaryl (i.e. 1-Naphthalenol, 1-(N-methylcarbamate)). The complete set of GM ratio results for primary demographic subgroups – including chemicals without any detected GM ratio above 1.5 – is provided in Table A4.

Demographic interaction results

In supplemental results, we also assessed GM ratios by demographic interactions of sex and race/ethnicity and demographic interactions of household income and race/ethnicity. These additional GM ratio results are available in Table A5, with subgroups’ distributions visualized in panel (a) of Figure A2. We briefly summarize key takeaways here. The elevated incidence of GM ratios greater than 1.5 for the non-Hispanic Black subgroup was particularly pronounced among males and lower-income individuals. In addition to the chemicals noted in the primary analysis, Black males and lower-income Black individuals presented with GM ratios greater than 1.5 for several more chemicals of note, including ethylene oxide (ratio = 1.79), ethylbenzene (ratio = 1.53), crotonaldehyde (ratio = 1.62), acrylamide (ratio = 1.76), and 4-biphenylamine (ratio = 1.56). Most of these are components of tobacco smoke, but the first two are also commonly used in industrial and manufacturing settings. GM ratios for non-Hispanic males of an “Other” race implied relatively high exposure to arsenic (ratio = 1.87) and melamine (ratio = 2.13).

Age-differentiated results

In supplemental results, we also assessed GM ratios by age group. These additional GM ratio results are available in Table A6, with subgroups’ distributions visualized in panel (b) of Figure A2. We briefly summarize key takeaways here, emphasizing exposure disparities that affect children. When considering GM ratios strictly differentiated by age group (age 19 and younger or age 20+), we did not identify any values greater than 1.5. Nevertheless, several carcinogens presented markedly higher GM ratios for the 19 and younger age group than the 20+ age group; these included di(2-ethylhexyl) phthalate (DEHP, ratio = 1.43), tris(2-chloroethyl) phosphate (TCEP, ratio = 1.44), and 2,4-dichlorophenoxyacetic acid (2,4D, ratio = 1.28). Joint consideration of age and race/ethnicity while assessing GM ratios turned up several interesting

¹⁶Interestingly, 2,4-dichlorophenol is also a metabolite of the still widely-used pesticide 2,4-dichlorophenoxyacetic acid (known more commonly as 2,4D); the elevated GM ratios for the metabolite 2,4-dichlorophenol contrasted with direct biomonitoring results for 2,4D that showed no evidence of exposure disparities across subgroups.

¹⁷The formal chemical name for NNK is 4-(N-Nitrosomethylamino)-1-(3-pyridyl)-1-butanone. The formal chemical name for NNAL is 4-(methylnitrosamino)-1-(3-pyridyl)-1-butanol.

results. The GM ratio for TCEP was highest among non-Hispanic White individuals aged 19 and younger (ratio = 1.54), and melamine exposure was highest on average for non-Hispanic “Other” individuals aged 19 and younger (ratio = 2.32). Many of the elevated relative exposure levels identified above for the broader non-Hispanic Black subgroup were driven by higher values for those aged 20 and greater, but there were exceptions. GM ratios for non-Hispanic Black individuals aged 19 and younger were greater than 1.5 for di(2-ethylhexyl) phthalate (ratio = 1.79) and malathion (ratio = 1.79). The former chemical is a widely-used plasticizer, while the latter is a common insecticide. Moreover, the sizable difference in exposure we identified for 1,4-dichlorobenzene appeared strongest among non-Hispanic Black individuals aged 19 and younger (ratio = 7.31, versus ratio = 5.81 for Black individuals aged 20+).

3.1.2 Arithmetic mean ratios

We next turn to a discussion of the arithmetic mean (AM) ratio results for the set of chemicals biomonitoring *with pooled samples*. There were a sufficient fraction of detected biomarker samples to calculate AM ratios for 21 carcinogens in total; 17 of these are individual PCB compounds, 2 are legacy organochlorine pesticides, 1 is a polybrominated biphenyl (PBB) compound and 1 is a chlorinated dibenzofuran.¹⁸ Of those, 9 carcinogens returned a GM ratio of 1.5 or greater for at least one primary demographic subgroup. Figure 2 provides an overview of these results.

Panel (a) presents the distributions of AM ratios across all of the 21 detected chemicals, shown separately for each primary demographic subgroup. And again, panel (b) lists the subset of individual chemicals for which at least one primary demographic subgroup had a AM ratio of 1.5 or greater. With the exception of relatively high exposure to the legacy organochlorine pesticide DDT (ratio = 2.01), the distribution of AM ratios for Hispanic individuals fell entirely below a value of 1. The AM ratio distribution for the non-Hispanic Asian subgroup, however, had a long right tail; we observed elevated exposure levels relative to national averages for DDT (ratio = 3.42), another legacy organochlorine pesticide beta-hexachlorocyclohexane (ratio = 9.22), and seven individual PCBs. For all other demographic subgroups, we did not find evidence of substantially different exposure levels among this set of 21 carcinogens.

Sex- and Age-differentiated results

In supplemental results, we also assessed AM ratios differentiated by sex and age group. These additional AM ratio results are available in Table A8 and Table A9, with subgroups’ distributions visualized in Figure A3. For the subset of chemicals considered here, two additional findings are worth noting. First, exposure levels were generally higher for the age 20+ subgroup. This held across all races and ethnicities and aligns with the fact that all 21 chemicals analyzed here are no longer in active use. Second, exposure to PCBs among the non-Hispanic Asian subgroup appeared more elevated for females, while exposure to DDT was relatively higher among males.

3.1.3 95th percentile ratios

Due to the limited number of individually-surveilled carcinogens for which we can calculate GM ratios, we next turn to a secondary biomonitoring analysis of 95th percentile ratios. These statistics inform us about cross-subgroup differences that may exist *among the most highly exposed individuals*. While there is no

¹⁸In our cross-analysis synthesis in Section 3.3, we treated all PCBs as a single carcinogen class, but in the results here we presented separate statistics for individual PCB compounds.

inherent reason that subgroups’ P95 and GM ratio measures should be correlated, one might be curious how well the former metric can serve as a proxy for the latter. In this particular context, we found limited evidence of such correlation.¹⁹ When calculating ratios disaggregated only by age or only by race/ethnicity, one could make a modest case for the P95 ratio acting as a meaningful signal of the GM ratio, but for all other disaggregations these correlations were weaker. Nevertheless, P95 ratios were informative of possible exposure disparities in their own right; EPA’s Guidelines for Human Exposure Assessment suggest consideration of high-end exposure levels to ensure protection of potentially exposed or susceptible subpopulations (EPA, 2019).

There were a sufficient fraction of detected biomarker samples to calculate P95 ratios for 75 carcinogen metabolites and parent chemicals (representing 62 carcinogens in total). Of those, 14 total carcinogens – measured as 8 parent chemicals and 6 metabolites – returned a GM ratio of 1.5 or greater for at least one demographic subgroup. Figure 3 provides an overview of these results.

Panel (a) presents the distributions of P95 ratios across all of the detected chemicals, shown separately for each primary demographic subgroup. These distributions were similar to those observed when assessing the GM ratios, though nuanced differences existed. The distribution of P95 ratios for lower income individuals was again skewed higher than the distribution for higher income individuals. For Hispanic, non-Hispanic Other, and non-Hispanic White subgroups, the distributions were again centered at median values close to 1, though the dispersion of the latter’s distribution was considerably tighter than the others. The P95 ratio distribution for the non-Hispanic Black subgroup again had a median value of greater than 1.25 and upper tail extremes larger than observed in the subgroup’s GM ratio distribution.

Panel (b) lists the subset of individual chemicals for which at least one primary demographic subgroup had a P95 ratio of 1.5 or greater. Many of the chemicals that had at least one GM ratio above 1.5 also appeared on this “high-end” exposure list. Among others, these included 1,4-dichlorobenzene (non-Hispanic Black ratio = 12.67), nitrofen (non-Hispanic Black ratio = 7.21), acrylonitrile (non-Hispanic Black ratio = 1.96), and arsenic (non-Hispanic “Other” ratio = 1.64). Contrasting these particular P95 ratios with their corresponding GM ratios indicated that cross-subgroup differences in high-end exposure to these chemicals were even more substantial than average differences. Additionally, several chemicals without flagged GM ratio differences of greater than 1.5 were found to have at least one P95 ratio greater than 1.5. The non-Hispanic Black subgroup showed evidence of elevated high-end exposure to a number of common industrial chemicals including ethylene thiourea (ratio = 2.77), ethylene oxide (ratio = 1.6), vinyl chloride (ratio = 1.6), acrylamide (ratio = 1.68), and perfluorooctanesulfonic acid (i.e. PFOS, ratio = 1.5), as well as the organophosphorus pesticide diazanon (ratio = 1.91). The lower-income demographic subgroup showed a difference in high-end exposure to furan (ratio = 1.51), while results for the non-Hispanic “Other” subgroup also suggested elevated high-end exposure to the pesticide dimethylarsinic acid (i.e. DMA, ratio = 1.66). The complete set of P95 ratio results for primary demographic subgroups – including chemicals without any detected P95 ratio above 1.5 – is provided in Table A10.

Demographic interaction results

In supplemental results, we also assessed P95 ratios by demographic interactions of sex and race/ethnicity and demographic interactions of household income and race/ethnicity. These additional P95 ratio results are available in Table A11, with subgroups’ distributions visualized in panel (a) of Figure A2. When considering demographic interactions, a broad range of chemicals presented with P95 ratios greater than 1.5 for at least

¹⁹See Figure A1 for visual evidence based on the set of carcinogens for which we could calculate both GM and P95 ratios.

one subgroup. In total this set included 44 carcinogens (represented in NHANES by 26 parent chemicals and 24 metabolites). Some chemicals of industrial and/or regulatory note that were flagged here but not in the previously-described GM ratio analyses nor the primary P95 ratio analysis include 1,3-butadiene, 1-bromopropane, perfluorooctanoic acid (i.e. PFOA), benzene, tetrachloroethylene, and diisononyl phthalate (i.e. DINP). Describing the results more broadly, though the most extreme P95 ratio values were well above their corresponding GM ratio values, on the whole, these more granular subgroups had P95 ratio distributions that tracked similarly to their GM ratio distributions. The prevalence of P95 ratios greater than 1.5 for the non-Hispanic Black subgroup again predominantly originated from elevated chemical exposure among males and lower-income individuals.

Age-differentiated results

In supplemental results, we also assessed P95 ratios by age group. These additional P95 ratio results are available in Table A12, with subgroups’ distributions visualized in panel (b) of Figure A2. When considering P95 ratios strictly differentiated by age group (age 19 and younger or age 20+), we did not identify any values greater than 1.5. Joint consideration of age and race/ethnicity while assessing P95 ratios did uncover some exposure differences. High-end melamine exposure was notably elevated for Hispanic individuals aged 19 and younger (ratio = 2.15), as were exposure to 1,4-dichlorobenzene and nitrofen (ratios = 3.28, 2.1). We also observed differential high-end exposure to di(2-ethylhexyl) phthalate and malathion for younger non-Hispanic “Other” individuals (ratios = 1.72, 1.64), and differential high-end exposure to 1,4-dichlorobenzene, nitrofen, and malathion for younger non-Hispanic Black individuals (ratios = 12.5, 4.95, 3.11).

3.1.4 Detected sample ratios

Before turning to our analysis of indirect exposure metrics, we briefly describe an additional set of supplemental results based on the NHANES data. In lieu of distributional statistics based on observed biomonitoring concentrations, we examined a categorical measurement of heterogeneous exposure levels across demographic subgroups in the form of *sample detection ratios*. The sample detection ratio for a given carcinogen and demographic subgroup was defined as the subgroup’s survey-weighted fraction of samples with concentrations above the detection limit divided by the total population’s survey-weighted fraction of samples with concentrations above the detection limit. This supplemental measure discretely indicates whether an individual was exposed to a carcinogen (above the detection limit) or less exposed (below the detection limit). Because this categorical metric only requires a single detectable sample for a carcinogen to be included in the analysis, we were able to assess 125 matched carcinogens and carcinogen-associated metabolites, of which 96 were based on individual-level biomonitoring samples and 29 were based on pooled samples.

We present a summary distributional visualization of these supplemental results in Figure A4, with complete results provided for primary demographic subgroups in Table A13, demographic interactions in Table A14, and age-differentiated subgroups in Table A15. Across the set of 125 matched carcinogens and metabolites, the non-Hispanic Black subgroup – in particular those older than 20 – had a higher median sample detection ratio than other primary demographic subgroups. Beyond this, there was little evidence suggesting systematic differences in sample detection ratios across subgroups.²⁰

²⁰Carcinogens for which at least one primary demographic subgroup’s sample detection ratio exceeded 1.5 include: **1,1,2,2-tetrachloroethane**, 1,3-butadiene, 1,4-dichlorobenzene, **4-methyl-2-pentanone**, **acetochlor**, acrylonitrile, arsenic acid, benzene, bromoform, carbon tetrachloride, chloroethane, cumene, diaznon, **dichloromethane**, furan, **Glu-P-1**, **Glu-P-2**, **lindane**, MeA-alpha-C, **N-nitroso-N-methylethylamine**, **N-nitrosodiethylamine**, **N-nitrosomorpholine**, **N-nitrosopiperidine**, **N-nitrosopyrrolidine**, PCB66, PCB81, TCDD (dioxin), tetrachloroethylene, **tetrahydrofuran**,

3.2 Indirect measures of exposure

Given that NHANES biomonitoring data covered a relatively small share of identified carcinogens, we now turn to indirect evidence of differences in exposure based on national-level data sets for ambient air, drinking water, and consumer products.

3.2.1 Ambient Air concentrations

We begin with an assessment of potential differences in residential ambient exposure to individual carcinogens via air releases from industrial sources. Recall that we overlaid the geographic location of demographic subgroups with the geographic locations of individual chemicals’ airborne releases and calculated spatially-weighted ambient concentrations for each subgroup. Our metric for identifying differences in exposure for a given carcinogen is the ratio of a subgroup’s *average* ambient air concentration and the national average ambient air concentration. We were able to match 186 candidate chemicals to the RSEI data, of which 136 ultimately had an average ambient concentration ratio of greater than 1.5 for at least one subgroup of interest.

Figure 4 summarizes several key findings from this analysis; the full set of results are in Appendix Table A16 for the interested reader.²¹ In panel (a), we graphed the distributions of relative concentration ratios for all 186 carcinogens, differentiating by subgroup. For each density plot, a black line indicates the distribution’s median value. For all racial/ethnic groups except non-Hispanic Black, the median concentration ratio value was below 1. In fact, for the non-Hispanic Asian, non-Hispanic American Indian/Alaskan Native, and non-Hispanic Pacific Islander subgroups, much of the entire distribution laid below 1, indicating that any relative exceedances in ambient exposure were limited to just a handful of chemicals. Conversely (and by mathematical construction), the median ambient air concentration ratio value for Black individuals was around 1.6, and the subgroup’s distribution had a long right tail. For 102 chemicals – well more than half of the those for which we have data – the average ambient air concentration for a Black individual was at least 1.5 times higher than the national average.

We next draw attention to panel (b), which shows the set of carcinogenic chemicals for which (i) total reported airborne emission volumes were greater than 150,000 pounds in the 2022 TRI and (ii) there was evidence of average exposure differences across population subgroups. These chemicals ranged in industrial and spatial extent of use. With the exception of sulfuric acid aerosols, which were primarily emitted from electric utility generation, many of the carcinogens listed were common to the chemical manufacturing sector. Indeed, across the 17 other chemicals highlighted, 47% of the ambient air emissions were reported from the chemical manufacturing sector, 12% from the metal manufacturing sector, and 11% from petroleum manufacturing. These emissions from the 17 chemicals overwhelmingly originated from facilities in the Southeast, Midwest, and Central-South, which respectively accounted for 24%, 17%, and 34% of total 2022 TRI emissions.

Panel (c) displays the set of carcinogens for which any subgroup had a concentration ratio greater than 4. As expected from panel (a), many of these exceptionally high ambient air concentration ratios were

trichloroethylene, and **Trp-P-1**. **Bolded** chemicals in above list are those for which there were insufficient detected samples for calculation of GM and P95 ratios.

²¹For both the ambient air and drinking water indirect exposure metrics, we also calculated spatially-weighted 95th percentile concentrations for each subgroup. A summary of the corresponding ambient air concentration ratios at 95th percentile values is presented in Table A17 for completeness. Results were qualitatively similar, though limited in coverage. This was due to the sizable share of TRI chemicals for which the national 95th percentile concentration was zero; concentration ratios could not be calculated when the denominator was zero.

indicative of non-Hispanic Black individuals’ markedly higher potential for residential exposure. Notably, however, none of the carcinogens listed in panel (c) were on the list in panel (b), suggesting that the strongest measures of potential exposure disparity generated by this statistical exercise derived from chemicals with highly local exposure patterns. In effect, these chemicals were emitted in lower total volumes and from fewer facilities, but their emissions were concentrated in neighborhoods with a higher-than-average fraction of non-Hispanic Black residents.

3.2.2 Drinking water

We next examined differences in exposure to individual carcinogens via public drinking water. For the 15 matched chemicals with sufficient detection rates for analysis, we calculated relative average concentration ratios that were nationally representative for households served by public water systems. Figure 5 summarizes the findings.²² We identified only a limited number of chemicals and subgroups for which substantial exposure differences existed. This may have been partially due to our statistical approach: any within-distribution-system differences in exposure were smoothed when we took a system-level average, and furthermore, the spatial scale of the analysis was implicitly more spatially aggregated due to water systems being larger than Census block groups. See Appendix B for further discussion on this.

In panel (a), we again graphed the distributions of relative concentration ratios, differentiating by demographic subgroup. The median concentration ratios for all seven subgroups were around 1. Only three individual concentration ratios were above the 1.5 threshold value for the American Indian and Pacific Islander subgroups. In panel (b), we present the complete set of drinking water relative concentration ratios, where we identified potential disparities in exposure to arsenic (ratio = 1.75) and chromium (VI) (ratio = 1.84) for American Indian individuals. These heavy metals are often naturally occurring but also likely affected by activities such as mining and energy production (Erickson et al., 2024); actual exposure levels may actually exceed the estimated values due to the high use rates of private groundwater wells for drinking water on Tribal lands (Spaur et al., 2021). For Pacific Islander individuals, the average drinking water concentration of chromium (VI) was almost two times higher than the national average (ratio = 1.99). This was largely driven by the residential concentration of Pacific Islander individuals across western states. Of the 50 CWSs serving the highest number of Pacific Islander residents, 46 were located in HI, NV, CA, UT, AZ, OR, and WA. CWSs in AZ had the highest state average concentration of chromium (VI) in drinking water. CWSs on tribal lands in the Pacific southwest had the second-highest average, CWSs in HI had the fourth-highest state average concentration, and CWSs in CA had the sixth-highest state average concentration. These higher chromium (VI) concentrations were naturally occurring in most cases.

3.2.3 Household products

Our final indirect exposure metric focused on consumer products that contain one or more carcinogens. Recall that a subgroup’s prevalence rate is defined as the total fraction of household-months in which a carcinogen appeared in the purchased consumption bundle. Using these rates, we calculated purchase prevalence ratios

²²A summary of the ($N = 17$) corresponding drinking water concentration ratios at 95th percentile values is presented in Table A18. Results were qualitatively similar, though the 95th percentile drinking water concentration for Hispanic individuals was substantially higher for dibromoacetic acid (a disinfection byproduct, ratio = 1.62) and trichloroethylene (ratio = 1.92). A summary of the ($N = 66$) drinking water *sample detection* ratios for matched carcinogens is presented in Table A19. Sample detection ratios results for drinking water should be interpreted conservatively, as detection limits for a given chemical may vary both across CWSs and within a CWS and be correlated with the size of population served and/or the population’s demographics. Nevertheless, broad trends in the sample detection ratio results suggest that Hispanic individuals, in particular, had a higher than average probability of carcinogens being detected in their drinking water.

for each of the 44 candidate carcinogens that appeared in the chemical-household data panel, again with national average purchase prevalence rates as the reference value. Figure 6 summarizes the key findings from our analysis. In panel (a), we graphed the distributions of relative prevalence ratios, differentiating by subgroup. Across the six demographic subgroups considered, the median prevalence ratio values were again consistently close to one. For the Black, Hispanic, and Asian subgroups we found that several carcinogens had prevalence ratios greater than 1.5, which we investigated further in panel (b).

This list includes the 21 chemicals for which at least one subgroup was found to have relatively elevated exposure, including highly-elevated benzophenone purchase prevalence ratios for Asian households. This chemical is a flavoring agent and fragrance enhancer and our results showed that Asian individuals were greater than 700% more likely to purchase products containing it than the average national consumer. Other chemicals with notably elevated prevalence ratios included oxidized asphalt (a common ingredient in coatings for construction, electronics, and packaging; ratio = 2.48) for Asian households, phenolphthalein (a common color-changing acid-base indicator and pharmaceutical drug ingredient in laxatives and weight loss products; ratio = 2.25) for Asian or Hispanic households and those with children, coal tar (used in skin care treatments and as a base for construction coatings and paint; ratio = 2.72) and 1,4-dichlorobenzene (ratio = 2.7) for Black households, and di(2-ethylhexyl) phthalate (ratio = 3.57) for Hispanic individuals.

Also noteworthy is the elevated melamine purchase prevalence ratio for Black households; melamine is commonly used in kitchenware and as a flame retardant in plastics, paints, and paper. This finding contrasts with the earlier biomonitoring results from NHANES, which indicated *lower than average* exposure to melamine for non-Hispanic Black individuals. A possible combined interpretation of these results is that despite Black households' relatively higher purchase rates of consumer products that contain melamine, the amounts of the chemical or uptake rates from this pathway either did not translate into elevated exposure or other exposure pathways dominated such that a disparity did not show up in urine, blood, or serum samples collected via NHANES. Further investigation of this interpretation is warranted, but the results nonetheless underscore the value of jointly assessing biomonitoring and consumer purchase data. See Table A20 for the complete set of results for all 44 matched chemicals.

3.3 Synthesis of results: carcinogens with the strongest evidence of exposure disparities

We conducted five main analyses of differences in exposure to individual carcinogens. To synthesize our findings across the analyses, we highlight carcinogens that met the set of inclusion criteria. Table 1 presents the 28 carcinogens which met our joint inclusion criteria. Individual flags in the table denote the analyses that did or did not result in at least one differential exposure ratio of 1.5 or greater for a specific chemical. Of these 28 chemicals, 6 are known carcinogens, 14 are probable carcinogens, 6 are possible carcinogens, and 2 are not classified by the IARC or RoC.

We found robust evidence of exposure disparities for a handful of chemicals across the multiple analyses. Arsenic was one such chemical; at least one demographic subgroup had an elevated exposure ratio in each of the five analyses that were conducted. Elevated exposure to arsenic for American Indian individuals was especially widespread. The results from the biomonitoring, ambient air, and drinking water analyses all supported a finding of potentially elevated exposure to the carcinogen for this subgroup. We also found evidence of elevated exposure to this chemical for females of childbearing age based on purchasing patterns identified in the household products analysis.

Other chemicals that demonstrated evidence of elevated exposure across multiple analyses for specific

subgroups via both biomonitoring and indirect measures include ethylene oxide, 1,4-dichlorobenzene, acrylonitrile, melamine, furan, and 1-naphthylamine. These chemicals are commonly used in both industrial processes and household products. Interestingly, di(2-ethylhexyl) phthalate and 1,2-propylene oxide had no biomonitoring ratio values greater than 1.5 across the primary demographic subgroups, but were flagged in indirect analyses for having at least one subgroup with ambient air concentration and household product prevalence ratios greater than 1.5. This motivated consideration of two supplemental sets of alternative inclusion criteria.

Alternative inclusion criteria

Our chosen synthesis inclusion criteria relied on researcher discretion in two key dimensions. The first was our reliance on NHANES-derived statistics for **only** primary demographic subgroups. Results for subgroups created by demographic interactions of sex, race/ethnicity, income, and age can tell a more complex story.²³ With this in mind, in Table A21 we present a supplemental synthesis list in which biomonitoring results based on demographic interactions were equally considered. The second was our reliance on a ratio threshold value of 1.5. As some carcinogens may have had demographic subgroups with ratio values that fell just below this strict threshold, we present a supplemental synthesis list constructed using a ratio threshold of 1.25 in Table A22.²⁴ In both supplemental tables, the list of carcinogens was qualitatively similar but included a much broader range of chemicals. The former list contained 53 carcinogens and the latter contained 69. While the additional chemicals added to the supplemental synthesis lists merit consideration in future research, we focus our attention on the default inclusion list in the remainder of this paper: this more streamlined synthesis list contains the chemicals for which we had the strongest evidence of cross-demographic differences in exposure.

Chemicals with limited exposure differences across subgroups

Lastly, we examined the extent to which our analyses demonstrated **no evidence** of exposure disparities for some carcinogens. Using the alternative results based on the more relaxed ratio threshold of 1.25 as a conservative approach, Table A23 compiles this list of chemicals. Only a single chemical showed fairly compelling evidence of homogeneously ubiquitous exposure across the population: formaldehyde. Given the carcinogen’s widespread use in home building materials and furniture, as well as its general ambient ubiquity due to industrial and vehicular combustion emissions, it is intuitive that our results point to near-universal exposure. We found that the evidence for the remaining set of chemicals in the table were inconclusive.²⁵ More than 10 other chemicals showed evidence of limited differences in exposure across subgroups based on results from a single indirect exposure media, but the lack of multidimensional robustness rendered any inference less compelling.

²³For example in the case of di(2-ethylhexyl) phthalate, age may be an important factor to consider. Our air concentration ratios for the chemical showed non-Hispanic Asian and Pacific Islander subgroups may face elevated ambient exposure, while the household product prevalence ratio was greater than 1.5 for Hispanic households. Biomonitoring results differentiated by race/ethnicity and age for Hispanic and non-Hispanic “Other” individuals aged 19 and younger had values greater than 1.5.

²⁴For example, 1,2-propylene oxide – which as discussed above had no biomonitoring metrics but two indirect exposure metrics greater than 1.5 – had a GM ratio exceeding the relaxed threshold of 1.25.

²⁵Limited biomonitoring results for two arsenic acids and 2,3,4,7,8-pentachlorodibenzofuran showed no strong evidence of potential exposure disparities across groups, but the former was complicated by the metabolization patterns of arsenic, and the latter lacked support from indirect exposure metrics.

3.4 Main exposure pathways for carcinogens with potential exposure disparities

We next conducted a review of data and literature describing exposure pathways and product categories for each carcinogen we identified as having potential exposure disparities in Table 1. There are two primary purposes of this exercise. The first is identification of additional carcinogen-specific exposure pathways beyond drinking water, ambient pollution, and household products that may have contributed to the differential exposure we observed in biomonitoring data. The second is to highlight the most common exposure pathways and major product categories across the set of carcinogens for which we identified evidence of notable differences in exposure across subgroups.

Figure 7 visually summarizes the available information on exposure pathways and product categories for chemicals with potential exposure disparities. The complete set of chemical-specific information collected about exposure pathways – including product categories, points of exposure, and exposure routes – as well as associated cancer types can be found in Table 3.

Panel (a) explores the general relevance of several exposure pathways, distinguishing between occupational and general population exposure. Among the carcinogens we identified with potential exposures disparities, we found that occupational exposures most frequently occur via manufacturing (20 of 28 chemicals), farming and agriculture (16/28), through processing of chemicals directly or as intermediaries (16/28), through industrial cleaning products (7/28), and through vehicle use, production, or maintenance (7/28). Turning to general population exposure, the most common pathways for exposure to this set of carcinogens included food and food additives (23 of 28 chemicals), tobacco products (14/28), personal care products (10/28), home maintenance products (10/28), home vehicle use or maintenance (8/28), home cleaning products (7/28), and art supplies (7/28). Additionally, we observed that environmental releases (13/28) and drinking water (13/28) were general population exposure pathways for a large portion of the carcinogens. Such broad incidence of these two pathways corroborated the findings from the drinking water and ambient air indirect exposure analyses presented earlier in this section.

Panel (b) characterizes the relative frequency of carcinogens’ association with different product categories. 21 of 28 chemicals were associated with use as an herbicide or pesticide. This high frequency included some legacy pesticide uses of carcinogens, but nonetheless aligned with the evidence above indicating that the occupational exposure pathway, farming/agriculture, was a possible source of differential carcinogen exposure. Other common product use categories that were potentially relevant to occupational exposure included metals and electronics (12 of 28 chemicals), plastics and rubber (15/28), and paint and adhesives (11/28). Common product use categories that were potentially relevant to general population exposure included food and food additives (15/28), pharmaceuticals (13/28), and textiles (14/28).

Recall that we did not collect information about, nor formally model how much exposure occurred via any specific pathway or product category. These results simply reflected the state of information available from our four primary sources regarding each individual carcinogen, and we made no additional assumptions about intensity of exposure or potential unlisted pathways. It was therefore not possible to make explicit claims about which particular pathways or product categories drove observed demographic differences in exposure. Taken as group, however, the pathways and product categories highlighted above, due to their frequent association with the 28 carcinogens on our list, likely merit further scrutiny in future work exploring exposure disparities.

3.5 Correlations between potential exposure disparities and cancer incidence rates

Disparities in cancer incidence by sex, race, and ethnicity are well-established (Singh and Jemal, 2017; Zavala et al., 2021; Siegel et al., 2024). While we cannot state the relative contribution of different exposure pathways to differences in body burden, nor how differences in body burden translate to disparities in cancer incidence, we can highlight the extent to which differences in exposure are positively correlated with differences in cancer incidence. Table 2 presents the five most recent years of national cancer incidence data (2017-2021) from the CDC (CDC, 2021b). These incidence rates were disaggregated by cancer type, sex, race, and ethnicity, and age-adjusted to account for potentially confounding subgroup age distributions. We included statistics for all leading and invasive cancer types. In what follows, we compared incidence rates across subgroups and noted cancer types that occurred with high frequency for a particular subgroup. To supplement our previously described findings, we contrasted these elevated cancer incidence rates with the cancer types typically caused by carcinogens that were shown to have exposure differences across subgroups.²⁶ Though any connections we drew between *current* cancer rates and our *concurrent* exposure metrics are purely correlational – most cancers have a latency period of several years – such findings may point to avenues worthy of investigation in future work.

There are a handful of cancer types for which multiple demographic subgroups showed substantially-higher-than-average incidence. Among non-Hispanic American Indian, non-Hispanic Asian/Pacific Islander, and Hispanic males, the incidence of liver-related cancers for males, respectively, were 50%, 68%, and 62% higher than the national male average. American Indian, Asian/Pacific Islander, and Hispanic females, respectively, had 100%, 82%, and 82% higher incidence of liver-related cancer than the national female average.²⁷ Stomach cancer was substantially more common among Asian, Black, and Hispanic individuals. For females, incidence rates were respectively 77%, 72%, and 74% higher than the female national average. For males, incidence rates were respectively 73%, 80%, and 58% higher than the male national average.²⁸ Cervical cancer incidence was also slightly higher than national average for American Indian (28%), Black (35%), and Hispanic (45%) females.

Turning to columns (4) and (5) of Table 3, liver-related cancers were associated with several chemicals for which we found large exposure differences across subgroups. In particular, epidemiological evidence has linked liver cancer with exposure to arsenic, 1,4-dichlorobenzene, diethanolamine, PCBs, vinyl chloride, and DDT.²⁹ When paired with findings from our direct and indirect exposure analyses, we inferred potential relationships between (1) Hispanic individuals’ historic and ongoing exposure to pesticides and/or ambient diethanolamine and the subgroup’s highly elevated liver cancer incidence, (2) American Indian individuals’ historic and ongoing exposure to arsenic and/or pesticides and the subgroup’s highly elevated liver cancer incidence, (3) Asian/Pacific Islander individuals’ historically elevated exposure to several pesticides and PCBs and the subgroup’s highly elevated liver cancer incidence, and (4) Black men’s historic and ongoing exposure to 1,4-dichlorobenzene, PCBs, and vinyl chloride, and the subgroup’s elevated liver cancer incidence.

²⁶Table A24 displays cancer *mortality* rates, again differentiated by cancer type, sex, race, and ethnicity. These statistics reflect the well-documented variation in mortality rates across demographic subgroups. We included this information for completeness and the interested reader, but did not analyze them further here – disparities in mortality from cancer result from an even more complex set of environmental, social, and exposure factors that are beyond the scope of this work.

²⁷The statistics also showed elevated liver-related cancer incidence among non-Hispanic Black males, though this 35% deviation from the national average was more modest.

²⁸The statistics also showed slightly elevated stomach-related cancer incidence among American Indian individuals: 32% higher than the female national average and 30% higher than the male national average.

²⁹Using animal-based evidence, 9 additional carcinogens with evidence of exposure disparities – furan, 4-methyl-2-pentanone, ethylene thiourea, nitrofen, PFOS, DEHP, Beta-HCH, NNK, dimethylarsinic acid – were associated with liver cancers.

Undertaking a similar exercise for chemicals with identified differences in exposure across subgroups that were associated with stomach cancers, we found epidemiological evidence of associations for arsenic, acrylonitrile, ethylene oxide, and diethanolamine.³⁰ When paired with findings from our direct and indirect exposure analyses, we inferred potential relationships between (1) Hispanic individuals’ historic and ongoing exposure to 1,4-dichlorobenzene, 1,2-propylene oxide, and diethanolamine and the subgroup’s highly elevated stomach cancer incidence, (2) American Indian individuals’ historic and ongoing exposure to arsenic and the subgroup’s elevated stomach cancer incidence, (3) Asian/Pacific Islander individuals’ historic and ongoing exposure to 1,2-propylene oxide and the subgroup’s highly elevated stomach cancer incidence, and (4) Black individuals’ historic and ongoing exposure to 1,4-dichlorobenzene, acrylonitrile, ethylene oxide, and diethanolamine, and the subgroup’s elevated stomach cancer incidence.

Table 2 also shows evidence of incidence disparities for other cancer types, and these cancer types are often associated with carcinogens for which we observed related potential exposure disparities. Among Black males, prostate cancer incidence – which epidemiological evidence has associated to diethanolamine and PFOS exposures – were 86% above the national average. Among Black and White males, respiratory cancer incidence – which epidemiological evidence has associated to carbaryl, vinyl chloride, 4-methyl-2-pentanone, diazinon, acrylonitrile, and arsenic exposures – were respectively 51% and 31% above the national male average. Among Black males and females and White males, larynx cancer incidence – which epidemiological evidence has associated with naphthalene exposure – were respectively 89%, 64%, and 27% above national averages. Lastly, among Black individuals, the incidence of colorectal cancers – which epidemiological evidence has associated with 2,4D, acrylonitrile, PCB, naphthalene, and acrolein exposures – were elevated compared to national averages for both males (43%) and females (38%).

To conclude this subsection, we compiled a full set of related information on exposure disparities, carcinogen-to-cancer-type associations, and cancer type incidence rates into a heatmap in Figure 8. Each cell in the heatmap denotes a demographic subgroup-cancer type pairing. Cell values (and color) denote the count of carcinogens for which we jointly found (1) subgroup-specific biomonitoring evidence of differences in exposure, and (2) epidemiological support for an association between the carcinogen and the cancer type. Visually, darker red cells are those with higher carcinogen counts, and cells with a heavy black border denote subgroup-cancer type pairings for which national statistics showed incidence was greater than the national average. Because the available cancer incidence statistics were decomposed by race/ethnicity and sex, the results in this table focused strictly on biomonitoring evidence of potentially exposure disparities for the corresponding subgroups.

Several components of the heatmap indicate alignment between our broader exposure findings and the cancer incidence statistics. Across subgroups, we found elevated incidence of liver cancers and potential disparities in exposure to at least one associated carcinogen for each group. Similarly, we found that Black males faced potential exposure disparities to multiple carcinogens associated with colorectal, liver, lung, and stomach cancers; national statistics for Black males suggest elevated incidence rates for these cancer types. Other results highlight the complexity in linking exposure metrics to cancer incidence. Among the subgroups considered here, we found broad-based potential disparities in exposure to carcinogens associated with blood cancers (leukemia, lymphomas), bladder cancers, and kidney cancers, though current data showed that only non-Hispanic American Indian females had a higher incidence of the latter cancer type. For other cancer types – cervical/uterine, esophageal, and gallbladder, among others – we observed differences in incidence

³⁰Using animal-based evidence, two additional carcinogens with observed exposure disparities were associated with stomach cancer: 1,4-dichlorobenzene and 1,2-propylene oxide.

across subgroups but lacked information on carcinogens that may have contributed through differing exposure rates. These findings emphasize that environmental factors are only a single contributor to the development of cancer.

4 Discussion

The results from our analyses in Section 3 strictly focused on carcinogens for which nationally representative exposure data were available. These analyses have some inherent methodological limitations. But perhaps more importantly, as shown in columns 3-7 of Table A1, we lacked information for the vast majority of carcinogens in each of our analyses. The lack of chemical breadth in available exposure information is well documented (Egeghy et al., 2012), but these data limitations continue to impede a full understanding of which carcinogens pose the greatest potential risk of exposure and subsequent health effects for specific subgroups. With an eye to improving future analyses of a similar nature, section 4.1 outlines the key limitations and caveats that pertain to our results, and section 4.2 describes some key data gaps that we identified while conducting our analyses.

4.1 Limitations

While we leveraged the available national-level data in a rigorous fashion, there remain several limitations and caveats related to the analyses. Across all analyses, we first emphasize that lack of information about a chemical – due either to limits of detection, or the nonexistence of environmental emissions and biomonitoring data – does not imply an absence of disparity in exposure. There were substantial big-picture data gaps that limited our analyses, a subject to which we return in the next section.

We begin with limitations related to the NHANES analyses. First, we emphasize the temporal variation in source data utilized across carcinogens. To present results relevant to current national exposure levels, we used only the most recent NHANES cycle for each carcinogen. Unfortunately, not every chemical was consistently monitored over time. While our results were largely based on biomonitoring from the past 10 years, results for some chemicals – namely, legacy pesticides including dieldrin and heptachlor epoxide B – were based on data from 2003-2004. These particular legacy chemicals slowly biodegrade in the environment, but it is possible that historic exposure disparities have diminished over the pursuant decades. Second, we emphasize our reliance on biomonitoring concentrations as a reasonable proxy for true exposure. This strong assumption ignores variation in elimination half-life, frequency of exposure, and numerous biological factors (Stanfield et al., 2022; Aylward et al., 2012, 2014). We aimed to mitigate these concerns by employing a statistical focus on comparisons of strictly within-chemical variation in subgroup exposure, but we acknowledge that the methods used here stopped well short of actual exposure modeling.

We next turn to limitations related to the RSEI ambient air concentrations analysis. Two caveats related to the underlying data are important to note. First, RSEI is built from TRI point-source air emissions data, and therefore is representative of possible exposure to chemicals emitted to the air from large industrial facilities in a subset of regulated industries and for a subset of chemicals. Ambient air concentrations of a chemical do not necessarily reflect locals’ individual-level exposure. Smaller airborne industrial releases of reportable chemicals were not represented in RSEI’s concentration measures, nor were releases from non-industrial or non-point source emitters (cars, biogenic sources, etc.). TRI are also self-reported data, and therefore sometimes estimated instead of directly measured. Second, in instances where very few facilities reported releases of a given chemical (e.g. bis(2-Chloro-1-methylethyl) ether, which was released by two

Gulf Coast facilities in 2022), subgroup concentration ratios were largely determined by a relatively small number of census block groups in nearby communities. Thus, though the calculated concentration ratios were derived from nationally representative, population-weighted averages – and therefore are an informative national measure of potential disparities – it is important to recognize that ambient exposure to some of these chemicals is highly localized.

There are three caveats to mention related to the drinking water exposure analysis. The first emphasizes an obvious but important point: the data underlying our results were sourced from drinking water quality samples collected by CWSs, and are only reflective of actual chemical exposure insofar as residents rely on tap water for personal consumption. Drinking water intake varies across demographics subgroups, and future exposure work of this nature should aim to quantitatively account for these differences (Brooks et al., 2017; Rosinger et al., 2022).³¹ The second relates to the national representativeness of the analysis; we could not assess tap water quality for households receiving water from non-reporting CWSs nor households reliant on private wells. We showed in Appendix C that for 12 of the 15 carcinogens analyzed, the drinking water concentration ratios were representative for at least 2/3 of the US population, including those residing in most major urban and suburban areas. The remaining three carcinogens analyzed are bromate, microcystin-LR, and N-nitrosodimethylamine; the statistics for these carcinogens were constructed from less nationally-representative data, and therefore we advise caution in their interpretation. Lastly, to the extent that CWSs’ non-reporting of detection limits – and thus, their true chemical concentrations – vary systematically with the demographics of their constituents, our assumptions regarding the imputation of sample values below level of detection limits could bias concentration ratio estimates.

With respect to the household product analysis, we note three caveats. First, the statistics in this setting were derived from a measure of chemical presence in households’ purchased bundles of goods. In the data, we did not directly observe the magnitude, frequency, and duration of product use, which household member used the product, or whether products were used in unintended ways. Therefore the statistics were a proxy for true exposure. Second, Stanfield et al. (2021) noted several potential limitations of their data; of most relevance for our application are (1) potential concerns about selection bias vis-à-vis the substantial number of products for which chemical ingredient information could not be matched, and (2) potential concerns about subgroups having differential rates of specialty, ethnic, or neighborhood store visits where products were missing UPC matches or were less likely to be scanned. In either case, our results based on a non-randomly selected set of data may potentially misrepresent true subgroup prevalence ratios. And third, the Nielson scanner data underlying this analysis were from 2012; in the intervening years, it is possible that product formulations and consumer buying patterns evolved such that exposures driven by previously observed differences in purchase prevalence were mitigated (or vice versa).

Our literature and data review concerning exposure pathways and associated product categories was limited in nature. To keep the analysis tractable, our review strictly considered information from four authoritative sources, and therefore our findings were constrained in scope. For less commonly used carcinogens, it is possible that some currently active exposure categories were omitted and some obsolete exposure pathways were included due to outdated and/or missing information from our sources. For example, we only noted environmental releases as an exposure pathway for carcinogens where naturally-occurring or industrial pollution were explicitly identified. In instances where such pollution was not explicitly noted in our data sources, it was omitted as a relevant exposure pathway. Furthermore, we did not collect information

³¹We note, however, that for dermal or inhalation exposure, exposure via showering, hand-washing, produce-washing, and other direct interactions with tap water may still be an important source of exposure via tap water.

about, nor model how much exposure occurs via any specific pathway or product category, and therefore we only made claims about potentially common pathways. A more complete exposure analysis is necessary to identify the particular pathway(s) and product(s) driving potential exposure disparities.

Consideration of temporal changes over a longer time-period could inform a more formal future exposure analysis. Changes in uses of chemicals over time are expected. This, in turn, changes emission patterns, levels observed in environmental emissions data, and observed biomonitoring data. Changes in consumer product use patterns, particularly for longer-term or frequent users of specific products will influence exposures in the indoor environment. Changes in industrial use patterns will influence longer-term occupational exposures and releases into the environment, such as through ambient air or drinking water.

It is also worth noting that while exposure pathways can often be cleanly delineated by the affected population (general versus occupational), some points of exposure for a given carcinogen may be less distinct. Products in a given category can be used in different settings, and therefore many product or use categories include both occupational and general population pathways of exposure. To illustrate, one example was the cleaning product category: janitors or dishwashers may be chronically exposed to a carcinogenic product in their day-to-day work, while the general population may use a consumer-focused version of the same cleaning product in their home (Calderon et al., 2022). Another example was the vehicle product use category. Occupational exposure could involve dermal contact or inhalation of carcinogenic vehicle maintenance products by automobile mechanics or long-distance truckers, while general population exposure to carcinogens while using vehicles could result from ambient off-gassing of flame retardants in seating (Hoehn et al., 2024). To the extent that information to make these nuanced joint delineations was unavailable, we may have misallocated a given exposure to a single affected population, rather than to both.

We conclude this section by noting that most carcinogens we studied were associated with multiple product categories. And similarly, many product categories were associated with more than 10 known carcinogens. As such, the potential for multiple exposures to interact through similar (or differing) modes of action is apparent. Though we stop short of a more formal analysis in this work, we emphasize the need for future research to contextualize cumulative exposure via occupational activities, consumer products, or other sources affecting the general population (Markowitz et al., 2013; Rodgers et al., 2018), account for behavioral risk factors associated with cancer like tobacco or alcohol use, and explore how exposure patterns correlate with identified windows of particular susceptibility (Terry et al., 2019; Shaffer et al., 2017).

4.2 Notable data gaps

One key takeaway from our empirical efforts is the degree to which the evaluation of heterogeneity in potential exposure to carcinogenic chemicals by race, ethnicity, income or sex was data-constrained, especially at the national level. As described in Section 2.2.1, only 106 of the 622 ubiquitous carcinogens analyzed were surveilled by NHANES between 1998 and 2018. Going one step further, only 36 of these carcinogens had both biomonitoring data and environmental monitoring via the TRI and drinking water sampling. If we were to make the most complete analysis possible, including only the set of carcinogens for which we were *also* able to assess exposure via household product purchases using proprietary data, the list shrinks to 10 chemicals.³² Reframing these limitations in the most charitable possible light: we were able to produce one or more usable direct or indirect exposure analyses for only 230 (37%) of 622 identified carcinogens. This is not sufficient for a thorough assessment of exposure disparities: reducing the size of these monitoring gaps

³²For the interested reader: these ten chemicals are: 1,4-dichlorobenzene, 1,4-dioxane, 2,4D, arsenic, cadmium, cobalt, DEHP, dichloromethane, ethylbenzene, and perchloroethylene.

is an immediate research need.

We ask two questions regarding the limited information we have at hand. First, were known carcinogens more likely to be monitored in the data we collected? The second panel of Table A1 lays out evidence showing that this was not systematically the case. Of the 96 candidate chemicals that are known carcinogens, 17% were directly monitored by NHANES. Comparable statistics for probable carcinogens, possible carcinogens, and other carcinogens (i.e. those solely listed as carcinogenic by CA Proposition 65, the EU, or the UK) are 17%, 13%, and 8%, respectively.³³ Second, were chemicals that had a particular product use category more or less likely to be monitored? Table A3 summarizes findings for a selected subset of product use categories. Among these common categories, carcinogens used in cleaning and safety products were biomonitoring with slightly higher frequency than those used in other categories. Chemicals used in home maintenance and personal care products were monitored in drinking water with the highest frequency. And perhaps unsurprisingly, chemicals found in cleaning products and personal care products were most frequently observed in the household products purchasing data. Taken together, however, these monitoring frequency differences across product categories were small, offering limited evidence that chemicals used in certain product categories had a higher propensity of being monitored.

A second key takeaway is that even when biomonitoring or environmental monitoring did occur, current laboratory analytical methods were a limiting factor. For more than half of the matched carcinogens in NHANES, we could not calculate subgroup-specific geometric mean ratios. Even when calculating 95th percentile ratios, we could only produce statistics for 80% of matched chemicals. Meeting the detection limit was even rarer when analyzing drinking water concentrations. Only 15 of the 70 matched carcinogens (21%) had a sufficient fraction of detected drinking water samples for production of nationally-representative concentration statistics. More broadly, our results illustrated that detection limit problems could be compounding when conducting a subgroup-focused statistical analysis: as subgroup categories became more and more disaggregated, we found that it was increasingly challenging to maintain a sufficient fraction of detected samples for calculation of subgroup-specific summary statistics.

A third key takeaway is related to the quality and accessibility of exposure pathway information. For most chemicals, we found that *some* degree of information was available on common exposure pathways and product uses. But despite a reliance on a structured information collection process spanning authoritative data sources, our efforts revealed the typically incomplete nature of compiled information. Exposure information was often segmented, with only a subset of production use categories or exposure routes listed by a single data source. For some chemicals, we found that PubChem contained a holistic set of information related to product uses, exposure pathways, and both human- and animal-linked cancer endpoints but for others the information available in PubChem was dated or more limited. In general, the vintage of information was especially problematic in relation to current use and exposure patterns. Much of the available information about occupational carcinogen exposure dated from the 1980s, and it was unclear whether many listed uses were still active today. Overall, the range of encountered data limitations demonstrated to us the potential value of coordinating a centralized exposure database that is regularly updated.³⁴

³³For ambient air quality, TRI reporting occurred for 23% of known carcinogens, compared to 41.2%, 25%, and 23%, respectively. For drinking water monitoring, 19% of known carcinogens were monitored, compared to 11%, 7%, and 8%, respectively.

³⁴As a preliminary step in identifying potential exposure disparities for the many carcinogens that lacked biomonitoring or indirect measures of exposure, we conducted a supplemental analysis of readily-available information on exposure pathways and major product categories for the complete set of *known carcinogens*. Perhaps surprisingly, most known carcinogens lacked national-level biomonitoring data. See Appendix B for results and a brief survey of findings from this analysis. We took particular note of carcinogens, exposure pathways, and product categories that shared suggestive associations with elevated cancer incidence rates.

In summary, we identified several data gaps that could benefit from future research attention. Opportunities to expand biomonitoring have grown in recent years; continued federal support in terms of grants and person-power could broaden the set of chemicals surveilled, and improve laboratory methods and detection limits for chemicals that are currently biomonitored. For similar reasons, innovation in state biomonitoring programs could pay large dividends. New state-level methods and exposure findings could help Federal health agencies in their efforts to prioritize additional chemicals for national-level surveillance. And lastly, we note that there is broad scope for improving indirect exposure research. For example, a nationally-representative survey that conducts chemical analysis of household dust wipes, indoor air samples, outdoor soil, and collects an inventory of consumer goods present in the home at the time of sampling could provide information that is highly complementary to more direct biomonitoring metrics.³⁵ A nationally-representative survey and database on foodborne chemical exposure could be another valuable resource for exploring the drivers of chemical exposure disparities. Finally, while it is beyond the scope of this paper to identify where the highest added value would be if resources were spent addressing some of these carcinogen-specific data gaps, we note that others have considered questions of a similar spirit using rigorous research methods that could be applied here (Pellizzari et al., 2019; Braun and Escher, 2023).

5 Conclusion

In this paper, we characterized the current state of readily-available and nationally-representative exposure information for a set of 622 ubiquitous carcinogens. Usable data on direct or indirect exposure were available for only 232 of these carcinogens. With the collected data, we calculated demographically-differentiated exposure statistics for individual carcinogens. Using these statistics, we identified a subset of 28 chemicals for which there is evidence of potential exposure disparities. We also reviewed readily-available, carcinogen-specific exposure information, highlighted common exposure pathways and major product categories among the subset of carcinogens, and noted carcinogen-demographic pairings where potential exposure disparities are correlated with elevated national cancer incidence rates. Our findings point to several exposure pathways and product categories that may merit particular attention during efforts to reduce differences in chemical exposure and cancer incidence. While conducting our analysis, we noted several data gaps and provided a discussion of where additional data gathering and future study may be useful.

Code and Data availability

All R code and non-proprietary data for this paper are available at https://github.com/adamtheising/ubiquitous_carcinogens.

³⁵The US Department of Housing and Urban Development already conducts the occasional American Healthy Home Survey, but to date, their chemical exposure analyses have been limited to lead, formaldehyde, and some pesticides.

References

- Austin, W., T. Bardot, and A. R. El-Khattabi (2024). Comparative analysis of service area boundaries and disparities in drinking water quality. *National Center for Environmental Economics Working Paper*.
- Aylward, L. L., S. M. Hays, R. Smolders, H. M. Koch, J. Cocker, K. Jones, N. Warren, L. Levy, and R. Bevan (2014). Sources of variability in biomarker concentrations. *Journal of Toxicology and Environmental Health, Part B* 17(1), 45–61.
- Aylward, L. L., C. R. Kirman, J. L. Adgate, L. M. McKenzie, and S. M. Hays (2012). Interpreting variability in population biomonitoring data: role of elimination kinetics. *Journal of exposure science & environmental epidemiology* 22(4), 398–408.
- Barr, K. J., C. L. Johnson, J. Cohen, P. D’Souza, E. I. Gallegos, C.-C. Tsai, A. L. Dunlop, E. J. Corwin, D. B. Barr, P. B. Ryan, et al. (2022). Legacy chemical pollutants in house dust of homes of pregnant african americans in atlanta. *Toxics* 10(12), 755.
- Belova, A., S. L. Greco, A. M. Riederer, L. E. Olsho, and M. A. Corrales (2013). A method to screen us environmental biomonitoring data for race/ethnicity and income-related disparity. *Environmental Health* 12, 1–17.
- Bertin, M., J. Bodin, N. Fouquet, N. Bonvallot, Y. Roquelaure, et al. (2018). Multiple exposures and coexposures to occupational hazards among agricultural workers: a systematic review of observational studies. *Safety and health at work* 9(3), 239–248.
- Bi, C., J. P. Maestre, H. Li, G. Zhang, R. Givhechi, A. Mahdavi, K. A. Kinney, J. Siegel, S. D. Horner, and Y. Xu (2018). Phthalates and organophosphates in settled dust and hvac filter dust of us low-income homes: association with season, building characteristics, and childhood asthma. *Environment international* 121, 916–930.
- Braun, G. and B. I. Escher (2023). Prioritization of mixtures of neurotoxic chemicals for biomonitoring using high-throughput toxicokinetics and mixture toxicity modeling. *Environment International* 171, 107680.
- Brender, J. D., J. A. Maantay, and J. Chakraborty (2011). Residential proximity to environmental hazards and adverse health outcomes. *American journal of public health* 101(S1), S37–S52.
- Brooks, C. J., S. L. Gortmaker, M. W. Long, A. L. Cradock, and E. L. Kenney (2017). Racial/ethnic and socioeconomic disparities in hydration status among us adults and the role of tap water and other beverage intake. *American journal of public health* 107(9), 1387–1394.
- Brown, J. A. (2024). Haz-map: Information on hazardous chemicals and occupational diseases. <https://haz-map.com/>.
- Calafat, A. M., X. Ye, L.-Y. Wong, A. M. Bishop, and L. L. Needham (2010). Urinary concentrations of four parabens in the us population: Nhanes 2005–2006. *Environmental health perspectives* 118(5), 679–685.
- Calderon, L., R. Maddalena, M. Russell, S. Chen, J. E. Nolan, A. Bradman, and K. G. Harley (2022). Air concentrations of volatile organic compounds associated with conventional and “green” cleaning products in real-world and laboratory settings. *Indoor air* 32(11), e13162.

- Chan, M., C. Mita, A. Bellavia, M. Parker, and T. James-Todd (2021). Racial/ethnic disparities in pregnancy and prenatal exposure to endocrine-disrupting chemicals commonly used in personal care products. *Current Environmental Health Reports* 8(2), 98–112.
- Dionisio, K. L., K. Phillips, P. S. Price, C. M. Grulke, A. Williams, D. Biryol, T. Hong, and K. K. Isaacs (2018). The chemical and products database, a resource for exposure-relevant data on chemicals in consumer products. *Scientific data* 5(1), 1–9.
- Dodson, R. E., M. Nishioka, L. J. Standley, L. J. Perovich, J. G. Brody, and R. A. Rudel (2012). Endocrine disruptors and asthma-associated chemicals in consumer products. *Environmental health perspectives* 120(7), 935–943.
- Dodson, R. E., J. O. Udesky, M. D. Colton, M. McCauley, D. E. Camann, A. Y. Yau, G. Adamkiewicz, and R. A. Rudel (2017). Chemical exposures in recently renovated low-income housing: Influence of building materials and occupant activities. *Environment international* 109, 114–127.
- Du, S., L. Rodenburg, N. Patterson, C. Chu, C. D. Riker, C. H. Yu, and Z. T. Fan (2020). Concentration of polychlorinated biphenyls in serum from new jersey biomonitoring study: 2016–2018. *Chemosphere* 261, 127730.
- Egeghy, P. P., R. Judson, S. Gangwal, S. Mosher, D. Smith, J. Vail, and E. A. C. Hubal (2012). The exposure data landscape for manufactured chemicals. *Science of the Total Environment* 414, 159–166.
- Erickson, M. L., C. J. Brown, E. J. Tomaszewski, J. D. Ayotte, J. K. Böhlke, D. B. Kent, and S. Qi (2024). Prioritizing water availability study settings to address geogenic contaminants and related societal factors. *Environmental Monitoring and Assessment* 196(3), 303.
- Hefferon, R., D. E. Goin, J. A. Sarnat, and A. E. Nigra (2024). Regional and racial/ethnic inequalities in public drinking water fluoride concentrations across the us. *Journal of exposure science & environmental epidemiology* 34(1), 68–76.
- Helm, J. S., M. Nishioka, J. G. Brody, R. A. Rudel, and R. E. Dodson (2018). Measurement of endocrine disrupting and asthma-associated chemicals in hair products used by black women. *Environmental research* 165, 448–458.
- Hoehn, R. M., L. G. Jahl, N. J. Herkert, K. Hoffman, A. Soehl, M. L. Diamond, A. Blum, and H. M. Stapleton (2024). Flame retardant exposure in vehicles is influenced by use in seat foam and temperature. *Environmental Science & Technology* 58(20), 8825–8834.
- International Agency for Research on Cancer (2024). List of classifications by cancer sites with sufficient or limited evidence in humans, iarc monographs volumes 1–137. https://monographs.iarc.who.int/wp-content/uploads/2019/07/Classifications_by_cancer_site.pdf.
- Isaacs, K. K., J. T. Wall, A. R. Williams, K. A. Hobbie, J. R. Sobus, E. Ulrich, D. Lyons, K. L. Dionisio, A. J. Williams, C. Grulke, et al. (2022). A harmonized chemical monitoring database for support of exposure assessments. *Scientific Data* 9(1), 314.
- Islami, F., J. Baeker Bispo, H. Lee, D. Wiese, K. R. Yabroff, P. Bandi, K. Sloan, A. V. Patel, E. C. Daniels, A. H. Kamal, et al. (2024). American cancer society’s report on the status of cancer disparities in the united states, 2023. *CA: A Cancer Journal for Clinicians* 74(2), 136–166.

- James-Todd, T. M., Y.-H. Chiu, and A. R. Zota (2016). Racial/ethnic disparities in environmental endocrine disrupting chemicals and women’s reproductive health outcomes: epidemiological examples across the life course. *Current epidemiology reports* 3, 161–180.
- Kapraun, D. F., J. F. Wambaugh, C. L. Ring, R. Tornero-Velez, and R. W. Setzer (2017). A method for identifying prevalent chemical combinations in the us population. *Environmental health perspectives* 125(8), 087017.
- Kim, S., J. Chen, T. Cheng, A. Gindulyte, J. He, S. He, Q. Li, B. A. Shoemaker, P. A. Thiessen, B. Yu, et al. (2023). Pubchem 2023 update. *Nucleic acids research* 51(D1), D1373–D1380.
- Kim, S., P. A. Thiessen, T. Cheng, B. Yu, and E. E. Bolton (2018). An update on pug-rest: Restful interface for programmatic access to pubchem. *Nucleic Acids Research* 46(W1), W563–W570.
- Kim, S., P. A. Thiessen, T. Cheng, B. Yu, B. A. Shoemaker, J. Wang, E. E. Bolton, Y. Wang, and S. H. Bryant (2016). Literature information in pubchem: associations between pubchem records and scientific articles. *Journal of cheminformatics* 8, 1–15.
- LaKind, J. S., J. R. Sobus, M. Goodman, D. B. Barr, P. Fürst, R. J. Albertini, T. E. Arbuckle, G. Schoeters, Y.-M. Tan, J. Teeguarden, et al. (2014). A proposal for assessing study quality: Biomonitoring, environmental epidemiology, and short-lived chemicals (bees-c) instrument. *Environment international* 73, 195–207.
- Latshaw, M. W., R. Degeberg, S. S. Patel, B. Rhodes, E. King, S. Chaudhuri, and J. Nassif (2017). Advancing environmental health surveillance in the us through a national human biomonitoring network. *International Journal of Hygiene and Environmental Health* 220(2), 98–102.
- Madrigal, J. M., A. Flory, J. A. Fisher, E. Sharp, B. I. Graubard, M. H. Ward, and R. R. Jones (2024). Sociodemographic inequities in the burden of carcinogenic industrial air emissions in the united states. *JNCI: Journal of the National Cancer Institute* 116(5), 737–744.
- Mariotto, A. B., L. Enewold, J. Zhao, C. A. Zeruto, and K. R. Yabroff (2020). Medical care costs associated with cancer survivorship in the united states. *Cancer epidemiology, biomarkers & prevention* 29(7), 1304–1312.
- Markowitz, S. B., S. M. Levin, A. Miller, and A. Morabia (2013). Asbestos, asbestosis, smoking, and lung cancer. new findings from the north american insulator cohort. *American journal of respiratory and critical care medicine* 188(1), 90–96.
- Martinez-Morata, I., B. C. Bostick, O. Conroy-Ben, D. T. Duncan, M. R. Jones, M. Spaur, K. P. Patterson, S. J. Prins, A. Navas-Acien, and A. E. Nigra (2022). Nationwide geospatial analysis of county racial and ethnic composition and public drinking water arsenic and uranium. *Nature communications* 13(1), 7461.
- McGoldrick, D. J. and E. W. Murphy (2016). Concentration and distribution of contaminants in lake trout and walleye from the laurentian great lakes (2008–2012). *Environmental Pollution* 217, 85–96.
- McKelvey, W., R. C. Gwynn, N. Jeffery, D. Kass, L. E. Thorpe, R. K. Garg, C. D. Palmer, and P. J. Parsons (2007). A biomonitoring study of lead, cadmium, and mercury in the blood of new york city adults. *Environmental Health Perspectives* 115(10), 1435–1441.

- McKelvey, W., J. B. Jacobson, D. Kass, D. B. Barr, M. Davis, A. M. Calafat, and K. M. Aldous (2013). Population-based biomonitoring of exposure to organophosphate and pyrethroid pesticides in new york city. *Environmental Health Perspectives* 121(11-12), 1349–1356.
- Mehta, S. S., I. Morin, K. Osborn, C. R. Lemeris, M. Conti, and R. M. Lunn (2023). An approach to assessing the influence of environmental and occupational cancer hazard identification on policy decision-making. *Environmental Health Perspectives* 131(12), 125001.
- Mitro, S. D., R. E. Dodson, V. Singla, G. Adamkiewicz, A. F. Elmi, M. K. Tilly, and A. R. Zota (2016). Consumer product chemicals in indoor dust: a quantitative meta-analysis of us studies. *Environmental science & technology* 50(19), 10661–10672.
- Nassif, J., A. M. Calafat, and K. M. Aldous (2021). The us national biomonitoring network—enhancing capability and capacity to assess human chemical exposures. *International Journal of Hygiene and Environmental Health* 237, 113828.
- Nguyen, V. K., A. Kahana, J. Heidt, K. Polemi, J. Kvasnicka, O. Jolliet, and J. A. Colacino (2020). A comprehensive analysis of racial disparities in chemical biomarker concentrations in united states women, 1999–2014. *Environment international* 137, 105496.
- Nguyen, V. K., L. Y. Middleton, L. Huang, N. Zhao, E. Verly Jr, J. Kvasnicka, L. Sagers, C. J. Patel, J. Colacino, and O. Jolliet (2023). Harmonized us national health and nutrition examination survey 1988–2018 for high throughput exposome-health discovery. *medRxiv*.
- Nigra, A. E., Q. Chen, S. N. Chillrud, L. Wang, D. Harvey, B. Mailloux, P. Factor-Litvak, and A. Navas-Acien (2020). Inequalities in public water arsenic concentrations in counties and community water systems across the united states, 2006–2011. *Environmental health perspectives* 128(12), 127001.
- O’Keefe, E. B., J. P. Meltzer, and T. N. Bethea (2015). Health disparities and cancer: racial disparities in cancer mortality in the united states, 2000–2010. *Frontiers in public health* 3, 51.
- Pellizzari, E. D., T. J. Woodruff, R. R. Boyles, K. Kannan, P. I. Beamer, J. P. Buckley, A. Wang, Y. Zhu, D. H. Bennett, and E. influences on Child Health Outcomes (2019). Identifying and prioritizing chemicals with uncertain burden of exposure: opportunities for biomonitoring and health-related research. *Environmental health perspectives* 127(12), 126001.
- Ravalli, F., Y. Yu, B. C. Bostick, S. N. Chillrud, K. Schilling, A. Basu, A. Navas-Acien, and A. E. Nigra (2022). Sociodemographic inequalities in uranium and other metals in community water systems across the usa, 2006–11: a cross-sectional study. *The Lancet Planetary Health* 6(4), e320–e330.
- Rodgers, K. M., J. O. Udesky, R. A. Rudel, and J. G. Brody (2018). Environmental chemicals and breast cancer: An updated review of epidemiological literature informed by biological mechanisms. *Environmental research* 160, 152–182.
- Rosinger, A. Y., A. I. Patel, and F. Weeks (2022). Examining recent trends in the racial disparity gap in tap water consumption: Nhanes 2011–2018. *Public health nutrition* 25(2), 207–213.
- Schildroth, S., K. M. Rodgers, M. Strynar, J. McCord, G. Poma, A. Covaci, and R. E. Dodson (2022). Per-and polyfluoroalkyl substances (pfas) and persistent chemical mixtures in dust from us colleges. *Environmental Research* 206, 112530.

- Shaffer, R. M., M. N. Smith, and E. M. Faustman (2017). Developing the regulatory utility of the exposome: mapping exposures for risk assessment through lifestage exposome snapshots (lens). *Environmental Health Perspectives* 125(8), 085003.
- Shaw, V., B. Zhang, M. Tang, W. Peng, C. Amos, and C. Cheng (2024). Racial and socioeconomic disparities in survival improvement of eight cancers. *BJC Reports* 2(1), 21.
- Siegel, R. L., A. N. Giaquinto, and A. Jemal (2024). Cancer statistics, 2024. *CA: a cancer journal for clinicians* 74(1).
- Singh, G. K. and A. Jemal (2017). Socioeconomic and racial/ethnic disparities in cancer mortality, incidence, and survival in the united states, 1950–2014: over six decades of changing patterns and widening inequalities. *Journal of Environmental and Public Health* 2017(1), 2819372.
- Spaur, M., M. A. Lombard, J. D. Ayotte, D. E. Harvey, B. C. Bostick, S. N. Chillrud, A. Navas-Acien, and A. E. Nigra (2021). Associations between private well water and community water supply arsenic concentrations in the conterminous united states. *Science of the Total Environment* 787, 147555.
- Stanfield, Z., C. K. Addington, K. L. Dionisio, D. Lyons, R. Tornero-Velez, K. A. Phillips, T. J. Buckley, and K. K. Isaacs (2021). Mining of consumer product ingredient and purchasing data to identify potential chemical coexposures. *Environmental Health Perspectives* 129(6), 067006.
- Stanfield, Z., R. W. Setzer, V. Hull, R. R. Sayre, K. K. Isaacs, and J. F. Wambaugh (2022). Bayesian inference of chemical exposures from nhanes urine biomonitoring data. *Journal of exposure science & environmental epidemiology* 32(6), 833–846.
- Stanfield, Z., R. W. Setzer, V. Hull, R. R. Sayre, K. K. Isaacs, and J. F. Wambaugh (2024). Characterizing chemical exposure trends from nhanes urinary biomonitoring data. *Environmental Health Perspectives* 132(1), 017009.
- Sun, M., E. Arevalo, M. Strynar, A. Lindstrom, M. Richardson, B. Kearns, A. Pickett, C. Smith, and D. R. Knappe (2016). Legacy and emerging perfluoroalkyl substances are important drinking water contaminants in the cape fear river watershed of north carolina. *Environmental science & technology letters* 3(12), 415–419.
- Tan, Y.-M., J. Sobus, D. Chang, R. Tornero-Velez, M. Goldsmith, J. Pleil, and C. Dary (2012). Reconstructing human exposures using biomarkers and other “clues”. *Journal of Toxicology and Environmental Health, Part B* 15(1), 22–38.
- Terry, M. B., K. B. Michels, J. G. Brody, C. Byrne, S. Chen, D. J. Jerry, K. M. Malecki, M. B. Martin, R. L. Miller, S. L. Neuhausen, et al. (2019). Environmental exposures during windows of susceptibility for breast cancer: a framework for prevention research. *Breast cancer research* 21, 1–16.
- US Census Bureau (2022). American community survey 5-year estimates. <https://www.census.gov/programs-surveys/acs/data.html>.
- US Centers for Disease Control and Prevention (2020a). Fourth national report on human exposure to environmental chemicals, updated tables, march 2021. volume three: Analysis of pooled serum samples for select chemicals, nhanes 2005-2016. <https://stacks.cdc.gov/view/cdc/105344>.

US Centers for Disease Control and Prevention (2020b). National health and nutrition examination survey data. <https://wwwn.cdc.gov/nchs/nhanes/Default.aspx>.

US Centers for Disease Control and Prevention (2021). Cdc wonder: United states cancer statistics. <https://wonder.cdc.gov/cancer.html>.

US Centers for Disease Control and Prevention (2024). Biomonitoring data tables for environmental chemicals. https://www.cdc.gov/exposurereport/data_tables.html.

US Environmental Protection Agency (2016a). The data management and quality assurance/quality control process for the third six-year review information collection rule dataset. Technical report, EPA-810-R-16-015.

US Environmental Protection Agency (2016b). Six-year review 3 compliance monitoring data (2006-2011). <https://www.epa.gov/dwsixyearreview/six-year-review-3-compliance-monitoring-data-2006-2011>.

US Environmental Protection Agency (2019). Guidelines for human exposure assessment. Technical report, EPA-100-B-19-001.

US Environmental Protection Agency (2020). Chemical data reporting data. <https://www.epa.gov/chemical-data-reporting/access-chemical-data-reporting-data>.

US Environmental Protection Agency (2022a). The data management and quality assurance/quality control process for the fourth six-year review information collection rule dataset. Technical report, EPA-810-R-22-001.

US Environmental Protection Agency (2022b). Microbial and disinfection byproduct data files (2012-2019) from epa's fourth six-year review information collection request. <https://www.epa.gov/dwsixyearreview/microbial-and-disinfection-byproduct-data-files-2012-2019-epas-fourth-six-year>.

US Environmental Protection Agency (2022c). Risk-screening environmental indicators model. <https://www.epa.gov/rsei/rsei-geographic-microdata-rsei-gm>.

US Environmental Protection Agency (2024a). Occurrence data from the unregulated contaminant monitoring rule (rounds 2-4). <https://www.epa.gov/dwucmr/occurrence-data-unregulated-contaminant-monitoring-rule>.

US Environmental Protection Agency (2024b). Toxic substances control act chemical substance inventory. <https://www.epa.gov/tsca-inventory/how-access-tsca-inventory>.

US National Cancer Institute (2024). Cancer statistics. <https://www.cancer.gov/about-cancer/understanding/statistics>.

US National Toxicology Program (2021). 15th report on carcinogens. <https://ntp.niehs.nih.gov/whatwestudy/assessments/cancer/roc>.

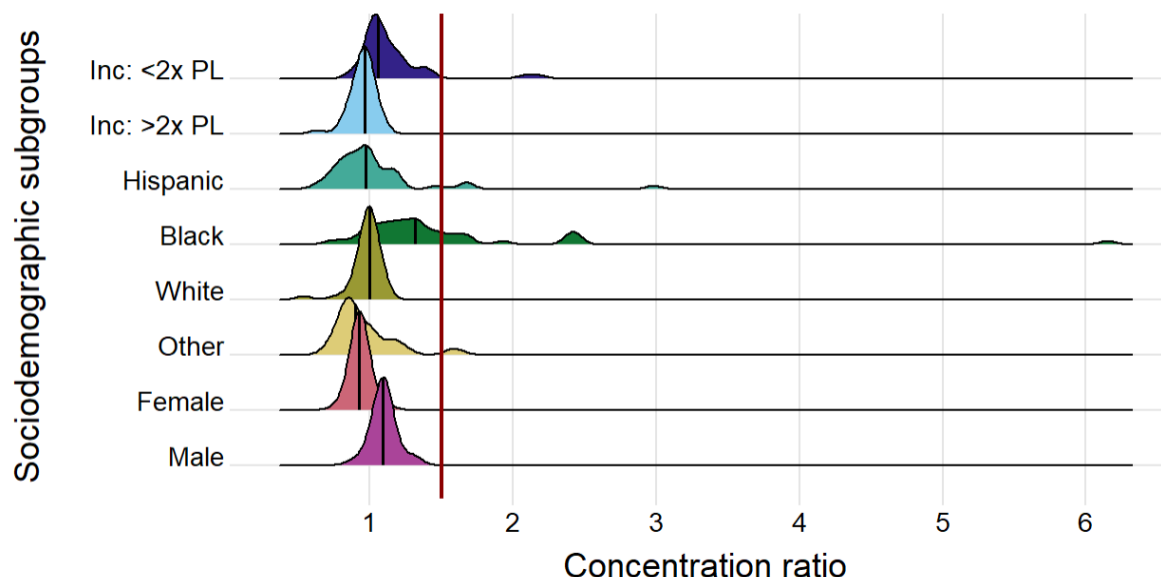
US National Toxicology Program (2022). Report on carcinogens data exploration dashboard. <https://public.tableau.com/app/profile/ntp.visuals/vizzes>. Internal-only dashboard containing animal carcinogen information. Contact NTP/NIH for access.

- Williams, A. J., C. M. Grulke, J. Edwards, A. D. McEachran, K. Mansouri, N. C. Baker, G. Patlewicz, I. Shah, J. F. Wambaugh, R. S. Judson, et al. (2017). The comptox chemistry dashboard: a community data resource for environmental chemistry. *Journal of Cheminformatics* 9, 1–27.
- Woodruff, T. J., A. R. Zota, and J. M. Schwartz (2011). Environmental chemicals in pregnant women in the united states: Nhanes 2003–2004. *Environmental health perspectives* 119(6), 878–885.
- Wu, S., S. Powers, W. Zhu, and Y. A. Hannun (2016). Substantial contribution of extrinsic risk factors to cancer development. *Nature* 529(7584), 43–47.
- Yu, C. H., C. D. Riker, S.-e. Lu, and Z. T. Fan (2020). Biomonitoring of emerging contaminants, perfluoroalkyl and polyfluoroalkyl substances (pfas), in new jersey adults in 2016–2018. *International Journal of Hygiene and Environmental Health* 223(1), 34–44.
- Zavala, V. A., P. M. Bracci, J. M. Carethers, L. Carvajal-Carmona, N. B. Coggins, M. R. Cruz-Correa, M. Davis, A. J. de Smith, J. Dutil, J. C. Figueiredo, et al. (2021). Cancer health disparities in racial/ethnic minorities in the united states. *British Journal of Cancer* 124(2), 315–332.

Figures

Figure 1: Summary of results: NHANES geometric mean analysis

(a) Distribution of geometric mean ratios across (N = 39) matched carcinogens, by demographic subgroup



(b) Selected results for chemicals with ratio ≥ 1.5

Chemical	Flags	Max Ratio	Demographic Subgroup
1,4-Dichlorobenzene*	3	6.15	Hispanic, Black, Inc: <2x PL
Nitrofen**	2	2.44	Hispanic, Black
2,4-Dichlorophenoxyacetic acid*	2	2.44	Hispanic, Black
4-(N-Methyl-N-nitrosamino)-1-(3-pyridyl)-1-butanone**	2	2.41	Black, Inc: <2x PL
Acrylonitrile**	1	2.39	Black
1-Naphthylamine	1	1.93	Black
Naphthalene**	1	1.69	Black
2-Naphthylamine	1	1.68	Black
Acrolein**	1	1.63	Black
Melamine	1	1.62	Other
1-Naphthalenol, 1-(N-methylcarbamate)**	1	1.57	Black
Arsenic	1	1.56	Other
2-Methylaniline	1	1.53	Black

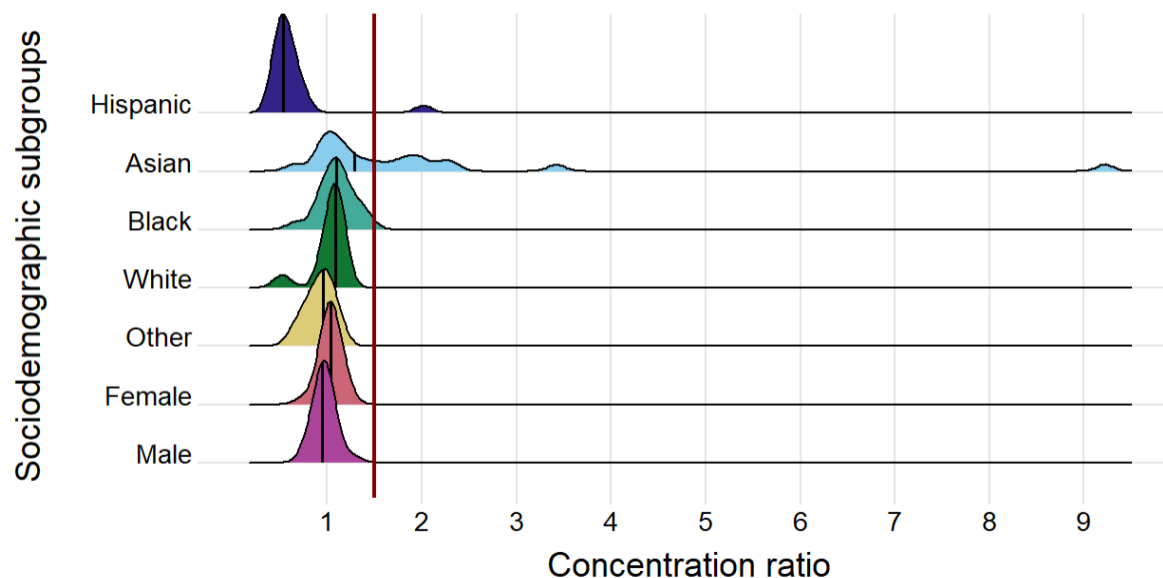
Notes: Panel (a) illustrates the full distribution of geometric mean concentration ratios across the 39 matched carcinogens, differentiated by demographic subgroup. Each subgroup's median geometric mean ratio is marked on their respective density plot with a black line. Panel (b) presents a selected set of results highlighting the set of chemicals for which at least one primary demographic subgroup was found to have a GMR of 1.5 or greater. Results in this figure are based on individual-level NHANES biomonitoring samples. *Flags* denote the number of subgroups for which the chemical had a prevalence ratio ≥ 1.5 . *Max Ratio* is the chemical's largest prevalence ratio value across all primary subgroups. The full set of geometric mean ratio results for these primary demographic subgroups are presented in Appendix Table A4. **: Carcinogens for which only metabolite biomonitoring samples were available. *: Carcinogens for which both direct and metabolite biomonitoring samples were available.

Demographic subgroups: Household income below 2 times the poverty line, Household income above 2 times the poverty line, Hispanic, non-Hispanic Black, non-Hispanic White, Other race including multi-racial, Female, Male.

Return to: Section 3.1.1.

Figure 2: Summary of results: NHANES pooled-sample mean analysis

(a) Distribution of arithmetic mean ratios across ($N = 21$) matched carcinogens, by demographic



(b) Selected results for chemicals with ratio ≥ 1.5

Chemical	Flags	Max Ratio	Demographics
DDT	2	3.42	Hispanic, Asian
Beta-Hexachlorocyclohexane	1	9.22	Asian
PCB146	1	2.33	Asian
PCB187	1	2.20	Asian
PCB126	1	2.02	Asian
PCB183	1	1.94	Asian
PCB178	1	1.84	Asian
PCB105	1	1.73	Asian
PCB167	1	1.56	Asian

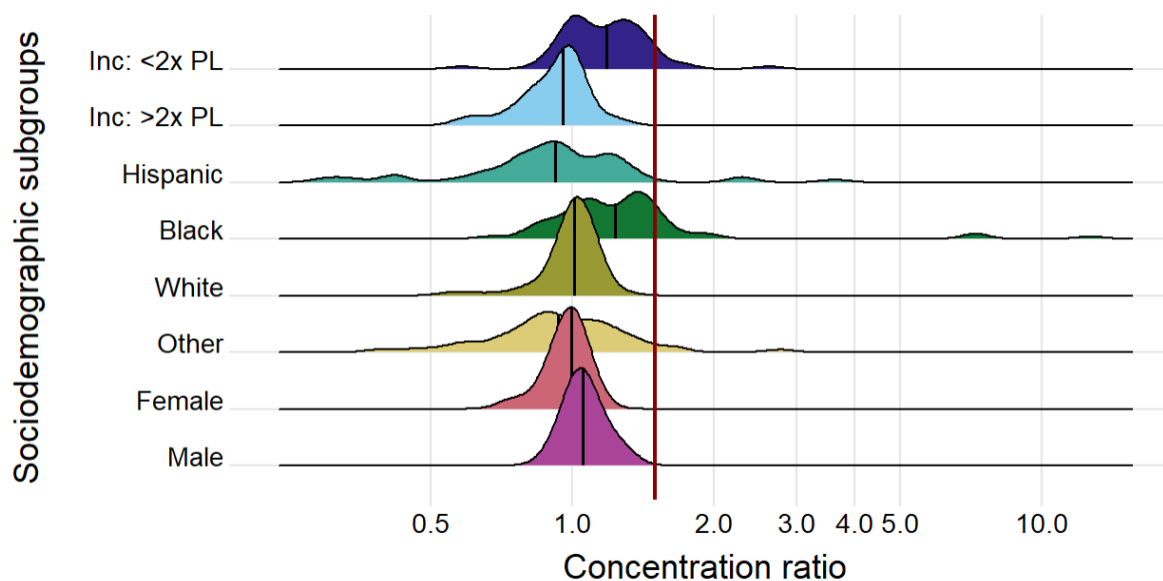
Notes: Panel (a) illustrates the full distribution of arithmetic mean concentration ratios across the 22 matched carcinogens, differentiated by demographic subgroup. Each subgroup's median arithmetic mean ratio is marked on their respective density plot with a black line. Panel (b) presents a selected set of results highlighting the set of chemicals for which at least one primary demographic subgroup was found to have a GMR of 1.5 or greater. Results in this figure are based on *pooled* NHANES biomonitoring samples. *Flags* denote the number of subgroups for which the chemical had a prevalence ratio ≥ 1.5 . *Max Ratio* is the chemical's largest prevalence ratio value across all primary subgroups. The full set of arithmetic mean ratio results for these primary demographic subgroups are presented in Appendix Table A7.

Demographic subgroups: Hispanic, non-Hispanic Asian, non-Hispanic Black, non-Hispanic White, Other race including multi-racial, Female sex, Male sex.

Return to: Section 3.1.2.

Figure 3: Summary of results: NHANES 95th percentile analysis

(a) Distribution of 95th percentile ratios across (N = 62) matched carcinogens, by demographic



(b) Selected results for chemicals with ratio ≥ 1.5

Chemical	Flags	Max Ratio	Demographics
1,4-Dichlorobenzene*	3	12.67	Hispanic, Black, Inc: <2x PL
Nitrofen**	3	7.21	Hispanic, Black, Inc: <2x PL
2,4-Dichlorophenoxyacetic acid*	3	7.21	Hispanic, Black, Inc: <2x PL
Ethylene thiourea	1	2.77	Other
Acrylonitrile**	1	1.96	Black
Diazinon**	1	1.91	Black
Acrylamide*	1	1.68	Black
Dimethylarsinic acid	1	1.66	Other
Arsenic	1	1.64	Other
Vinyl chloride**	1	1.60	Black
Ethylene oxide*	1	1.60	Black
Furan	1	1.51	Inc: <2x PL
1-Naphthylamine	1	1.51	Black
Perfluorooctanesulfonic acid	1	1.50	Black

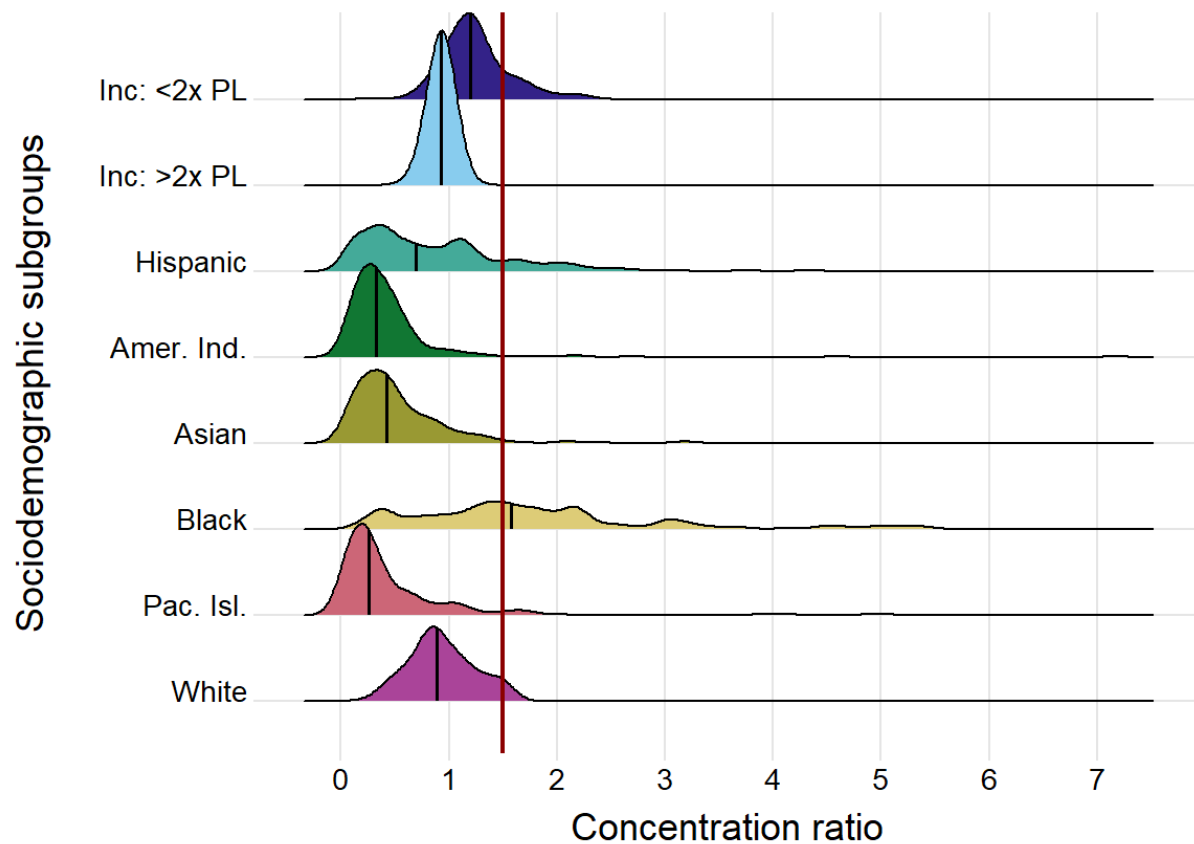
Notes: Panel (a) illustrates the full distribution of 95th percentile concentration ratios across the 62 matched carcinogens, differentiated by demographic subgroup. Each subgroup's median P95 ratio is marked on their respective density plot with a black line. Panel (b) presents a selected set of results highlighting the set of chemicals for which at least one primary demographic subgroup was found to have a GMR of 1.5 or greater. Results in this figure are based on individual-level NHANES biomonitoring samples. *Flags* denote the number of subgroups for which the chemical had a prevalence ratio ≥ 1.5 . *Max Ratio* is the chemical's largest prevalence ratio value across all primary subgroups. The full set of 95th percentile ratio results for these primary demographic subgroups are presented in Appendix Table A10. **: Carcinogens for which only metabolite biomonitoring samples were available. *: Carcinogens for which both direct and metabolite biomonitoring samples were available.

Demographic subgroups: Household income below 2 times the poverty line, Household income above 2 times the poverty line, Hispanic, non-Hispanic Black, non-Hispanic White, Other race including multi-racial, Female sex, Male sex.

Return to: Section 3.1.3.

Figure 4: Summary of results: air toxics analysis

(a) Distribution of relative air toxics exposure across (N = 186) matched carcinogens, by demographic



(b) Selected chems with high emissions and potential disparity

Chemical	Air emissions	Facilities	Demographics
Sulfuric acid	48,416,426.0	577	PI
4-Methyl-2-pentanone	3,294,525.7	476	Bl
Benzene	3,251,165.8	1,172	His, Bl
Dichloromethane	2,864,140.7	231	Bl
1,3-Butadiene	1,218,625.8	216	His
Naphthalene	1,157,890.4	1,288	His
Vinyl acetate	899,712.9	132	His
1-Bromopropane	869,649.6	62	Asn, PI
Cumene	846,218.1	578	His, Bl
Tetrachloroethylene	834,234.4	196	His
Vinyl chloride	469,094.4	34	Bl
1,2-Dichloroethane	457,365.2	52	Bl
Acrylonitrile	354,361.9	89	Bl
1,2-Propylene oxide	290,254.0	94	His
Chloroform	258,528.3	79	Bl
Vinyl fluoride	192,790.1	4	<2x PL, AmInd, Bl
Chloroethane	182,415.0	40	<2x PL, Bl
Ethylene oxide	166,356.4	146	Bl

(c) Select chems and subgroups with largest disparities in average ambient exposure (Ratio > 4)

Chemical	Ratio Value	Demographics
Bis(chloroethyl) ether	7.17	AmInd
1,4-Dichloro-2-butene	5.45	Bl
Diazinon	5.36	Bl
N-Nitrosodiphenylamine	5.28	Bl
Malathion	5.23	Bl
Propargite	5.11	Bl
4-Biphenylamine	5.08	Bl
1-Bromopropane	5.02	PI
1,2-Phenylenediamine	4.93	Bl
Nitrapyrin	4.92	PI
2-Methoxy-5-methylaniline	4.90	Bl
Furan	4.69	Bl
Acifluorfen sodium	4.68	Bl
Arsenic	4.58	AmInd
Parathion	4.50	Bl
Propoxur	4.47	Bl
Chloroprene	4.41	Bl
Rhodamine B	4.34	His
1,2-Benzenediol	4.15	PI
1,2-Dibromoethane	4.10	Bl

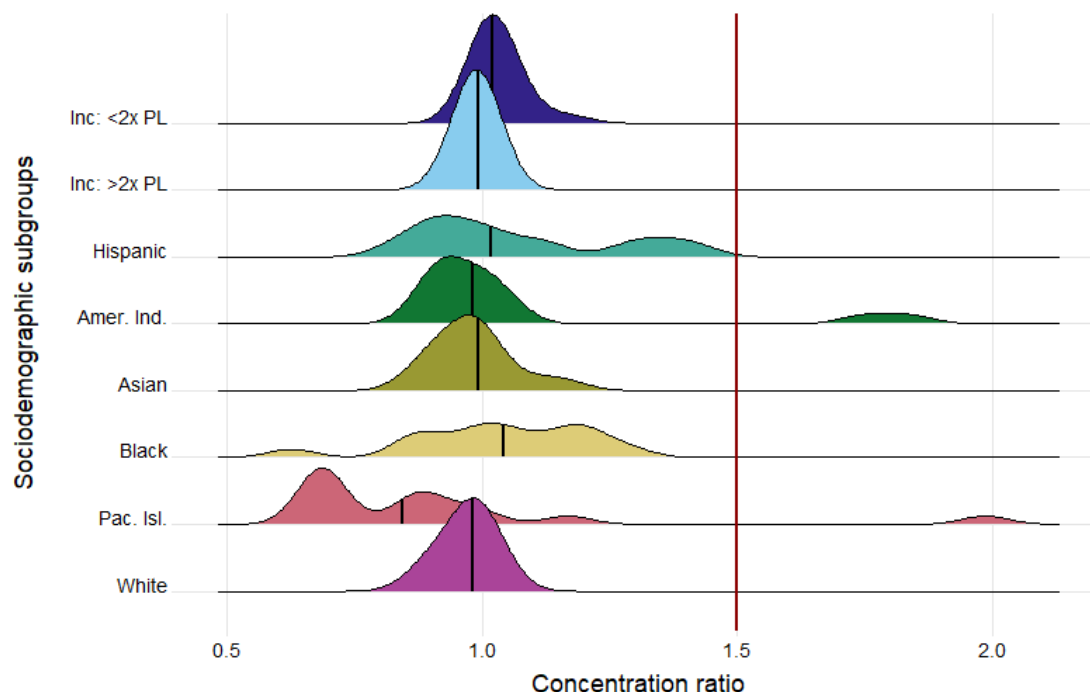
Notes: Panel (a) illustrates the across-chemical distribution of ambient average concentration ratios, differentiated by demographic subgroup. The ambient concentration ratio is an indirect metric summarizing demographic disparities in exposure to each carcinogen, and is measured by the ratio of a demographic subgroup’s *average* ambient concentration of the chemical and its national average ambient concentration. The median ambient concentration ratio for each subgroup line are denoted by the black vertical lines, while the threshold value of 1.5 is denoted by the red vertical line. Panel (b) shows a selected set of chemicals with high levels of national air releases (>150,000 pounds in 2022 TRI), and potential evidence of exposure disparities for at least one subgroup. Panel (c) highlights a selected set of chemicals which have an ambient concentration ratio value above 4 for at least one subgroup. The full set of ambient concentration ratios results is presented in Appendix Table A16.

Demographic subgroups: Hispanic (*His*), non-Hispanic American Indian or Alaskan Native (*AmInd*), non-Hispanic Asian (*Asn*), non-Hispanic Black (*Bl*), non-Hispanic Pacific Islander (*PI*), non-Hispanic White, household income below 2 times the poverty line (<2x *PL*), household income above 2 times the poverty line.

Return to: Section 3.2.1.

Figure 5: Summary of results: drinking water analysis

(a) Distribution of relative drinking water exposure across (N = 15) matched carcinogens, by demographic



(b) Drinking water average concentration ratios: complete results

Chemical	Hispanic	Amer. Ind.	Asian	Black	Pac. Isl.	White	<2x PL	>2x PL
1,4-DIOXANE	1.1	0.9	0.94	1.01	0.89	0.97	1.02	0.99
ARSENIC	1.02	1.75	0.87	0.87	0.88	1.03	0.99	1
BROMATE	0.94	0.93	1	1.04	1	1.02	1	1
CHROMIUM-6	1.42	1.84	1.16	0.63	1.99	0.9	1.07	0.97
DIBROMOACETIC ACID	1.28	0.93	0.96	0.86	0.97	0.94	1.04	0.98
DICHLOROACETIC ACID	0.89	0.99	0.99	1.22	0.68	0.99	1.02	0.99
DICHLOROMONOBROMOMETHANE	0.98	1	0.96	1.11	0.67	0.99	1	1
HAA5	0.91	0.9	1.02	1.19	0.69	0.99	1.01	0.99
HAA6BR	1.13	1.03	0.91	0.98	0.71	0.97	1.02	0.99
HAA9	0.95	0.93	0.99	1.15	0.68	0.98	1.01	0.99
MICROCYSTIN-LR	1.32	0.89	0.89	0.89	0.89	0.9	1.16	0.95
NDMA	1.02	0.98	0.93	1.06	0.84	0.98	1.03	0.99
BROMOFORM	1.38	0.92	0.99	0.97	1.17	0.89	1.04	0.98
TRICHLOROACETIC ACID	0.87	1.02	1.1	1.29	0.69	0.97	1.02	0.99
CHLOROFORM	0.82	1.05	1.01	1.2	0.68	1.02	1.01	0.99

Notes: Panel (a) illustrates the full distribution of chemical drinking water concentration ratios, differentiated by demographic subgroup. This concentration ratio is our metric of drinking water exposure disparities for each carcinogen, measured by the ratio of a demographic subgroup's *average* concentration of the chemical and its national average drinking water concentration. Each subgroups' median national concentration ratio is marked on the figure with a black line. Panel (b) presents subgroups' drinking water concentration ratios for each of the 15 chemicals that were monitored and detectable. **Bold** values denote concentration ratios ≥ 1.25 . Underlined values denote concentration ratios ≥ 1.5 .

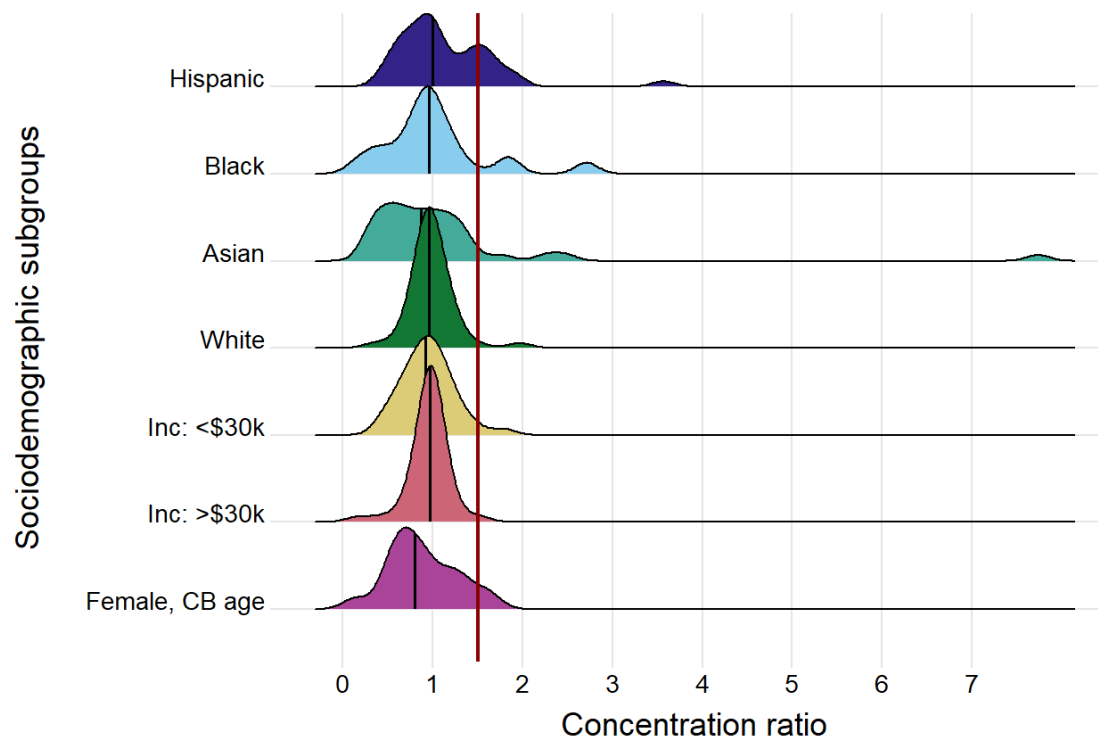
Demographic subgroups: Hispanic, non-Hispanic American Indian or Alaskan Native, non-Hispanic Asian, non-Hispanic Black, non-Hispanic Pacific Islander, non-Hispanic White, household income below 2 times the poverty line, household income above 2 times the poverty line.

Additional results: 95th percentile ratio results. Sample detection ratio results.

Return to: Section 3.2.2

Figure 6: Summary of results: household product purchases analysis

(a) Distribution of relative household product exposure across (N = 44) matched carcinogens, by demographic



(b) Selected summary results for chemicals with ratio ≥ 1.5

Chemical	Flags	Max Ratio	Demographics
Phenolphthalein	3	2.25	Hispanic, Asian, Female CBA
Di(2-ethylhexyl) phthalate	2	3.57	Hispanic, Inc: >\$30k
Cadmium	2	1.55	Hispanic, Female CBA
Benzophenone	1	7.74	Asian
Coal tar	1	2.72	Black
1,4-Dichlorobenzene	1	2.70	Black
Asphalt, oxidized	1	2.48	Asian
1-Chloro-4-(trifluoromethyl)benzene	1	1.97	White
Furan	1	1.90	Hispanic
1,2-Propylene oxide	1	1.90	Hispanic
Melamine	1	1.86	Black
Mineral oil - includes paraffin oil	1	1.83	Black
Nickel (II) oxide	1	1.81	Black
C.I. Solvent Yellow 14	1	1.79	Inc: <\$30k
Diethanolamine	1	1.76	Asian
4-Methyl-2-pentanone	1	1.74	Hispanic
Amides, coco, N,N-bis(hydroxyethyl)	1	1.63	Hispanic
Diuron	1	1.60	Hispanic
Arsenic	1	1.60	Female CBA
Ethylene oxide	1	1.58	Hispanic

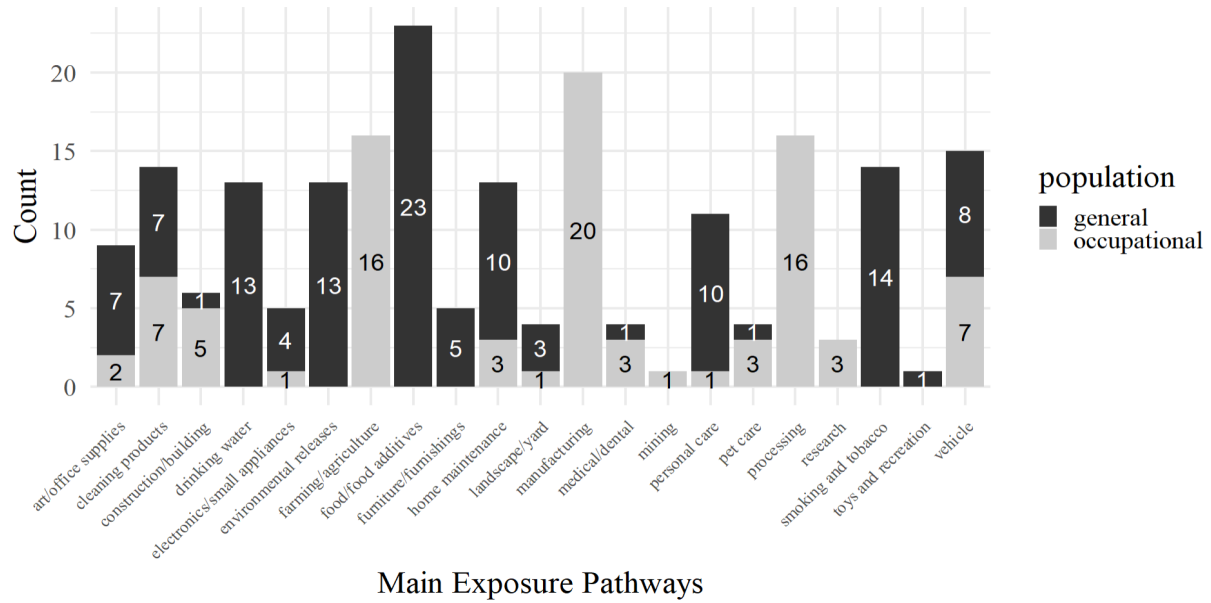
Notes: Panel (a) illustrates the full distribution of chemical purchase prevalence ratios, differentiated by demographic subgroup. This prevalence ratio is our metric of disparities in exposure to individual carcinogens found in consumer products, and is measured by the ratio of a demographic subgroup's *average* propensity for purchasing products with a chemical and the national average propensity for doing so. Each subgroups' median prevalence ratio is marked on the figure with a black line. Panel (b) presents subgroups' purchase prevalence ratios for the selected set of chemicals with a ratio value of 1.5 or greater for at least one demographic subgroup. *Flags* denote the number of subgroups for which the chemical had a prevalence ratio ≥ 1.5 . *Max Ratio* is the chemical's largest prevalence ratio value across subgroups. The full set of purchase prevalence ratio results is presented in Appendix Table A20.

Demographic subgroups: Due to the original paper's data construction, racial subgroups (Asian, Black, White) are not exclusively non-Hispanic ethnicity. All racial/ethnic subgroup assignments are based on the female head of household. Household-income subgroups differentiated by those earning above and below \$30,000 in the year of data collection. Female, CBA denotes whether there is a female of child-bearing age (<45 years old) in the household.

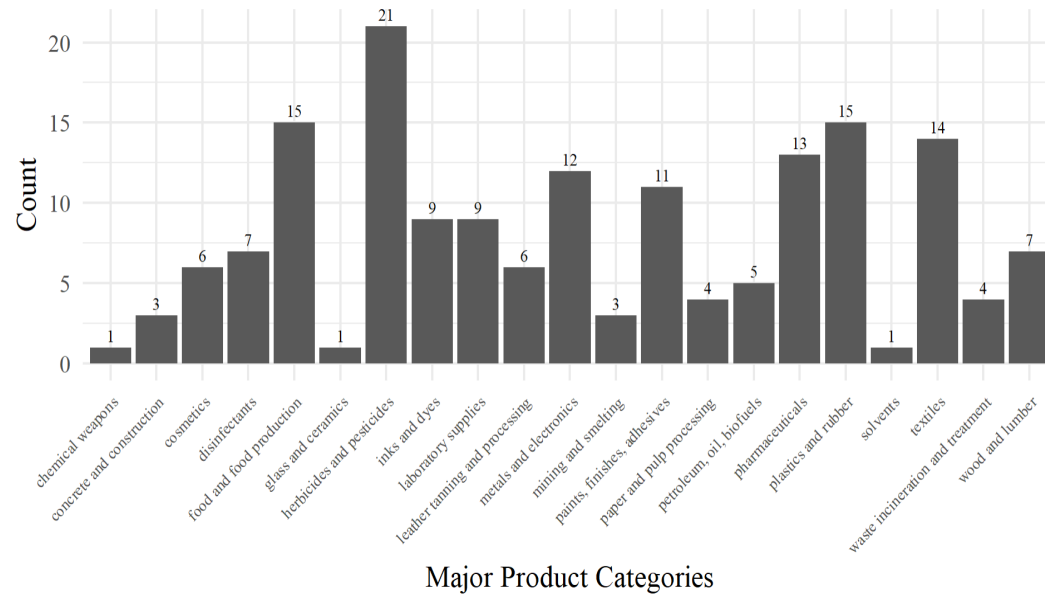
Return to: Section 3.2.3.

Figure 7: Carcinogens with exposure disparities: likely pathways and product categories

(a) Frequency of identified uses (general/occupational) amongst carcinogens with exposure disparities



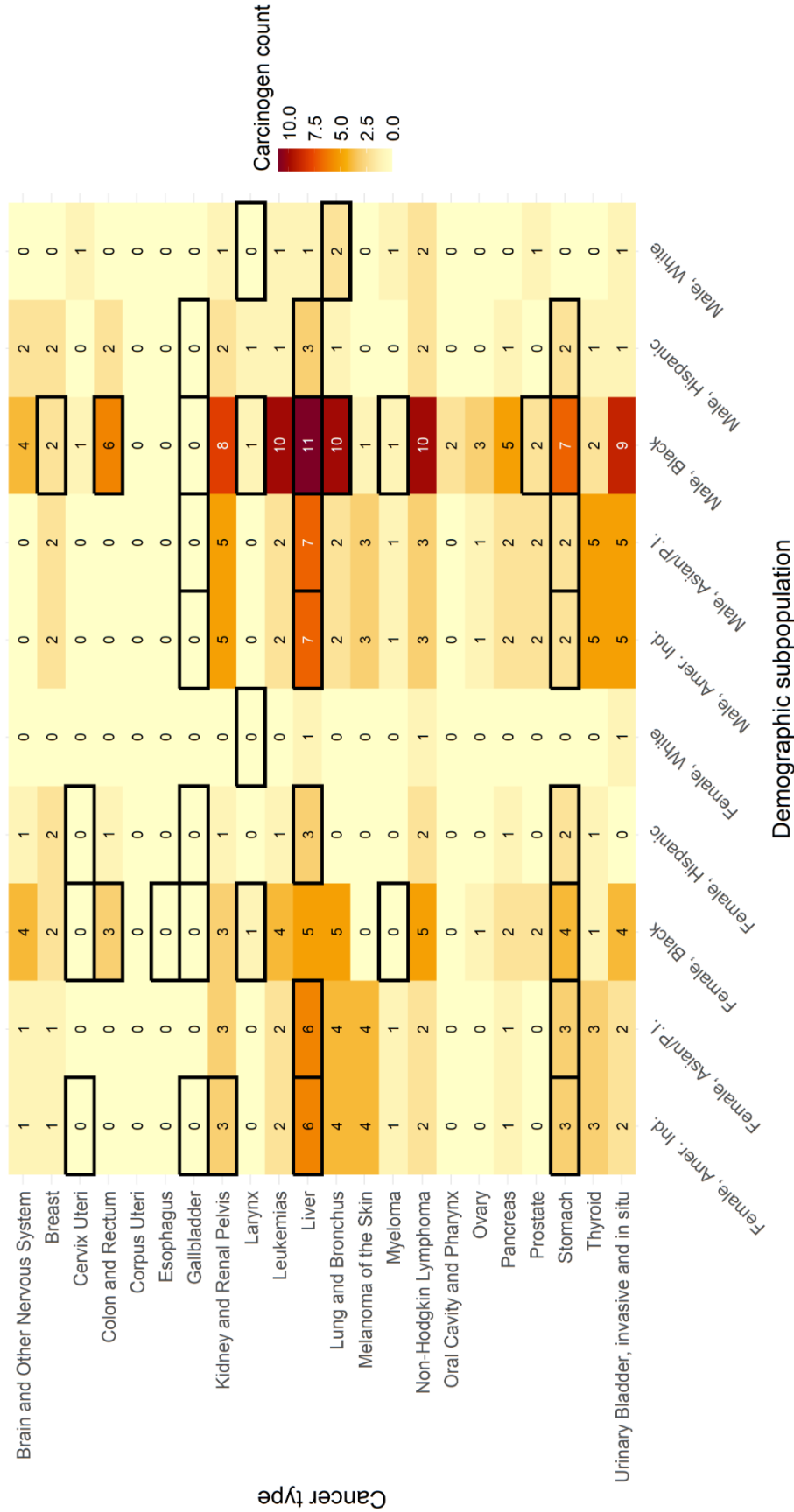
(b) Product category frequency amongst carcinogens with exposure disparities



Notes: Panel (a) illustrates the relative prevalence of particular **exposure pathways** for the set of carcinogens with likely exposure disparities. The figure distinguishes between general population and occupational pathways. Panel (b) presents the count of carcinogens with likely exposure disparities (out of N=28) which were found to be associated with a given **product category**. See Section 2.4 and Section 3.4 for description of exposure pathway information collection and product categorization methods.

Return to: Section 3.4.

Figure 8: Correlation heatmap: carcinogen exposure disparities and elevated cancer incidence.



Notes: Heatmap of carcinogen exposure and associated cancer types. A cell's black bordering denotes an elevated rate of cancer incidence (25% or more above the national average) for a given cancer type and associated subgroup. A cell's color (and its corresponding numeric value inside the cell) denotes the number of carcinogens which are (1) associated with the particular cancer type and (2) present at elevated levels in biomonitoring data for the relevant demographic subgroup**. A darker cell with black bordering therefore jointly indicates a cancer type and demographic subgroup for which we identified overlaps between cancer type and chemical exposure disparities.

**:. We denote a subgroup's exposure as elevated if either the calculated GM ratio or P95 ratio exceed a value of 1.5. Carcinogen counts for American Indian and Asian/Pacific Islander subgroups are jointly based on the "non-Hispanic, Other" race group in the NHANES microdata.

Return to: Section 3.5.

Tables

Table 1: Primary synthesis: carcinogens with evidence of potential exposure disparities

Chemical	NHANES Avg	NHANES P95	HHP	DW	RSEI	NHANES Cycle
Arsenic	✓	✓	✓	✓	✓	2015-2016
1-Naphthylamine	✓	✓			✓	2013-2014
Acrylonitrile**	✓	✓			✓	2015-2016
Furan		✓	✓		✓	2017-2018
1,4-Dichlorobenzene*	✓	✓	✓		✗	2017-2018
2,4-Dichlorophenoxyacetic acid*	✓	✓	✗		✓	2015-2016
Ethylene oxide*	✗	✓	✓		✓	2015-2016
4-Methyl-2-pentanone			✓		✓	2017-2018
Diazinon**		✓			✓	2013-2014
Diethanolamine			✓		✓	
Ethylene thiourea		✓			✓	2007-2008
Nitrofen**	✓	✓				2015-2016
PCBs†	✓				✓	2015-2016
Vinyl chloride**		✓			✓	2015-2016
2-Methylaniline	✓	✗			✓	2013-2014
Acrylamide*	✗	✓			✓	2015-2016
Melamine	✓	✗	✓			2003-2004
Perfluorooctanesulfonic acid	✗	✓			✓	2017-2018
1,2-Propylene oxide**	✗	✗	✓		✓	2015-2016
1-Naphthalenol, 1-(N-methylcarbamate)**	✓	✗	✗		✓	2015-2016
Di(2-ethylhexyl) phthalate**	✗	✗	✓		✓	2017-2018
Naphthalene**	✓	✗	✗		✓	2015-2016
Beta-Hexachlorocyclohexane†	✓					2015-2016
DDT†	✓					2015-2016
2-Naphthylamine	✓	✗				2013-2014
4-(N-Methyl-N-nitrosamino)-1-(3-pyridyl)-1-butanone**	✓	✗				2013-2014
Dimethylarsinic acid	✗	✓				2017-2018
Acrolein**	✓	✗			✗	2015-2016

Notes: Cross-analysis summary of chemical-specific disparities in exposure, based on ratio inclusion threshold value of 1.5. ✓: evidence of potential exposure disparities for at least of the analysis's primary subgroups. ✗: no evidence of potential disparities in exposure. Chemicals listed in first column are parent carcinogens. NHANES results may be based on metabolite or pooled concentrations. **: Carcinogens for which only metabolite biomonitoring results were available. *: Carcinogens for which both direct and metabolite biomonitoring results were available. †: NHANES average analysis based on pooled data.

Column descriptions: NHANES Avg refers to the GM and AM ratio results in Section 3.1.1 and Section 3.1.2. NHANES P95 refers to the P95 ratio results in Section 3.1.3. HHP refers to the household product prevalence ratio results in Section 3.2.3. DW refers to the drinking water concentration ratio results in Section 3.2.2. RSEI refers to the ambient air concentration ratio results in Section 3.2.1. NHANES Cycle notes the biomonitoring data cycle years used to calculate chemical-specific statistics.

Parent Chems: {1-naphthol**: 1-Naphthalenol, 1-(N-methylcarbamate), Naphthalene}; {2-naphthol**: Naphthalene}; {2,4-dichlorophenol**: Nitrofen, 2,4-Dichlorophenoxyacetic acid}; {2,4,5-trichlorophenol**: beta-Hexachlorocyclohexane, Lindane, 1,2,3,4,5,6-Hexachlorocyclohexane, Pentachlorophenol}; {2,5-dichlorophenol**: 1,4-Dichlorobenzene}; {Bis(2-chloroethyl) phosphate**: Tris(2-chloroethyl) phosphate}; {Dimethyl phosphate**: Malathion, Z-Tetrachlorvinphos, Dichlorvos}; {Malathion dicarboxylic acid**: Malathion}; {Mandelic acid**: Ethylbenzene, Styrene}; {Mono-(2-ethyl-5-hydroxyhexyl) phthalate**: Di(2-ethylhexyl) phthalate}; {Mono-(2-ethyl-5-oxohexyl) phthalate**: Di(2-ethylhexyl) phthalate}; {Mono-2-ethyl-5-carboxypentyl phthalate**: Di(2-ethylhexyl) phthalate}; {N-Acetyl-S-(1-cyano-2-hydroxyethyl)-L-cysteine**: Acrylonitrile}; {N-acetyl-S-(2-carbamoyl-ethyl)-L-cysteine**: Acrylamide}; {N-acetyl-S-(2-Carboxyethyl)-L-cysteine**: Acrolein}; {N-acetyl-S-(2-cyanoethyl)-L-cysteine**: Acrylonitrile}; {N-acetyl-S-(2-hydroxy-3-butenyl)-L-cysteine**: 1,3-Butadiene}; {N-acetyl-S-(2-Hydroxyethyl)-L-cysteine**: Acrylonitrile, Vinyl chloride, Ethylene oxide}; {N-acetyl-S-(3-Hydroxypropyl)-L-cysteine**: Acrolein}; {N-acetyl-s-(3-hydroxypropyl-1-methyl)-L-cysteine**: Crotonaldehyde}; {N-Acetyl-S-(n-propyl)-L-cysteine**: 1-Bromopropane}; {NNAL**: NNK}; {Oxypyrimidine**: Diazinon};

Return to: Section 3.3, Section 3.4.

Table 2: National cancer incidence by race/ethnicity and sex

Cancer Type													
Brain and Other Nervous System	5.3	4.1	3.2	3.6	4.7	6.3	7.4	5.7	4.5	4.8	6	8.8	
Breast	129.8	99.3	95.4	125.3	97.2	132.4	1.3	0.6	0.7	1.9	0.8	1.3	
Cervix Uteri	7.5	9.6	6.7	10.1	10.9	7.3							
Colon and Rectum	32.1	37.6	28.5	44.2	31.4	37.8	41.4	47.9	38.4	59.1	44.2	49.8	
Corpus Uteri	26.8	21.5	18.4	23.7	21.9	25.8							
Esophagus	1.8	1.9	1.1	2.5	1.2	1.9	7.8	7.4	3.8	8	5.2	8.8	
Gallbladder	1.3	2.7	1.5	1.9	2.6	1.1	0.8	1.5	1.1	1.3	1.3	0.7	
Kidney and Renal Pelvis	12.0	16	5.1	12.6	12.2	11.1	23.4	28.4	10.8	24.2	21.4	21.8	
Larynx	1.1	1.3	0.3	1.8	0.7	1.5	4.9	5.5	2.2	9.2	5	6.2	
Leukemias	11.0	9.1	6.4	8.9	9.6	11.4	17.8	13.6	10	13.7	13.7	19.1	
Liver	3.4	6.8	6.3	4	6.3	2.4	10.9	16.3	18.3	14.7	17.7	8.2	
Lung and Bronchus	48.5	51.1	28.3	49.8	25.7	57.6	59.6	65.2	45.7	89.8	41.5	77.9	
Melanoma of the Skin	18.4	6.5	1.2	1	4.6	21.5	28.7	9.7	1.5	1.1	5	32.7	
Myeloma	5.8	5.6	3.1	11.8	5.6	4.6	8.6	7.2	4.8	15.8	7.9	7.4	
Non-Hodgkin Lymphoma	15.4	13.3	11	12.1	15.6	16.9	22.3	16.3	15.9	17.4	20.5	24.4	
Oral Cavity and Pharynx	6.6	5.7	5.3	5.3	4.4	6.8	18.0	15	11.5	15.3	10.9	18.7	
Ovary	10.1	10.9	9.4	9.5	10.7	12.5							
Pancreas	11.9	10.1	8.7	14.5	10.6	10.7	15.3	11.8	10.2	17.4	12.4	14.3	
Prostate							113.1	93.5	69.3	210.5	109.1	128.1	
Stomach	4.7	6.2	8.3	8.1	8.2	3.6	8.3	10.8	14.4	14.9	13.1	7.9	
Thyroid	19.0	14.8	19.6	11.4	18.6	18.7	6.9	4.5	6.3	3.3	5.1	6.8	
Urinary Bladder, invasive and in situ	8.1	5.2	3.7	6.7	5.3	10	32.5	20.5	15.3	19.7	20.3	40.3	
Dem subgroup:													
Sex:	Female			Female			Female			Female			
Race/Ethnicity:	All			Amer Ind			Asian/PI			Black			
	Male			Amer Ind			Asian/PI			Black			
	Male			Male			Male			Male			
	White			White			White			White			

Notes: Table presents national cancer incidence rates (per 100,000 population). Data obtained from CDC, 2021b in October 2024, and are based on incidence during the years 2017-2021. Cancer sites presented in this table include all leading and invasive cancer types. All statistics are age-adjusted rates, based on 2000 U.S. standard population. Race/ethnicity subgroups are: non-Hispanic American Indian and Alaska Native, non-Hispanic Asian and Pacific Islander, non-Hispanic Black, Hispanic, non-Hispanic White. **Bold font** designates subgroup-specific incidence rates that are 25% or more above the corresponding national average. Underlined values denote concentration ratios subgroup-specific incidence rates that are 50% or more above the corresponding national average.

Return to: Section 3.5.

Table 3: Exposure pathways and cancer types for chemicals with exposure disparities

Chemical	Major Prod. Categories	Exp. route	Cancers: animal	Cancers: human	General exposure	Occupational exposure
Arsenic	Mining And Smelting; Herbicides And Pesticides; Metals And Electronics	Ingestion	Leukemia; Lung; Lymphoma; Stomach	Bladder; Skin; Lung; Stomach; Digestive; Liver; Kidney; Prostate; Lymphatic; Hemtopoietic	Electronics/ Small Appliances; Cleaning Products; Art/ Office Supplies; Home Maintenance; Vehicle; Food/ Food Additives; Smoking And Tobacco; Drinking Water; Environmental Releases	Manufacturing; Processing; Vehicle; Farming/ Agriculture; Medical/ Dental
1-Naphthylamine	Plastics And Rubber; Textiles; Laboratory Supplies; Herbicides And Pesticides; Pharmaceuticals	Inhalation; Dermal		Bladder	Food/ Food Additives; Smoking And Tobacco	Manufacturing; Processing
Acrylonitrile	Paints, Finishes, Adhesives; Plastics And Rubber; Inks And Dyes; Pharmaceuticals; Textiles	Inhalation; Ingestion; Dermal	Brain; Breast; CNS; Prostate	Bladder; Lung; Lymphatic; Stomach; Colon; Respiratory	Art/ Office Supplies; Home Maintenance; Smoking And Tobacco; Furniture/ Furnishings; Food/ Food Additives; Vehicle	Manufacturing; Processing; Home Maintenance
Furan	Paints, Finishes, Adhesives; Herbicides And Pesticides; Pharmaceuticals; Wood And Lumber; Petroleum, Oil, Biofuels; Food And Food Production	Inhalation; Dermal	Liver; Bile-Duct; Adrenal; Leukemia		Smoking And Tobacco; Food/ Food Additives; Environmental Releases; Drinking Water	Manufacturing
1,4-Dichlorobenzene	Food And Food Production; Herbicides And Pesticides; Disinfectants; Inks And Dyes; Pharmaceuticals; Plastics And Rubber; Metals And Electronics; Leather Tanning And Processing; Textiles; Laboratory Supplies	Inhalation; Ingestion; Dermal	Stomach; Kidney; Leukemia; Adrenal; Liver	Leukemia	Cleaning Products; Personal Care; Vehicle; Food/ Food Additives; Drinking Water; Environmental Releases	Vehicle; Manufacturing; Processing; Farming/ Agriculture

Continued, next page...

Table 3: Exposure pathways and cancer types for chemicals with exposure disparities

Chemical	Major Prod. Categories	Exp. route	Cancers: animal	Cancers: human	General exposure	Occupational exposure
2,4- Dichlorophenoxyacetic Acid	Herbicides And Pesticides; Food And Food Production	Ingestion; Inhalation		Non-Hodgkin's Lymphoma; Sarcoma; Colorectal; Gastrointestinal; Breast; Brain	Landscape/ Yard	Farming/ Agriculture; Cleaning Products; Manufacturing; Processing
Ethylene Oxide	Laboratory Supplies; Herbicides And Pesticides; Food And Food Production; Disinfectants; Cosmetics	Inhalation; Ingestion	Leukemia; Brain; Breast; Lung; Lymphoma; Uterus; Mesothelium	Leukemia; Stomach; Lymphatic; Breast; Myeloma	Environmental Releases; Smoking And Tobacco; Personal Care; Home Maintenance; Food/ Food Additives; Vehicle	Medical/ Dental; Manufacturing; Farming/ Agriculture
4-Methyl-2-Pentanone	Paints, Finishes, Adhesives; Pharmaceuticals; Plastics And Rubber; Textiles; Inks And Dyes; Solvents; Cosmetics; Wood And Lumber; Metals And Electronics; Food And Food Production	Inhalation; Dermal; Ingestion	Leukemia; Renal; Kidney; Liver	Bladder; Lung	Art/ Office Supplies; Food/ Food Additives; Cleaning Products; Furniture/ Furnishings; Electronics/ Small Appliances; Personal Care; Home Maintenance; Pet Care	Art/ Office Supplies; Cleaning Products; Vehicle; Personal Care
Diazinon	Herbicides And Pesticides	Dermal	Leukemia; Lymphoma	Lung; Leukemia; Lymphoma	Food/ Food Additives; Drinking Water; Environmental Releases; Landscape/ Yard	Farming/ Agriculture; Pet Care
Diethanolamine	Disinfectants; Concrete And Construction; Mining And Smelting; Herbicides And Pesticides; Textiles; Pharmaceuticals; Cosmetics; Inks And Dyes; Paper And Pulp Processing; Petroleum, Oil, Biofuels; Paints, Finishes, Adhesives; Plastics And Rubber; Metals And Electronics	Dermal; Inhalation	Liver; Renal	Stomach; Respiratory; Liver; Pancreas; Prostate; Leukemia	Cleaning Products; Food/ Food Additives; Art/ Office Supplies; Personal Care; Electronics/ Small Appliances	Cleaning Products; Construction/ Building; Manufacturing; Processing; Vehicle; Electronics/ Small Appliances

Continued, next page...

Table 3: Exposure pathways and cancer types for chemicals with exposure disparities

Chemical	Major Prod. Categories	Exp. route	Cancers: animal	Cancers: human	General exposure	Occupational exposure
Ethylene Thiourea	Plastics And Rubber; Paints, Finishes, Adhesives; Disinfectants; Textiles; Metals And Electronics; Waste Incineration And Treatment; Herbicides And Pesticides	Inhalation; Ingestion; Dermal	Liver; Thyroid; Leukemia	Thyroid	Food/ Food Additives; Drinking Water; Environmental Releases	Manufacturing; Processing; Vehicle; Farming/ Agriculture
Nitrofen	Herbicides And Pesticides; Food And Food Production	Inhalation; Ingestion; Dermal	Liver; Pancreas; Bile-duct		Food/ Food Additives	Farming/ Agriculture; Manufacturing
Polychlorinated Biphenyls (PCBs)	Metals And Electronics; Paints, Finishes, Adhesives; Herbicides And Pesticides; Inks And Dyes; Food And Food Production; Laboratory Supplies; Plastics And Rubber; Waste Incineration And Treatment	Inhalation	Liver; Bile-duct; Lung; Oral cavity; Pancreas; Spleen; Uterus	Melanoma; Liver; Bile-Duct; Gastro-Intestinal; Lymphoma; Breast	Environmental Releases; Drinking Water; Food/ Food Additives	Farming/ Agriculture; Manufacturing; Research
Vinyl Chloride	Plastics And Rubber; Food And Food Production; Metals And Electronics; Wood And Lumber; Leather Tanning And Processing	Ingestion; Inhalation; Dermal	Breast; Kidney; Liver; Bile-duct; Lung; Nasal cavity; Skin; Stomach	Liver; Brain; Lung; Lymphoma; Leukemia	Smoking And Tobacco; Environmental Releases; Drinking Water; Food/ Food Additives; Vehicle; Home Maintenance	Manufacturing; Processing; Vehicle; Construction/ Building
2-Methylaniline	Textiles; Pharmaceuticals; Herbicides And Pesticides	Dermal; Inhalation; Ingestion		Bladder	Personal Care; Smoking And Tobacco	Manufacturing; Research
Acrylamide	Waste Incineration And Treatment; Inks And Dyes; Pharmaceuticals; Cosmetics; Paper And Pulp Processing; Textiles; Mining And Smelting; Petroleum, Oil, Biofuels; Food And Food Production; Concrete And Construction; Paints, Finishes, Adhesives	Ingestion; Dermal; Inhalation	Breast; Thyroid; Testicular; Adrenal; Oral; Uterus Lung	Pancreatic; Breast	Food/ Food Additives; Vehicle; Home Maintenance; Smoking And Tobacco; Landscape/ Yard; Personal Care; Drinking Water	Mining; Processing; Manufacturing; Farming/ Agriculture; Art/ Office Supplies; Home Maintenance

Continued, next page...

Table 3: Exposure pathways and cancer types for chemicals with exposure disparities

Chemical	Major Prod. Categories	Exp. route	Cancers: animal	Cancers: human	General exposure	Occupational exposure
Melamine	Laboratory Supplies; Paints, Finishes, Adhesives; Wood And Lumber; Leather Tanning And Processing; Metals And Electronics; Textiles; Paper And Pulp Processing; Plastics And Rubber	Ingestion; Inhalation; Dermal	Bladder	Bladder; Kidney	Furniture/ Furnishings; Food/ Food Additives	Cleaning Products; Construction/ Building; Manufacturing; Processing
Perfluorooctanesul- fonic Acid	Paints, Finishes, Adhesives; Metals And Electronics; Food And Food Production; Textiles; Paper And Pulp Processing; Petroleum, Oil, Biofuels; Herbicides And Pesticides	Ingestion; Inhalation; Dermal	Liver; Thyroid; Testicular; Breast; Pancreas	Bladder; Testicular; Kidney; Thyroid; Prostate; Breast; Ovarian	Food/ Food Additives; Environmental Releases; Home Maintenance; Cleaning Products; Furniture/ Furnishings; Drinking Water	Farming/ Agriculture; Manufacturing
1,2-Propylene Oxide	Food And Food Production; Pharmaceuticals; Disinfectants; Textiles; Paints, Finishes, Adhesives; Herbicides And Pesticides; Food And Food Production; Plastics And Rubber; Concrete And Construction	Dermal; Inhalation	Nasal/Sinus; Adrenal; Stomach; Mammary	Lymphatic	Personal Care; Food/ Food Additives; Vehicle; Home Maintenance; Cleaning Products	Farming/ Agriculture; Research; Processing; Cleaning Products
1-Naphthalenol	Inks And Dyes; Cosmetics; Pharmaceuticals; Plastics And Rubber; Leather Tanning And Processing; Metals And Electronics; Herbicides And Pesticides	Inhalation; Dermal	Nasal/Sinus; Lung	Bladder; Lung; Breast	Personal Care; Art/ Office Supplies; Food/ Food Additives; Smoking And Tobacco	Farming/ Agriculture; Processing
Di(2-Ethylhexyl) Phthalate	Plastics And Rubber; Herbicides And Pesticides; Inks And Dyes; Metals And Electronics; Laboratory Supplies; Food And Food Production; Cosmetics; Paints, Finishes, Adhesives	Ingestion; Inhalation; Medical Procedure	Liver; Pancreas; Testicular	Breast; Ovarian; Thyroid; Testicular	Furniture/ Furnishings; Vehicle; Home Maintenance; Personal Care; Art/ Office Supplies; Construction/ Building; Food / Food Additives; Toys And Recreation; Environmental Releases; Drinking Water	Medical/ Dental; Construction/ Building; Processing

Continued, next page...

Table 3: Exposure pathways and cancer types for chemicals with exposure disparities

Chemical	Major Prod. Categories	Exp. route	Cancers: animal	Cancers: human	General exposure	Occupational exposure
Naphthalene	Herbicides And Pesticides; Leather Tanning And Processing; Wood And Lumber; Textiles; Disinfectants; Petroleum, Oil, Biofuels; Plastics And Rubber; Pharmaceuticals	Inhalation; Ingestion	Lung; Nasal/Sinus; Brain	Larynx; Colorectal	Medical/ Dental; Home Maintenance; Smoking And Tobacco; Environmental Releases; Electronics/ Small Appliances; Food/ Food Additives	Vehicle; Cleaning Products; Construction/ Building; Home Maintenance; Manufacturing; Processing
Beta- Hexachlorocyclohexane	Laboratory Supplies; Herbicides And Pesticides	Inhalation; Dermal	Liver	Leukemia; Melanoma	Food/ Food Additives; Environmental Releases; Drinking Water	Farming/ Agriculture
DDT	Laboratory Supplies; Pharmaceuticals; Food And Food Production; Herbicides And Pesticides	Ingestion	Liver; Lung; Lymphoma; Thyroid	Breast; Non-Hodgkin's Lymphoma; Testicular; Liver; Bile-duct	Food/ Food Additives; Drinking Water; Smoking And Tobacco; Personal Care	Farming/ Agriculture; Manufacturing; Pet Care
2-Naphthylamine	Plastics And Rubber; Textiles	Dermal; Inhalation; Ingestion	Liver; Bile-duct; Bladder	Bladder	Cleaning Products; Smoking And Tobacco; Environmental Releases	Manufacturing
4-(N-Methyl-N-Nitrosamino)-1-(3-Pyridyl)-1-Butanone	Laboratory Supplies	Inhalation	Nasal/Sinus; Lung; Liver; Bile-duct; Adrenal; Leukemia; Lymphoma; Larynx; Pancreas	Lung	Smoking And Tobacco	
Dimethylarsinic Acid	Herbicides And Pesticides; Pharmaceuticals; Chemical Weapons; Wood And Lumber	Inhalation	Lung; Bladder; Kidney; Liver; Thyroid			Landscape/ Yard; Farming/ Agriculture; Pet Care; Processing

Continued, next page...

Table 3: Exposure pathways and cancer types for chemicals with exposure disparities

Chemical	Major Prod. Categories	Exp. route	Cancers: animal	Cancers: human	General exposure	Occupational exposure
Acrolein	Herbicides And Pesticides; Wood And Lumber; Leather Tanning And Processing; Plastics And Rubber; Disinfectants; Glass And Ceramics; Textiles; Inks And Dyes; Food And Food Production; Metals And Electronics; Waste Incineration And Treatment	Inhalation; Ingestion	Nasal/Sinus; Lung	Lung; Colorectal	Art/ Office Supplies; Smoking And Tobacco; Food/ Food Additives; Drinking Water	Farming/ Agriculture; Processing; Manufacturing; Cleaning Products

Notes: Compiled information on exposure sources/types, product categories, and associated cancer types for the complete list of 28 carcinogens which presented the strongest evidence of exposure differences across demographic subgroups. Column (1) is the chemical name. Column (2) is the list of major product categories in which the chemical is used. Column (3) lists relevant exposure routes for the carcinogen. Column (4) lists cancer types linked with the chemical in animal testing. Column (5) lists cancer types linked with the chemical in human/epidemiological studies. Column (6) lists general population exposure sites. Column (7) lists occupational exposure sites. **Bold** text in columns (4) and (5) indicate that evidence supporting an animal-based or human-based link between carcinogen exposure and the cancer site is included in NTP's Report on Carcinogens dashboard (ROC, 2022) or the IARC's list of classifications by cancer sites with sufficient or limited evidence in humans (IARC, 2024). Other evidence sources include: CompTox (Williams et al., 2017), PubChem (Kim et al., 2023), and Haz-Map (Brown, 2024). See Section 3.4 and Appendix C for further details of literature and data review methods.

Return to: Section 3.4, Section 3.5.

Appendices for “Identifying disparities in U.S. exposure to ubiquitous carcinogens”

Adam Theising, Tina Bardot, Ann Wolverton

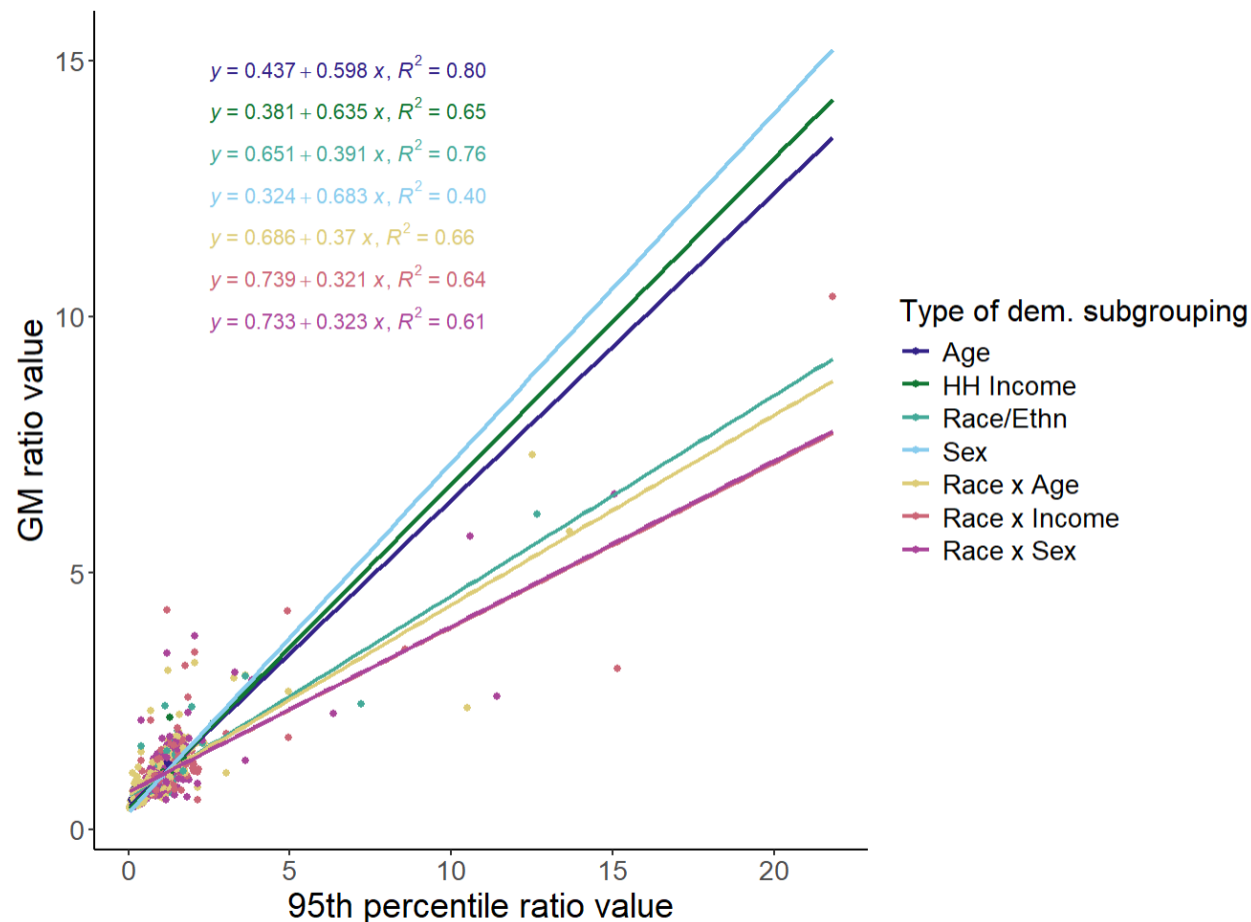
December 2024

Appendices

A	Supplemental figures and tables	1
B	Exposure pathways for known carcinogens with scarce biomonitoring	61
C	Information on data construction and exposure literature review	80
D	Summary of carcinogen biomonitoring by states	85
E	Full list of candidate carcinogens	90

A Supplemental figures and tables

Figure A1: Correlations between NHANES GM ratio and P95 ratio values

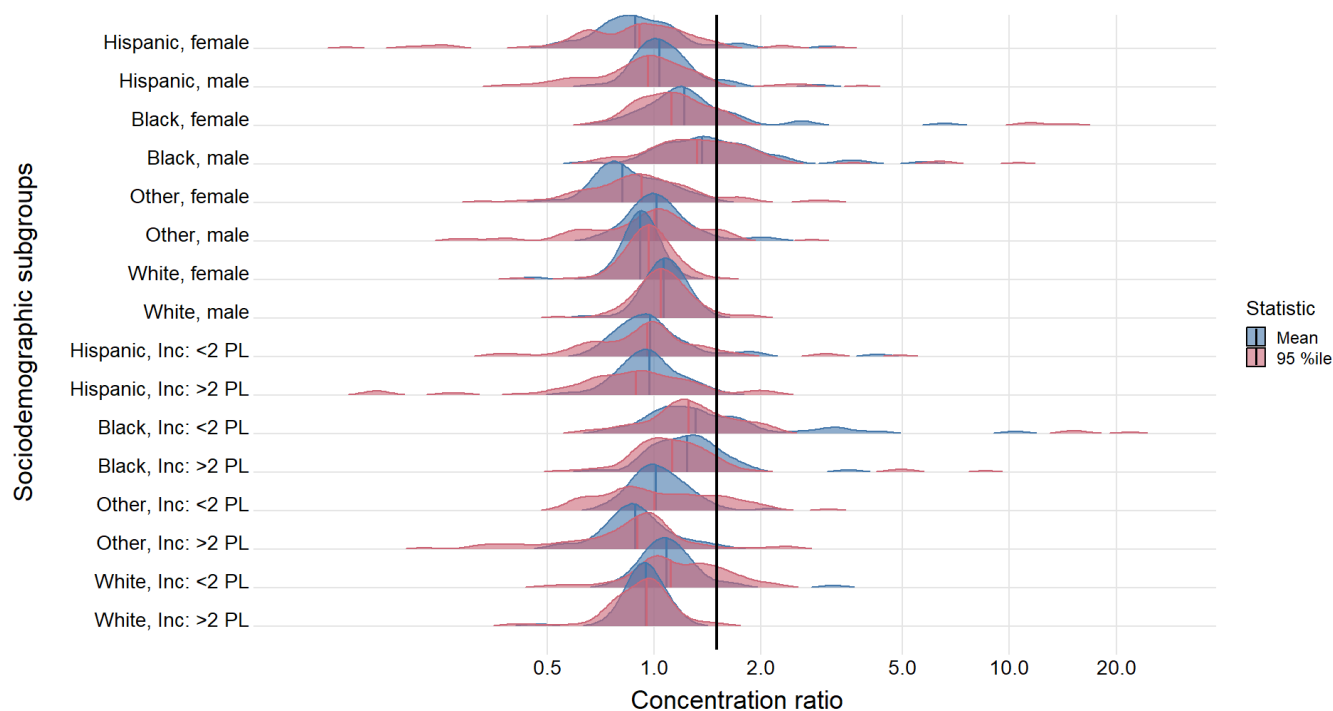


Notes: Correlation between calculated NHANES geometric mean ratio values and 95th percentile ratio values for different demographic subgroups. Each observation in these correlation regressions is a **complete** quartet of characteristics {Demographic subgroup, Chemical, GM ratio value, 95th %ile value}. If a subgroup-specific geometric mean concentration could not be calculated due to detection limits, the observation cannot be used in a correlation regression. Demographic subgroups include age (<19, 20+), HH income (<2x poverty line, >2x poverty line), Race/Ethnicity (Hispanic, non-Hispanic Black, non-Hispanic White, Other), Sex (Female, Male), and their interactions (denoted by $(\cdot) \times (\cdot)$). Correlation regressions fit on 94 observations for Age subgroups, 95 observations for HH Income subgroups, 190 observations for Race/Ethnicity subgroups, 95 observations for Sex subgroups, 371 observations for Race $\times$ Age subgroups, 378 observations for Race $\times$ Income subgroups, and 374 observations for Race $\times$ Sex subgroups.

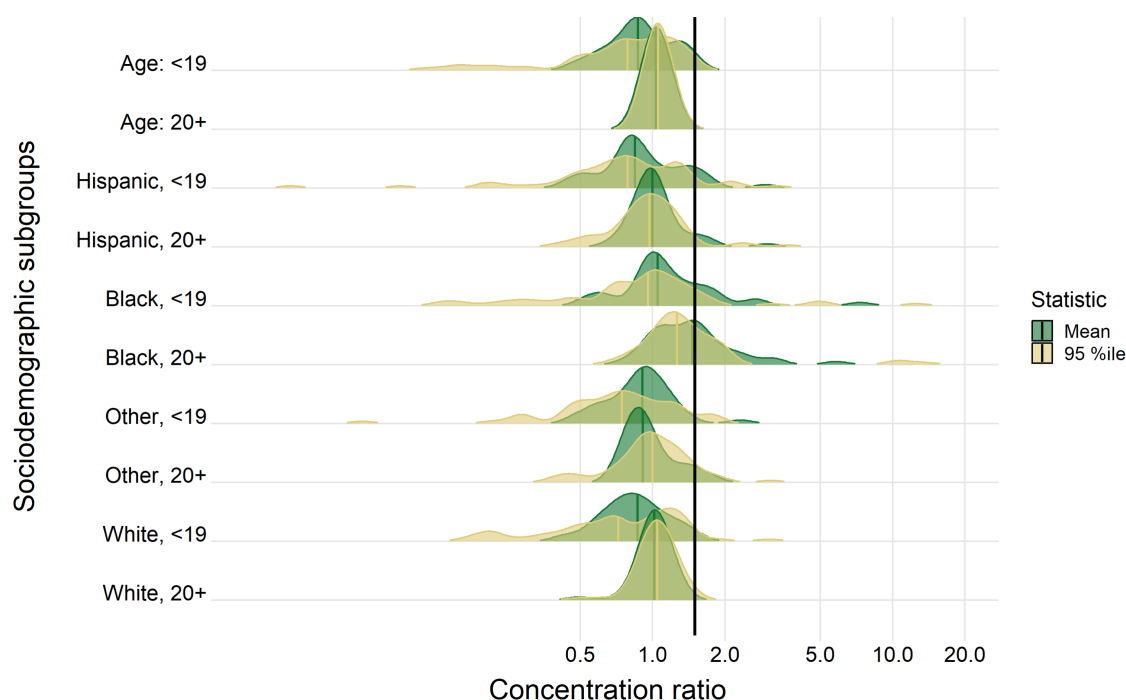
Return to: Section 3.1.3.

Figure A2: Cross-chemical summary of demographic heterogeneity analysis using individual-level NHANES data

(a) Distribution of geometric mean and 95th %ile ratios across (N = 39, 62) carcinogens, by demographic interactions



(b) Distribution of geometric mean and 95th %ile ratios across (N = 39, 62) carcinogens, by age-differentiated demographics

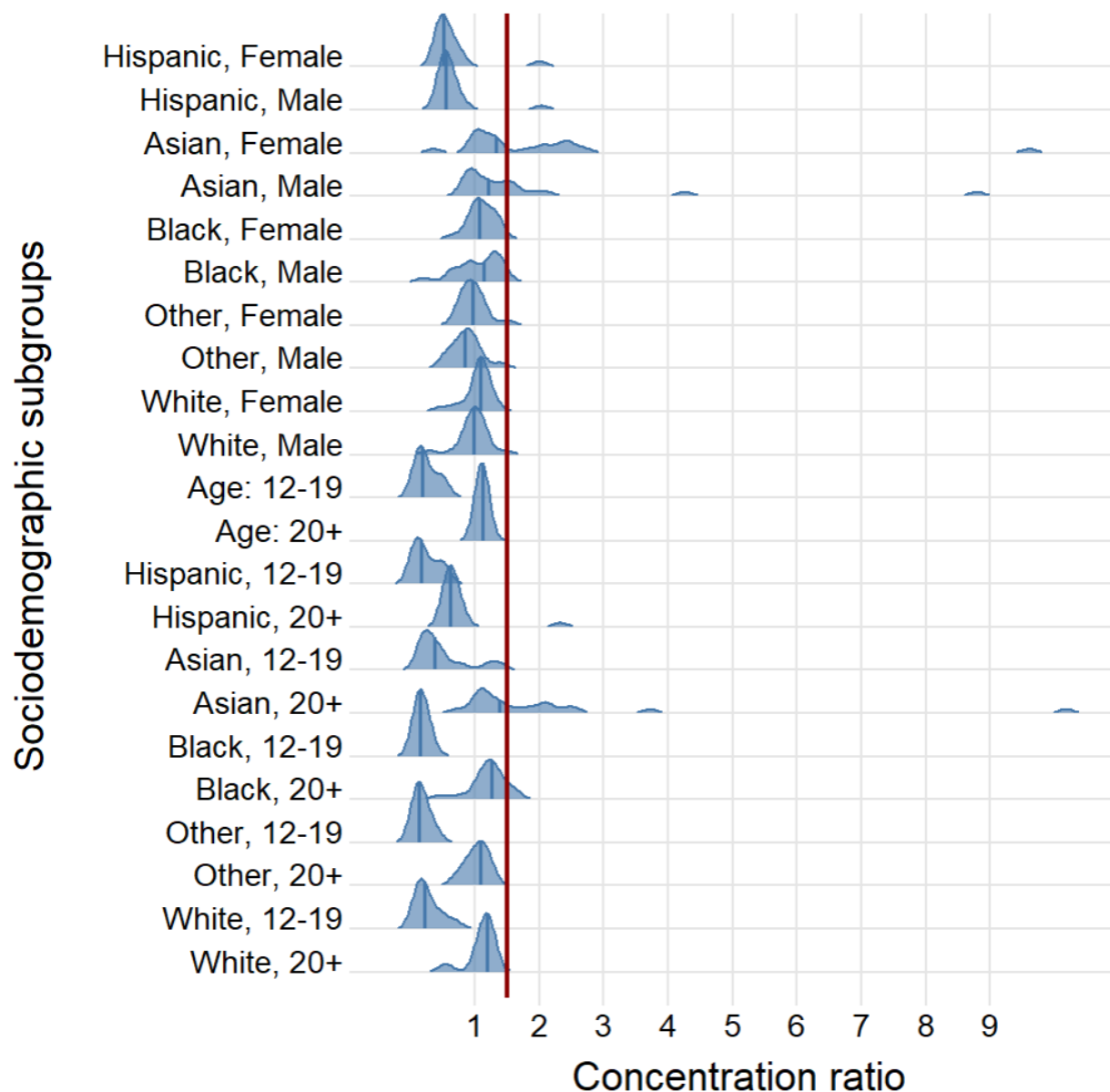


Notes: Panel (a) illustrates the full distribution of geometric mean (N = 39) and 95th percentile (N = 62) concentration ratios across matched carcinogens, differentiated by demographic subgroup. Subgroups in this panel include interactions of race/ethnicity and sex, and interactions of race/ethnicity and household income. The full, chemical-level results for these demographic subgroups are presented in Appendix Table A5 and Table A11. Panel (b) illustrates the full distribution of geometric mean (N = 39) and 95th percentile (N = 62) concentration ratios across matched carcinogens, presented by age subgroup, including age subgroups differentiated by race/ethnicity. The full, chemical-level results for these demographic subgroups are presented in Appendix Table A6 and Table A12. In both panels, subgroups' median GMR and PR₉₅ are marked on their respective density plot.

Demographic subgroups: Household income below 2 times the poverty line, Household income above 2 times the poverty line, Hispanic, non-Hispanic Black, non-Hispanic White, Other race including multi-racial, Female sex, Male sex.

Return to: Section 3.1.

Figure A3: Distribution of arithmetic mean concentration ratios across ($N = 22$) carcinogens for *pooled* NHANES samples, by demographic subgroup

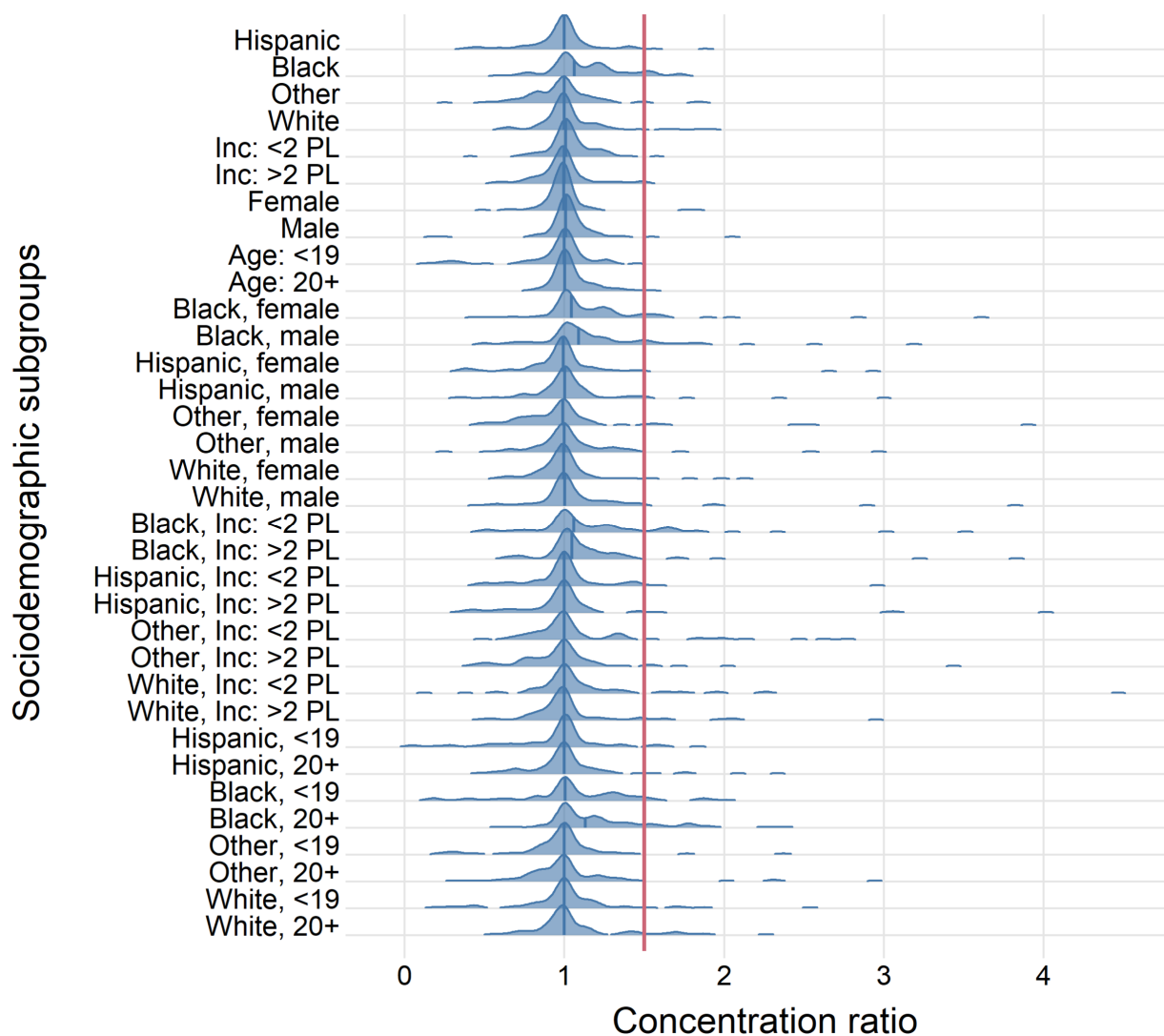


Notes: This figure illustrates the full distribution of arithmetic mean concentration ratios across the 22 matched carcinogens from pooled NHANES samples, differentiated by demographic subgroup. Subgroups in this figure include interactions of race/ethnicity and sex, and interactions of race/ethnicity and age. Each subgroup's median arithmetic mean ratio is marked on their respective density plot. The full set of arithmetic mean ratio results for these demographic subgroups are presented in Appendix Table A8 and Table A9.

Demographic subgroups: Hispanic, non-Hispanic Asian, non-Hispanic Black, non-Hispanic White, Other race including multi-racial, Female sex, Male sex.

Return to: Section 3.1.2.

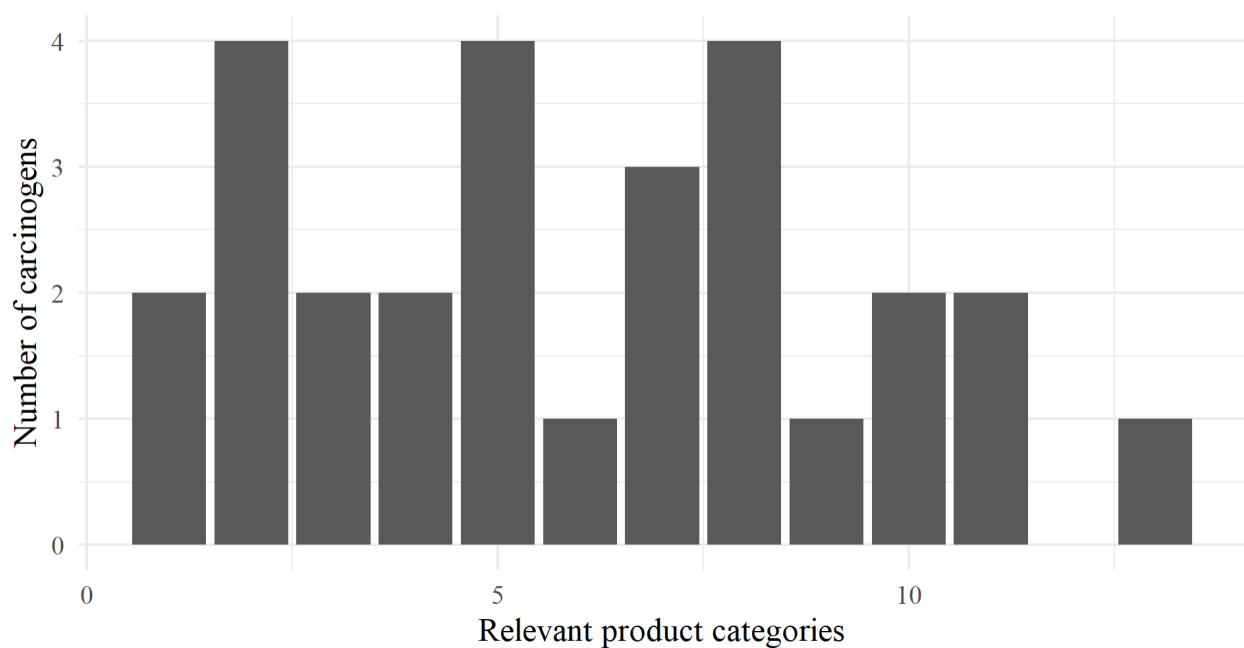
Figure A4: Distribution of sample detection ratios across ($N = 82$) carcinogens for individual NHANES samples, by demographic subgroup



Notes: This figure illustrates the full distribution of sample detection ratios across the 82 matched carcinogens, differentiated by demographic subgroup. The sample detection ratio for a given chemical and subgroup is given as the subgroup's survey-weighted fraction of samples with concentrations above the detection limit divided by the total population's survey-weighted fraction of samples with concentrations above the detection limit. Implicitly, for a chemical to be included in this analysis, at least one sample in the individual-level NHANES data must be detectable. The full set of sample detection ratio results for these demographic subgroups are presented in Appendix Table A13, Table A14, and Table A15. Each subgroup's median sample detection ratio is marked on their respective density plot.

Demographic subgroups: Household income below 2 times the poverty line, Household income above 2 times the poverty line, Hispanic, non-Hispanic Black, non-Hispanic White, Other race including multi-racial, Female sex, Male sex.
Return to: Section 3.1.4.

Figure A5: Exposure sourcing: number of identified product categories in which each chemical is used



Notes: Histogram of the count of associated product categories for each of the ($N = 28$) carcinogens found to have likely exposure disparities. For example, two carcinogens are associated with a single product category, four carcinogens are associated with two different product categories, ..., and one carcinogen is associated with thirteen different product categories. See Section 3.4 for description of exposure pathway information collection and product categorization methods.

Return to: Section 2.4, Section 3.4.

Table A1: Summary of carcinogen list sourcing, classification, and data coverage

Panel A: Source List	(1) All	(2) Cand.	(3a) Dir. BM	(3b) Metab. BM	(4) RSEI	(5) DW	(6) HHP	(7) Any Data
IARC	41	30	5	4	6	2	5	12
ROC	47	28	1	0	5	1	1	7
Prop65	173	141	12	5	38	14	4	45
SVHC	35	24	1	0	0	0	0	1
IARC & ROC	5	5	0	0	0	0	1	1
IARC & Prop65	161	141	16	5	36	10	12	48
IARC & SVHC	0	0	0	0	0	0	0	0
ROC & Prop65	49	45	2	1	9	4	2	12
ROC & SVHC	14	14	0	0	0	0	0	0
Prop65 & SVHC	1	1	0	0	1	0	0	1
ROC, Prop65 & SVHC	2	2	0	0	0	0	0	0
IARC, ROC & Prop65	164	156	38	14	67	30	14	79
IARC, Prop65 & SVHC	7	7	0	0	2	0	0	2
IARC, ROC & SVHC	0	0	0	0	0	0	0	0
All Lists	28	28	10	3	22	6	5	22
Total	727	622	85	32	186	67	44	230

Panel B: Classification	(1) All	(2) Cand.	(3) Dir. BM	(4) Metab. BM	(5) RSEI	(6) DW	(7) HHP	(8) Any Data
Known Carcinogen	112	96	16	6	22	18	9	30
Probable Carcinogen	234	218	37	16	90	25	16	103
Possible Carcinogen	172	142	19	5	35	10	15	50
Other	209	166	13	5	39	14	4	47
Total	727	622	85	32	186	67	44	230

Notes: Summary of carcinogen list sources (Panel A) and carcinogen classification (Panel B) for subsets of the 727 listed carcinogens and the 622 candidate carcinogens scoped into the analysis. All numeric values are applicable carcinogens counts. Column (1): all potential carcinogens. Column (2): all carcinogen candidates found to be ubiquitous. Column (3): set of carcinogens directly biomonitoried by NHANES. Column (4): carcinogens indirectly biomonitoried by metabolite. Column (5): carcinogens whose industrial releases are monitored in TRI/RSEI. Column (6) carcinogens monitored in drinking water by at least one PWS. Column (7): carcinogens matched as ingredients in purchased products for household product analysis. Column (8): set of carcinogens for which data was available (and detectable) in at least one analysis.

Return to: Section 2.1.2.

Table A2: Summary of carcinogen biomonitoring detection in NHANES

Chemical	GM	AM	P95	Detect
1,1,1,2-Tetrachloroethane				
1,1,1-Trichloroethane				T
1,1,2,2-Tetrachloroethane				T
1,1,2-Trichloroethane				
1,1-Dichloroethane				
1,1-Dichloroethylene				
1,2,3,4,5,6-Hexachlorocyclohexane			T	T
1,2,3-Trichloropropane				
1,2-Dibromo-3-chloropropane				
1,2-Dibromoethane				
1,2-Dichloropropane				
1,2-Propylene oxide	T		T	T
1,3-Butadiene	T		T	T
1,4-Dichlorobenzene	T		T	T
1,4-Dioxane				
1-Bromopropane	T		T	T
1-Naphthalenol, 1-(N-methylcarbamate)	T		T	T
1-Naphthylamine	T		T	T
2,3,4,7,8-Pentachlorodibenzofuran		T		T
2,3,7,8-Tetrachlorodibenzo-p-dioxin				T
2,4,6-Trichlorophenol			T	T
2,4-Dichlorophenoxyacetic acid	T		T	T
2,6-Dimethylaniline	T		T	T
2-Amino-1-methyl-6-phenylimidazo [4,5-b]pyridine			T	T
2-Amino-3-methyl-3H-imidazo[4,5-f]quinoline				T
2-Amino-alpha-carboline			T	T
2-Anisidine	T		T	T
2-Methylaniline	T		T	T
2-Naphthylamine	T		T	T
2-Phenylphenol			T	T
4,4'-Diaminobiphenyl methane			T	T
4-(N-Methyl-N-nitrosamino)-1-(3-pyridyl)-1-butanone	T		T	T
4-Biphenylamine	T		T	T
4-Methyl-2-pentanone				T
Acetochlor				T
Acrolein	T		T	T
Acrylamide	T		T	T
Acrylonitrile	T		T	T
Alachlor				T
Arsenic	T		T	T
Arsenic acid			T	T
Benzene			T	T
Beryllium				
Bromodichloromethane			T	T
Bromoform			T	T

Continued, next page...

Table A2: Summary of carcinogen biomonitoring detection in NHANES

Chemical	GM	AM	P95	Detect
Cadmium	T		T	T
Carbon tetrachloride				T
Chloroethane				T
Chloroform			T	T
Cobalt	T		T	T
Crotonaldehyde	T		T	T
Cumene				T
DDT		T		T
Di(2-ethylhexyl) phthalate	T		T	T
Diazinon			T	T
Dichloromethane				T
Dichlorvos	T		T	T
Dieldrin			T	T
Diisononyl phthalate			T	T
Dimethylarsinic acid	T		T	T
Ethylbenzene	T		T	T
Ethylene oxide	T		T	T
Ethylene thiourea			T	T
Formaldehyde	T		T	T
Furan			T	T
Glu-P-1				T
Glu-P-2				T
Heptachlor epoxide B			T	T
Hexachlorobenzene				
Hexachloroethane				
Lead	T		T	T
Lindane				T
Malathion	T		T	T
MeA-alpha-C			T	T
Melamine	T		T	T
Methylarsonic acid			T	T
Mirex				T
N'-Nitrosornicotine			T	T
N,N-Dimethylformamide	T		T	T
N-Nitroso-N-methylethylamine				T
N-Nitrosodiethylamine				T
N-Nitrosomorpholine				T
N-Nitrosopiperidine				T
N-Nitrosopyrrolidine				T
Naphthalene	T		T	T
Nickel	T		T	T
Nitrobenzene				
Nitrofen	T		T	T
Nitromethane	T		T	T
Pentachlorophenol			T	T

Continued, next page...

Table A2: Summary of carcinogen biomonitoring detection in NHANES

Chemical	GM	AM	P95	Detect
Perfluorooctanesulfonic acid	T		T	T
Perfluorooctanoic acid	T		T	T
Polybrominated biphenyls (PBB)		T		T
Polychlorinated biphenyls		T		T
Propoxur				
Styrene	T		T	T
Tetrachloroethylene			T	T
Tetrahydrofuran				T
Trichloroethylene				T
Tris(2-chloroethyl) phosphate	T		T	T
Trp-P-1				T
Vinyl bromide				
Vinyl chloride			T	T
Z-Tetrachlorvinphos	T		T	T
beta-Hexachlorocyclohexane		T		T
p,p'-DDE				

Notes: Summary of analyzability of the ($N = 106$) matched carcinogens with available NHANES biomonitoring data. A value of **T** in Columns (2-5) indicates that the chemical was measurable for that particular statistic. *GM* denotes the geometric mean ratio. *AM* denotes the arithmetic mean ratio. *P95* denotes the 95th percentile ratio. *Detect* denotes the detected sample ratio.

Return to: Section 2.2.1.

Table A3: Summary of carcinogens' product use categories and their data coverage

Product Uses	(1) All	(2) Cand.	(3) Dir. BM	(4) Metab. BM	(5) RSEI	(6) DW	(7) HHP	(8) Any Data
Batteries	69	68	17	6	31	11	18	40
Cleaning and safety	91	87	27	14	46	20	32	58
Construction and building materials	103	101	29	15	53	20	30	66
Food and drug	95	90	16	8	32	16	16	41
Furniture and furnishings	59	57	15	8	30	9	20	40
Home maintenance	128	124	33	17	68	27	33	83
Industrial products	112	110	31	15	59	24	33	75
Laboratory supplies	308	298	67	25	143	56	38	170
Medical/dental	55	54	17	6	31	11	23	37
Personal care	91	86	22	15	43	17	29	57
Pesticides	46	44	15	10	30	13	18	36
Raw materials	186	180	45	22	94	36	39	113
Specialty occupational products	118	114	28	13	60	23	29	71
Vehicle	75	73	24	14	42	18	29	54

Notes: Summary of carcinogens' product use categories and the corresponding exposure data coverage for subsets of the 727 listed carcinogens and the 622 candidate carcinogens scoped into the analysis. All numeric values are applicable carcinogens counts. Column (1): all potential carcinogens. Column (2): all carcinogen candidates found to be ubiquitous. Column (3): set of carcinogens directly biomonitored by NHANES. Column (4): carcinogens indirectly biomonitored by metabolite. Column (5): carcinogens whose industrial releases are monitored in TRI/RSEI. Column (6) carcinogens monitored in drinking water by at least one PWS. Column (7): carcinogens matched as ingredients in purchased products for household product analysis. Column (8): set of carcinogens for which data was available (and detectable) in at least one analysis.

Return to: Section 2.4, Section 3.4.

Table A4: NHANES GM biomarker concentration ratios: primary results

Chemical													
1-Naphthylamine	1.45	0.82	0.77	1.93	0.98	0.79	0.92	1.08					
1-naphthol**	1.11	0.9	0.79	1.57	1.01	0.82	0.92	1.09					
2,4-Dichlorophenoxyacetic acid	0.97	1.02	0.89	1	1.04	0.9	0.89	1.13					
2,4-dichlorophenol**	1.36	0.82	1.67	2.44	0.76	0.74	0.97	1.03					
2,5-dichlorophenol**	2.08	0.63	2.98	6.15	0.54	0.77	0.91	1.1					
2,6-Dimethylaniline	1.05	0.98	0.96	1.26	1.01		0.97	1.03					
2-Anisidine	0.98	1.02	0.88	1.01	1.06	0.83	0.89	1.12					
2-Methylaniline	1.12	0.94	0.98	1.53	0.95	0.85	0.92	1.09					
2-Naphthylamine	1.23	0.89	0.84	1.68	0.98	0.82	0.92	1.09					
2-naphthol**	1.21	0.87	1.47	1.69	0.85	0.73	0.96	1.05					
4-Biphenylamine	1.21	0.9	0.84	1.45	1	0.84	0.96	1.04					
Acrylamide	1.06	0.97	0.95	1.13	1.01	0.92	0.94	1.07					
Arsenic	0.96	1.01	1.07	1.32	0.87	1.56	0.92	1.1					
Bis(2-chloroethyl) phosphate**	1.19	0.91	0.99	1.12	0.98	1.03	0.93	1.07					
Cadmium	1.05	0.96	0.77	1.1	1.03	1.2	1.14	0.87					
Cobalt	1	0.99	0.97	0.94	1.01	1	1.07	0.93					
Dimethyl phosphate**	1.05	0.97	1.16	1.18	0.95	0.85	0.97	1.04					
Dimethyl thiophosphate**	0.92	1.06	1	1.05	1.04	0.74	0.91	1.1					
Dimethylarsinic acid	1.01	0.94	1.2	1.1	0.89	1.27	0.95	1.05					
Ethylene oxide	1.23	0.87	0.92	1.46	0.95	1.04	0.92	1.09					
Formaldehyde	0.98	1.01	1	0.94	1.01	0.99	1	1					
Lead	1	0.99	0.83	1.01	1.03	1.17	0.88	1.15					
Mandelic acid**	1.06	0.96	0.99	1.35	0.97	0.86	0.89	1.13					
Melamine	1.05	0.99	0.71	0.73	1.1	1.62	0.92	1.09					
Mono-(2-ethyl-5-hydroxyhexyl) phthalate**	1.15	0.92	1.06	1.32	0.94	0.93	0.9	1.11					
Mono-(2-ethyl-5-oxohexyl) phthalate**	1.17	0.91	1.09	1.26	0.94	0.92	0.93	1.08					
Mono-2-ethyl-5-carboxypentyl phthalate**	1.18	0.9	1.13	1.19	0.94	0.94	0.93	1.08					
N-Acetyl-S-(n-propyl)-L-cysteine**	1.03	0.97	1.17	1.33	0.88	1.18	0.92	1.09					
N-acetyl-S-(2-Carboxyethyl)-L-cysteine**	1.07	0.95	0.92	1.63	0.96	0.84	0.87	1.16					
N-acetyl-S-(2-carbamoyl-ethyl)-L-cysteine**	1.06	0.96	0.98	1.45	0.97	0.83	0.86	1.17					
N-acetyl-S-(2-cyanoethyl)-L-cysteine**	1.38	0.82	0.7	2.39	0.99	0.69	0.77	1.32					
N-acetyl-S-(2-hydroxypropyl)-L-cysteine**	1.05	0.96	0.92	1.27	1	0.85	0.88	1.15					
N-acetyl-S-(3,4-dihydroxybutyl)-L-cysteine**	1.01	0.97	0.97	1.22	1	0.84	0.9	1.11					
N-acetyl-S-(3-Hydroxypropyl)-L-cysteine**	1.12	0.93	0.92	1.43	0.95	1.06	0.82	1.22					
N-acetyl-S-(N-methylcarbamoyl)-L-cysteine**	1.01	0.98	0.85	1.04	1.09	0.75	0.91	1.11					
Dem. Subgroup:													
Inc/PL ratio:	<2x P.L. >2x P.L.												
Race/eth:	Hispanic NH Black NH White NH Other												
Sex:	Female Male												

Continued, next page...

Table A4: NHANES GM biomarker concentration ratios: primary results

Chemical										
N-acetyl-s-(3-hydroxypropyl-1-methyl)-L-cysteine**	1.1	0.93	0.87	1.34	1.01	0.87	0.88	1.15		
NNAL**	2.19		0.62	2.41	1			1.33		
Nickel	1.17	0.9	1.01	1.22	0.96		1	0.97	1.03	
Nitromethane	0.99	1	1.02	1.1	0.96		1.14	1.02	0.98	
Perfluorooctanesulfonic acid	0.88	1.09	0.75	1	1.1		0.93	0.8	1.26	
Perfluorooctanoic acid	0.88	1.1	0.84	0.83	1.09		0.98	0.89	1.13	
Phenylglyoxylic acid**	1.01	0.98	0.89	1.29	1		0.9	0.89	1.12	
Dem. Subgroup:										
Inc/PL ratio:	<2x P.L.		>2x P.L.							
Race/eth:			Hispanic	NH Black	NH White	NH Other		Female	Male	
Sex:										

Notes: Presentation of the primary set of geometric mean ratios for the carcinogens and associated metabolites (N = 42) for which censoring by detection limits did not prevent statistical calculations. As summarized in Figure 1, after mapping metabolites to their parent chemicals, 39 separate carcinogens are represented by these results. **Bold** values denote concentration ratios >=1.25. Underlined values denote concentration ratios >=1.5. Statistics based on individual-level NHANES biomonitoring samples.

Demographic subgroups: Household income below 2 times the poverty line, Household income above 2 times the poverty line, Hispanic, non-Hispanic Black, non-Hispanic White, Other race including multi-racial, Female sex, Male sex.

Parent Chems: {1-naphthol: *1-Naphthalenol*, *1-(N-methylcarbamate)***}; {1-naphthol, 2-naphthol: *Naphthalene***}; {2,4-dichlorophenol: *2,4-Dichlorophenoxyacetic acid***}; {2,4-dichlorophenol: *Nitrofen***}; {2,5-dichlorophenol: *1,4-Dichlorobenzene***}; {2-naphthol: *Naphthalene***}; {Bis(2-chloroethyl) phosphate: *Tris(2-chloroethyl) phosphate***}; {Dimethyl phosphate: *Dichlorvos***}; {Dimethyl phosphate: *Malathion***}; {Dimethyl phosphate: *Z-Tetrachlorovinphos***}; {Dimethyl thiophosphate: *Malathion***}; {Mandelic acid: *Ethylbenzene***}; {Mono-(2-ethyl-5-hydroxyhexyl) phthalate, Mono-(2-ethyl-5-oxohexyl) phthalate, Mono-2-ethyl-5-carboxypentyl phthalate: *Di(2-ethylhexyl) phthalate***}; {N-Acetyl-S-(n-propyl)-L-cysteine: *1-Bromopropane***}; {N-acetyl-S-(2-Carboxyethyl)-L-cysteine, N-acetyl-S-(3-Hydroxypropyl)-L-cysteine: *Acrolein***}; {N-acetyl-S-(2-carbamoyl-ethyl)-L-cysteine: *Acrylamide***}; {N-acetyl-S-(2-cyanoethyl)-L-cysteine: *Acrylonitrile***}; {N-acetyl-S-(2-hydroxypropyl)-L-cysteine: *1,2-Propylene oxide***}; {N-acetyl-S-(3,4-dihydroxybutyl)-L-cysteine: *1,3-Butadiene***}; {N-acetyl-S-(N-methylcarbamoyl)-L-cysteine: *N,N-Dimethylformamide***}; {N-acetyl-S-(3-hydroxypropyl-1-methyl)-L-cysteine: *Crotonaldehyde***}; {NNAL: *4-(N-Methyl-N-nitrosamino)-1-(3-pyridyl)-1-butanone***}; {Phenylglyoxylic acid: *Styrene***}; {Phenylglyoxylic acid: *Ethylbenzene***}

Return to: Section 3.1.1.

Table A5: NHANES GM biomarker concentration ratios: demographic interactions

Chemical		0.68	0.88	<u>1.65</u>	<u>2.28</u>	0.77	0.91	1.05	0.81	0.71	<u>2.57</u>	1.24	1.11	<u>1.63</u>	0.81
1-Naphthylamine		0.68	0.91	<u>1.35</u>	<u>1.87</u>	0.7	0.96	1.05	0.83	0.75	<u>1.73</u>	<u>1.46</u>	0.98	0.7	1.18
1-naphthol**		0.69	0.91	<u>1.35</u>	<u>1.87</u>	0.7	0.96	1.05	0.83	0.75	<u>1.73</u>	<u>1.46</u>	0.98	0.7	1.18
2,4-Dichlorophenoxyacetic acid		0.8	0.98	0.93	1.1	1.08	0.92	1.19	0.88	0.9	0.98	1.04	0.93	0.9	1.02
2,4-dichlorophenol**		<u>1.74</u>	<u>1.61</u>	<u>2.6</u>	<u>2.26</u>	0.75	0.73	0.7	<u>1.87</u>	1.21	<u>3.14</u>	<u>1.8</u>	0.87	0.67	0.89
2,5-dichlorophenol**		<u>3.07</u>	<u>2.9</u>	<u>6.54</u>	<u>5.71</u>	0.7	0.85	0.64	<u>4.25</u>	<u>1.54</u>	<u>10.39</u>	<u>3.52</u>	1.13	0.54	0.8
2,6-Dimethylaniline		1.02	1.19	<u>1.34</u>			0.99	1.03	0.88	<u>1.26</u>	<u>1.32</u>	1.07		1.09	0.98
2-Anisidine		0.79	0.97	0.92	1.1	0.77	0.89	0.93	1.21	0.98	0.95	1.05	0.9	0.79	1.09
2-Methylaniline		0.91	1.05	<u>1.4</u>	<u>1.69</u>	0.79	0.92	1.04	0.96	1.05	<u>1.64</u>	<u>1.37</u>	1.04	0.76	1.07
2-Naphthylamine		0.76	0.92	<u>1.56</u>	<u>1.81</u>	0.79	0.86	0.9	1.07	0.85	<u>1.98</u>	<u>1.28</u>	0.89	0.8	<u>1.31</u>
2-naphthol**		<u>1.42</u>	<u>1.51</u>	<u>1.72</u>	<u>1.65</u>	0.67	0.79	0.81	0.9	<u>1.55</u>	<u>1.75</u>	<u>1.63</u>	0.82	0.67	0.97
4-Biphenylamine		0.76	0.92	<u>1.35</u>	<u>1.56</u>	0.96	0.71	0.95	1.05	0.83	<u>1.65</u>	1.18	0.88	0.83	<u>1.35</u>
Acrylamide		0.9	1	<u>1.31</u>		0.89	0.96	1.07	0.96	0.94	<u>1.27</u>	1	0.96	0.91	1.08
Arsenic		0.96	1.2	<u>1.31</u>	<u>1.34</u>	<u>1.87</u>	0.8	0.95	0.98	1.2	1.16	<u>1.54</u>	<u>1.49</u>	<u>1.55</u>	0.79
Bis(2-chloroethyl) phosphate**		0.92	1.07	1.12	1.11	0.97	1.09	0.9	1.06	0.99	1	1.21	1.01	1.2	0.96
Cadmium		0.87	0.68	1.17	1.02	<u>1.42</u>	1.02	1.18	0.89	0.74	1.1	1.02	<u>1.37</u>	1.08	1.18
Cobalt		1.09	0.86	0.99	0.87	1.11	0.91	1.07	0.96	1	0.96	0.95	0.96	1.05	1.02
Dimethyl phosphate**		1.12	1.21	1.05	<u>1.36</u>	0.81	0.91	0.94	0.96	1.23	1.07	<u>1.36</u>	0.96	0.81	0.98
Dimethyl thiophosphate**		1.03	0.98	0.91	1.23	0.68	0.81	0.93	1.17	1.01	0.89	0.91	<u>1.28</u>	0.78	0.92
Dimethylarsinic acid		1.13	<u>1.27</u>	1.14	1.05	<u>1.27</u>	<u>1.26</u>	0.96	1.16	1.18	1.07	1.03	1.05	<u>1.33</u>	0.92
Ethylene oxide		0.87	0.98	1.23	<u>1.79</u>	1.08	1.01	0.86	1.05	0.98	<u>1.78</u>	1.18	<u>1.25</u>	0.92	1.24
Formaldehyde		1.01	1	0.95	0.94	0.98	0.99	1.01	0.99	1.02	0.95	0.93	1	0.98	1.02
Lead		0.71	0.96	0.92	1.13	1.06	<u>1.29</u>	0.89	1.19	0.85	0.79	1	0.95	<u>1.29</u>	1.04
Mandelic acid**		0.88	1.12	1.21	<u>1.53</u>	0.74	1	0.87	1.09	0.99	1	<u>1.34</u>	<u>1.35</u>	0.89	1.06
Melamine		0.57	0.88	0.81	0.64	1.1	<u>2.13</u>	1.03	1.19	0.81	0.57	0.68	<u>2.13</u>	<u>1.35</u>	1.24
Mono-(2-ethyl-5-hydroxyhexyl) phthalate**		0.98	1.15	1.23	<u>1.42</u>	0.86	1.02	0.84	1.06	1.15	<u>1.36</u>	<u>1.32</u>	1.13	0.86	1.1
Mono-(2-ethyl-5-oxohexyl) phthalate**		1.04	1.14	1.19	<u>1.34</u>	0.87	1	0.86	1.04	1.21	<u>1.3</u>	<u>1.28</u>	1.13	0.86	1.11
Mono-2-ethyl-5-carboxypentyl phthalate**		1.1	1.17	1.13	<u>1.26</u>	0.87	1.03	0.87	1.03	<u>1.25</u>	<u>1.27</u>	1.16	1.13	0.86	1.12
N-Acetyl-S-(n-propyl)-L-cysteine**		1.1	1.24	<u>1.33</u>	<u>1.32</u>	0.97	<u>1.44</u>	0.81	0.97	1.05	<u>1.32</u>	<u>1.4</u>	1.14	1.2	0.92
N-acetyl-S-(2-Carboxyethyl)-L-cysteine**		0.79	1.07	<u>1.46</u>	<u>1.85</u>	0.65	1.1	0.85	1.09	0.93	0.9	<u>1.64</u>	<u>1.57</u>	0.82	1.07
N-acetyl-S-(2-carbamoyl-ethyl)-L-cysteine**		0.84	1.13	1.23	<u>1.76</u>	0.65	1.08	0.84	1.12	0.97	<u>1.56</u>	<u>1.36</u>	0.76	0.87	1.05
N-acetyl-S-(2-cyanoethyl)-L-cysteine**		0.53	0.94	<u>1.62</u>	<u>3.78</u>	0.51	0.95	0.78	1.26	0.71	0.72	<u>3.46</u>	1.02	0.55	<u>1.63</u>
N-acetyl-S-(2-hydroxypropyl)-L-cysteine**		0.81	1.03	1.13	<u>1.47</u>	0.67	1.09	0.89	1.14	0.95	0.88	<u>1.31</u>	<u>1.26</u>	0.9	1.12
N-acetyl-S-(3,4-dihydroxybutyl)-L-cysteine**		0.86	1.1	1.12	<u>1.35</u>	0.69	1.01	0.91	1.09	0.96	0.97	1.23	1.18	0.78	1.04
N-acetyl-S-(3-Hydroxypropyl)-L-cysteine**		0.76	1.11	1.18	<u>1.76</u>	0.83	<u>1.37</u>	0.78	1.15	0.93	0.92	<u>1.49</u>	1.3	1.21	0.98
N-acetyl-S-(N-methylcarbamoyl)-L-cysteine**		0.76	0.96	0.93	1.18	0.63	0.89	1.01	1.18	0.83	0.89	0.99	0.74	0.74	1.21
Dem. Subgroup:															
Inc/PL ratio:															
Race/eth:		Hsp	Hsp	Blk	Blk	Oth	Oth	Whit	Hsp	Hsp	Blk	Blk	Oth	Oth	>2
Sex:		F	M	F	M	F	M	F	M	M	M	M	M	M	Whit

Continued, next page...

Table A7: NHANES pooled biomarker mean concentration ratios: primary results

Chemical							
Beta-Hexachlorocyclohexane	0.63	<u>9.22</u>		0.49	0.66	1.2	0.79
DDT	<u>2.01</u>	<u>3.42</u>	0.98	0.55	1.12	0.94	1.07
PBB153	0.63	0.64	0.97	1.13	1.12	0.75	1.27
PCB074	0.51	1.05	1.05	1.13	0.8	1.21	0.78
PCB105	0.61	<u>1.73</u>	1.12	1.04	0.65	1.11	0.88
PCB126	0.74	<u>2.02</u>	0.9	1.01		1.08	0.91
PCB146	0.51	<u>2.33</u>	1.44	0.94	0.77	1.05	0.94
PCB156	0.45	0.87	0.97	1.16	0.99	1.05	0.94
PCB157		0.94		1.15	0.95		0.94
PCB167		<u>1.56</u>	1.13	1.07		1.14	0.86
PCB169	0.64	1.4	0.89	1.08		0.99	1.02
PCB170	0.54	1.19	1.1	1.08	1.04	1.01	0.99
PCB178	0.51	<u>1.84</u>	1.21	1.02	0.86	1.02	0.98
PCB183	0.59	<u>1.94</u>	1.34	0.98	0.84	1.05	0.94
PCB187	0.53	<u>2.2</u>	1.35	0.96	0.82	1.05	0.94
PCB194	0.48	1.02	1.06	1.12	1.05	0.94	1.06
PCB196	0.49	1.15	1.1	1.1	0.96	0.98	1.02
PCB199	0.48	1.28	1.12	1.09	0.97	0.98	1.02
PCB206	0.49	0.97	1.17	1.1	1	1.01	0.99
PCB209	0.49	1	1.25	1.09	0.96	1.09	0.91
PeCDF (furan)	0.74	1.12	0.66	1.12		0.94	1.06
Dem. Subgroup:							
Race/eth:	Hispanic	NH Asian	NH Black	NH White	Other	Female	Male
Sex:							

Notes: Presentation of the primary set of arithmetic mean ratios for the carcinogens and associated metabolites ($N = 21$) for which censoring by detection limits did not prevent statistical calculations. Statistics based on pooled NHANES biomonitoring samples. **Bold** values denote concentration ratios >1.25 . Underlined values denote concentration ratios >1.5 .

Demographic subgroups: Hispanic, non-Hispanic Black, non-Hispanic White, Other race including multi-racial, Female sex, Male sex.

Return to: Section 3.1.2.

Table A8: NHANES pooled biomarker mean concentration ratios: demographic interactions

Chemical													
Beta-Hexachlorocyclohexane	0.77	0.49	<u>9.61</u>	<u>8.79</u>			0.2	0.69	0.29	0.81			
DDT	<u>2</u>	<u>2.03</u>	<u>2.72</u>	<u>4.25</u>	1.26		0.66	0.46	0.66	<u>1.52</u>	0.64		
PBB153	0.52	0.74	0.36	0.95	1		0.94	0.79	1.48	<u>0.84</u>	1.41		
PCB074	0.56	0.46	1.2	0.9	1.37		0.64	1.36	0.89	1.07	0.53		
PCB105	0.66	0.57	<u>2.05</u>	1.39	1.29		0.91	1.13	0.94	0.76	0.53		
PCB126	0.78	0.7	2.36	<u>1.64</u>	0.91		0.89	1.09	0.92	0.7			
PCB146	0.47	0.55	<u>2.52</u>	<u>2.11</u>	1.41		1.48	1.01	0.87	0.84	0.71		
PCB156	0.43	0.47	0.94	0.8	1.02		0.9	1.23	1.1	1.18	0.8		
PCB157			1.01	0.87	1.09				1.09	1.15			
PCB167		0.54	<u>1.8</u>	1.29	1.24			1.23	0.91				
PCB169		0.69	1.34	1.47			0.99	1.1	1.06				
PCB170	0.51	0.57	1.27	1.11	1.06		1.15	1.1	1.06	1.06	1.02		
PCB178		0.58	<u>2.09</u>	1.56	1.06		1.39	1.07	0.98	0.9			
PCB183	0.56	0.61	<u>2.32</u>	1.55	1.36		1.31	1.02	0.93	0.88	0.8		
PCB187	0.47	0.6	<u>2.47</u>	<u>1.91</u>	1.27		1.46	1.04	0.87	0.84	0.8		
PCB194	0.43	0.53	1.05	0.99	0.89		1.26	1.08	1.17	0.98	1.13		
PCB196	0.45	0.53	1.26	1.02	1		1.23	1.09	1.12	0.95	0.96		
PCB199	0.44	0.53	1.4	1.16	1		1.26	1.08	1.09	0.97	0.97		
PCB206		0.51	1.03	0.91	1.07		1.29	1.14	1.07	1.03	0.97		
PCB209		0.48	1.08	0.9	1.14		1.39	1.23	0.95	1.07	0.85		
PeCDF (furan)	0.65	0.83	1.01	1.21	0.67		0.65	1.07	1.18		0.91		
Dem. Subgroup:													
Race/eth:	Hispanic	Hispanic	NH Asian	NH Asian	NH Black	NH Black	NH Black	NH White	NH White	Other	Other	Other	Other
Sex:	Female	Male	Female	Male	Female	Male	Female	Female	Male	Female	Male	Female	Male

Notes: Demographic interaction analysis showing arithmetic mean ratios for the carcinogens and associated metabolites (N = 21) for which censoring by detection limits did not prevent statistical calculations. Statistics based on pooled NHANES biomonitoring samples. Subgroups in this table include interactions of race/ethnicity and sex, and interactions of race/ethnicity and household income. **Bold** values denote concentration ratios ≥ 1.25 . Underlined values denote concentration ratios ≥ 1.5 .
Demographic subgroups: Hispanic, non-Hispanic Asian, non-Hispanic Black, non-Hispanic White, Other race including multi-racial, Female sex, Male sex.
Return to: Section 3.1.2.

Table A9: NHANES pooled biomarker mean concentration ratios: age-differentiated results

Chemical													
Beta-Hexachlorocyclohexane													
DDT	0.5	1.07	1.11	0.67	1.39	10.18	0.44	0.54	0.73				
PBB153	0.23	1.11	1.13	0.74	1.21	3.72	1.13	0.55	1.24				
PCB074	0.52	1.07	1.11	0.58	0.28	0.7	1.11	1.25	1.28				
PCB105	0.52	1.07	1.11	0.69	0.53	1.15	1.19	1.23	0.89				
PCB126	1.08	1.11	1.13	0.83	0.53	1.88	1.26	1.08	0.71				
PCB146	0.2	1.11	1.13	0.6	0.45	2.16	1	1.06					
PCB156	0.18	1.12	1.13	0.53	0.19	2.56	1.66	1.03	0.87				
PCB157	1.1	1.1	1.1	0.62	0.22	0.96	1.11	1.28	0.1				
PCB167	1.1	1.1	1.1	0.62	0.22	1.03	1.13	1.25	1.06				
PCB169	1.1	1.1	1.1	0.71	0.22	1.7	1.29	1.14					
PCB170	0.14	1.12	1.13	0.64	0.22	1.49	0.99	1.18	1.19				
PCB178	1.12	1.12	1.13	0.6	0.41	1.31	1.27	1.19	0.97				
PCB183	1.11	1.13	1.13	0.68	0.37	2.03	1.39	1.13	0.95				
PCB187	0.14	1.13	1.13	0.64	0.37	2.13	1.54	1.07	0.92				
PCB194	0.08	1.13	1.13	0.57	0.16	2.43	1.56	1.06	1.2				
PCB196	0.1	1.13	1.13	0.58	0.2	1.13	1.22	1.25	1.08				
PCB199	1.13	1.13	1.13	0.58	0.22	1.26	1.27	1.22	1.1				
PCB206	1.12	1.13	1.13	0.57	1.06	1.39	1.29	1.21	1.13				
PCB209	1.13	1.13	1.13	0.57	1.08	1.34	1.22	1.22	1.09				
PeCDF (furan)	0.41	1.09	1.09	0.8	0.79	1.45	1.21	1.21	0.37				
Dem. Subgroup:													
Age:	12-19	20+	Hispanic	20+	12-19	NH Asian	20+	12-19	NH Black	20+	12-19	NH White	20+
Race/eth:			Hispanic										

Notes: Age-based analysis showing arithmetic mean ratios for the carcinogens and associated metabolites (N = 21) for which censoring by detection limits did not prevent statistical calculations. Statistics based on pooled NHANES biomonitoring samples. Demographic subgroups presented by age, including age subgroups differentiated by race/ethnicity. **Bold** values denote concentration ratios ≥ 1.25 . Underlined values denote concentration ratios ≥ 1.5 .

Demographic subgroups: Hispanic, non-Hispanic Asian, non-Hispanic Black, non-Hispanic White, Other race including multi-racial

Return to: Section 3.1.2.

Table A10: NHANES P95 biomarker concentration ratios: primary results

Chemical	1.17	0.61	1.34	3.4	0.6	0.69	0.9	1.02
1,4-Dichlorobenzene	1.17	0.61	1.34	3.4	0.6	0.69	0.9	1.02
1-Naphthylamine	1.41	0.72	0.41	1.51	1.03	0.55	0.9	1.04
1-naphthol**	1.32	0.8	0.75	1.43	1.06	0.67	1.01	1
2,4,5-trichlorophenol**	1	1	0.67	0.67	1	1	1	1
2,4,6-Trichlorophenol	1.18	0.91	1.18	1.36	0.91	0.91	1	1
2,4-Dichlorophenoxyacetic acid	0.98	1	0.92	0.83	1.04	0.9	0.88	1.08
2,4-dichlorophenol**	1.71	0.62	2.3	7.21	0.55	0.7	1.05	1
2,5-dichlorophenol**	2.61	0.33	3.62	12.67	0.22	0.32	1.14	0.91
2,6-Dimethylaniline	1.01	1.04	0.76	0.89	1.18	0.68	1.14	0.89
2-Amino-1-methyl-6-phenylimidazo [4,5-b]pyridine	0.58	1.21	0.92	1.1	1.06	0.86	0.91	1.09
2-Amino-alpha-carboline	1.37	0.72	0.41	1.37	1.04	0.45	0.94	1.05
2-Anisidine	0.91	1.06	0.86	0.81	1.06	0.85	0.93	1.05
2-Methylaniline	1.08	0.8	0.88	1.2	0.94	0.93	0.93	1.04
2-Naphthylamine	1.22	0.83	0.55	1.42	1.05	0.74	0.94	1.05
2-Phenylphenol	1	1	1	1.4	1	0.6	1	1
2-naphthol**	1.21	0.87	1.07	1.44	0.94	0.85	1.03	1
4,4'-Diaminobiphenyl methane	1.02	1.02	1.07	1.11	0.99	1.03	0.99	1.02
4-Biphenylamine	1.26	0.84	0.61	1.35	1.09	0.78	1.06	0.95
Acrylamide	1.27	0.88	0.86	1.68	0.99	0.74	0.84	1.27
Arsenic	0.85	0.97	0.91	1.4	0.81	1.64	0.96	1.12
Arsenic acid	1.01	1.01	1.16		0.94	0.93	1	0.99
Benzene	1.45	0.54	0.42	1.27	1.06	1.22	0.93	1.06
Bis(2-chloroethyl) phosphate**	1.34	0.82	0.88	0.91	1.04	1.22	0.83	1.26
Bromodichloromethane	1	1	1	1	0.91	1.09	1	1
Bromoform	1.3	1	1.2	0.8	1	1	1	1.1
Cadmium	1.21	0.81	0.62	1.05	1.05	1.12	1.14	0.89
Chloroform	1.03	1	0.91	1.15	1	1.26	0.94	1.03
Cobalt	1.02	1	0.95	0.86	1.05	0.86	1.09	0.88
Crotonaldehyde	1.07	0.99	1.44	1	0.9	1.22	0.99	1.07
Dieldrin	0.89	1.13	0.77	0.84	1.03	1.37	0.89	1.15
Diisononyl phthalate	1	1	1.33	1.06	0.94	0.78	0.94	1.06
Dimethyl dithiophosphate**	0.67	1.3	0.63	0.89	1.3	0.5	0.87	1.32
Dimethyl phosphate**	1.25	0.95	1.25	1.32	0.98	0.59	0.94	1.07
Dimethyl thiophosphate**	1	1	0.73	1.13	1.06	0.5	0.74	1.07
Dimethylarsinic acid	0.95	0.9	1.15	1.11	0.79	1.66	1.06	0.9
Dem. Subgroup:								
Inc/PL ratio:	<2x P.L.	>2x P.L.						
Race/eth:			Hispanic	NH Black	NH White	NH Other	Female	Male
Sex:								

Continued, next page...

Table A10: NHANES P95 biomarker concentration ratios: primary results

Chemical		1.23	0.73	0.58	1.08	1.09	1.19	0.92	1.08
Ethylbenzene		1.12	0.94	0.55	1.25	1.1	0.68	0.9	1.05
Ethylene oxide		1.4	0.79	1.14	1.47	0.7	2.77	0.95	1
Ethylene thiourea		1	1.01	1.03	0.99	1	0.98	1	1
Formaldehyde		1.51	0.57	0.43	1.09	1.05	1.38	0.93	1.08
Furan		1.04	1	0.8	0.98	1.08	0.88	1.04	0.98
Heptachlor epoxide B		1.09	0.96	0.87	1.13	0.96	1.13	0.87	1.15
Lead		0.88	1.01	0.65	0.65	1.14	0.8	1.07	0.9
Malathion dicarboxylic acid**		1.23	0.85	0.97	1.35	0.93	0.88	0.98	1.02
Mandelic acid**		1.42	0.61	0.32	1.49	1.1	0.5	0.78	1.22
MeA-alpha-C		1.11	1	1.35	1	0.96	0.39	1	1.11
Melamine		0.97	0.97	1.12	0.99	0.93	1.11	0.99	1.01
Methylarsonic acid		1.08	0.91	1.02	1.23	1	1.07	0.92	1.11
Mono-(2-ethyl-5-hydroxyhexyl) phthalate**		1.13	0.9	1.07	1.15	0.95	1.02	0.91	1.09
Mono-(2-ethyl-5-oxohexyl) phthalate**		1.04	0.9	1.3	1.32	0.84	1.18	0.84	1.14
Mono-2-ethyl-5-carboxypentyl phthalate**		1.09	0.8	1.12	1.18	0.88	1.18	0.9	1.12
N ² -Nitrosornicotine		1.38	0.73	0.29	1.1	1.12	0.86	0.73	1.26
N-Acetyl-S-(1-cyano-2-hydroxyethyl)-L-cysteine (CYHA)**		1.48	0.7	0.57	1.67	0.98	0.63	0.87	1.13
N-Acetyl-S-(n-propyl)-L-cysteine**		0.9	1.01	1.19	1.04	0.95	1.28	1.09	0.96
N-Acetyl-S-(phenyl)-L-cysteine**		0.94	1.03	1.23	0.82	0.98	1.1	0.93	1.05
N-acetyl-S-(2-Carboxyethyl)-L-cysteine**		1.05	0.97	0.75	1.45	1.02	0.66	0.94	1.11
N-acetyl-S-(2-Hydroxyethyl)-L-cysteine**		1.42	0.8	0.85	1.6	0.82	1.1	1.05	0.96
N-acetyl-S-(2-carbamoyl-ethyl)-L-cysteine**		1.2	0.86	0.93	1.46	0.9	0.95	0.8	1.17
N-acetyl-S-(2-cyanoethyl)-L-cysteine**		1.34	0.82	0.61	1.96	0.95	0.68	0.76	1.34
N-acetyl-S-(2-hydroxy-3-butenyl)-L-cysteine**		1.3	0.67	0.85	1.44	1.02	0.59	0.91	1.14
N-acetyl-S-(2-hydroxypropyl)-L-cysteine**		1.01	0.84	0.75	0.89	1.16	0.84	1.04	0.95
N-acetyl-S-(3,4-dihydroxybutyl)-L-cysteine**		1.05	0.94	0.9	1.05	1.03	0.91	1.01	1
N-acetyl-S-(3-Hydroxypropyl)-L-cysteine**		1.09	0.96	0.69	1.45	1.07	0.79	1.05	1
N-acetyl-S-(N-methylcarbamoyl)-L-cysteine**		1.16	0.9	0.67	1.21	1.13	0.78	0.86	1.1
N-acetyl-s-(3-hydroxypropyl-1-methyl)-L-cysteine**		1.2	0.87	0.65	1.31	1.11	0.83	0.93	1.09
NNAL**		1.29	0.7	0.34	1.13	1.14	0.76	0.74	1.22
Nickel		1.1	0.84	0.94	1.1	0.99	0.97	0.98	1.02
Nitromethane		1.04	0.95	0.95	1.24	0.94	1.06	1.02	0.96
Oxypyrimidine**		1.42	0.86	0.96	1.91	0.84	0.86	1	1.02
Pentachlorophenol		0.94	1.02	0.8	1.12	1.02	0.77	0.88	1.26
Dem. Subgroup:									
Inc/PL ratio:		<2x P.L.	>2x P.L.	Hispanic	NH Black	NH White	NH Other	Female	Male
Race/eth:									
Sex:									

Continued, next page...

Table A10: NHANES P95 biomarker concentration ratios: primary results

Chemical										
Perfluorooctanesulfonic acid	0.96	1.03	0.68	<u>1.5</u>	0.95	1.36	0.9	1.08		
Perfluorooctanoic acid	0.97	1.11	0.79	0.89	1	1.46	1	1		
Phenylglyoxylic acid**	1	0.96	0.84	1.27	1.03	0.82	0.94	1.04		
Styrene	1.15	0.97	0.64	1.27	1.14	0.7	0.83	1.15		
Tetrachloroethylene		1.21	0.78	1.03	1.07		0.74	1.18		
Dem. Subgroup:										
Inc/PL ratio:	<2x P.L. >2x P.L.									
Race/eth:	Hispanic NH Black NH White NH Other									
Sex:	Female Male									

Notes: Presentation of the primary set of 95th percentile ratios for the carcinogens and associated metabolites (N = 75) for which censoring by detection limits did not prevent statistical calculations. As summarized in Figure 3, after mapping metabolites to their parent chemicals, 62 separate carcinogens are represented by these results. **Bold** values denote concentration ratios >=1.25. Underlined values denote concentration ratios >=1.5. Statistics based on individual-level NHANES biomonitoring samples.

Demographic subgroups: Household income below 2 times the poverty line, Household income above 2 times the poverty line, Hispanic, non-Hispanic Black, non-Hispanic White, Other race including multi-racial, Female sex, Male sex.

Parent Chemicals: {1-naphthol: 1-Naphthalenol, 1-(N-methylcarbamate)**}; {1-naphthol, 2-naphthol: Naphthalene**}; {2,4-dichlorophenol: 2,4-Dichlorophenoxyacetic acid**}; {2,4-dichlorophenol: Nitrofen**}; {2,5-dichlorophenol: 1,4-Dichlorobenzene**}; {2-naphthol: Naphthalene**}; {Bis(2-chloroethyl) phosphate: Tris(2-chloroethyl) phosphate**}; {Dimethyl phosphate: Dichlorvos**}; {Dimethyl phosphate: Malathion**}; {Dimethyl phosphate: Malathion**}; {Dimethyl thiophosphate: Malathion**}; {Dimethyl dicarboxylic acid: Malathion**}; {Mandelic acid: Styrene**}; {Mandelic acid: Ethylbenzene**}; {Mono-(2-ethyl-5-hydroxyhexyl) phthalate, Mono-(2-ethyl-5-oxohexyl) phthalate, Mono-(2-ethyl-5-carboxypentyl) phthalate: Di(2-ethylhexyl) phthalate**}; {N-Acetyl-S-(1-cyano-2-hydroxyethyl)-L-cysteine (CYHA), N-acetyl-S-(2-Hydroxyethyl)-L-cysteine, N-acetyl-S-(2-Cyanoethyl)-L-cysteine: Acrylonitrile**}; {N-Acetyl-S-(n-propyl)-L-cysteine: 1-Bromopropane**}; {N-Acetyl-S-(phenyl)-L-cysteine: Benzene**}; {N-acetyl-S-(2-Hydroxyethyl)-L-cysteine, N-acetyl-S-(3-Hydroxypropyl)-L-cysteine: Acrolein**}; {N-Acetyl-S-(2-Hydroxyethyl)-L-cysteine: Vinyl Chloride**}; {N-acetyl-S-(2-Hydroxyethyl)-L-cysteine, N-acetyl-S-(3-Hydroxypropyl)-L-cysteine: Acrolein**}; {N-Acetyl-S-(2-Hydroxyethyl)-L-cysteine: Vinyl Chloride**}; {N-acetyl-S-(2-Hydroxyethyl)-L-cysteine, N-acetyl-S-(2-Cyanoethyl)-L-cysteine: Ethylene Oxide**}; {N-acetyl-S-(2-Cyanoethyl)-L-cysteine: N,N-Dimethylformamide**}; {N-acetyl-S-(2-Hydroxyethyl)-L-cysteine, N-acetyl-S-(3-Hydroxypropyl)-L-cysteine: N,N-Dimethylformamide**}; {N-acetyl-S-(2-Hydroxyethyl)-L-cysteine, N-acetyl-S-(3-Hydroxypropyl)-L-cysteine, N-acetyl-S-(3,4-dihydroxybutyl)-L-cysteine: 1,3-Butadiene**}; {N-acetyl-S-(2-Hydroxypropyl)-L-cysteine: N,N-Dimethylformamide**}; {N-acetyl-S-(3-Hydroxypropyl)-L-cysteine, N-acetyl-S-(3,4-dihydroxybutyl)-L-cysteine: 1,3-Butadiene**}; {N-acetyl-S-(N-methylcarbamoyl)-L-cysteine: 1,2-Pyridyl-1-butanone**}; {Oxypyrimidine: Diazonon**}; {Phenylglyoxylic acid: Styrene**}; {Phenylglyoxylic acid: Ethylbenzene**}

Return to: Section 3.1.3.

Table A11: NHANES P95 biomarker concentration ratios: demographic interactions

Chemical	1.17	1.34	3.87	3.13	0.86	0.66	0.41	0.74	1.34	0.73	4.07	2.12	1.01	0.47	0.74	0.44
1,4-Dichlorobenzene	0.27	0.55	1.19	1.86	0.55	0.57	0.91	1.21	0.67	0.26	1.86	1.13	0.63	0.6	1.62	0.74
1-Naphthylamine	0.63	0.75	1.17	1.54	0.67	0.83	1.24	1.04	0.85	0.51	1.31	1.52	0.71	0.52	2.16	0.8
1-naphthol**	0.67	0.67	1	0.67	0.67	1	1	1	0.67	0.67	0.67	0.67	1.67	0.33	1	1
2,4,5-trichlorophenol**	1.09	1.18	1.45	1.18	0.82	0.91	1	0.91	1	1.45	1.73	0.91	0.82	0.91	1.09	0.82
2,4,6-Trichlorophenol	0.8	1.13	0.83	0.86	0.9	0.89	0.95	1.15	0.83	1.05	0.83	0.94	0.75	0.99	1.06	1
2,4-Dichlorophenoxyacetic acid	2.29	2.57	11.42	6.36	0.82	0.38	0.58	0.53	3.04	1.36	15.14	4.95	0.82	0.31	0.6	0.53
2,4-dichlorophenol**	3.29	3.84	15.05	10.59	0.25	0.32	0.22	0.21	4.94	1.98	21.81	8.58	0.43	0.18	0.57	0.18
2,5-dichlorophenol**	0.66	0.88	0.69	3.62	0.72	0.68	1.55	0.89	0.61	1.79	1.19	0.69	0.87	0.56	1.18	1.25
2,6-Dimethylaniline	0.92	0.92	0.7	1.23	0.94	0.6	0.9	1.2	0.87	0.9	0.63	1.28	0.91	0.71	0.49	1.27
2-Amino-1-methyl-6-phenylimidazo [4,5-b]pyridine	0.26	0.49	1.26	1.53	0.47	0.3	1.01	1.12	0.54	0.16	1.38	1.1	0.9	0.43	1.53	0.81
2-Amino-alpha-carboline	0.61	1.04	0.82	0.8	0.79	0.86	1.05	1.09	0.82	1.04	0.75	0.9	0.84	0.86	1.05	1.07
2-Anisidine	0.88	0.95	1.2	1.45	0.88	1.15	0.92	1.02	0.95	0.82	1.4	0.9	0.81	0.88	1.08	0.75
2-Methylaniline	0.5	0.56	1.52	1.27	0.73	0.77	0.94	1.21	0.62	0.54	1.52	1.24	0.65	0.91	1.47	0.85
2-Naphthylamine	1	1	1.2	1.4	0.6	0.6	1	1	1	1	1.2	1.4	0.6	0.4	1	1
2-Phenylphenol	1.06	1.2	1.47	1.39	0.96	0.74	0.94	0.9	1.03	0.96	1.66	1.32	0.87	0.8	1.23	0.82
2-naphthol**	1.14	1.01	1.11	1.11	0.89	1.24	0.97	0.99	1.07	1.04	1.22	1.02	1.21	1.02	0.87	1
4,4'-Diaminobiphenyl methane	0.65	0.59	1.61	1.32	1.02	0.61	1.09	1.07	0.71	0.54	1.46	1.21	0.64	0.79	1.55	0.84
4-Biphenylamine	0.62	0.98	1.07	1.78	0.77	0.73	0.85	1.27	0.86	0.8	2.01	1	0.95	0.59	1.27	0.92
Acrylamide	0.91	0.98	1.4	1.67	1.64	1.58	0.77	0.83	0.73	1.06	1.3	1.43	1.82	1.47	0.57	0.85
Arsenic	1.09	1.23			0.93	0.95	1	0.91	1.14	1.23		0.94		1.07	1.05	1
Arsenic acid	0.35	0.51	1.09	1.41	1.1	1.38	1.07	1.05	0.44	0.39	1.13	1.07	1.45	0.87	1.73	0.43
Benzene	0.75	0.98	0.86	1.26	1.87	1.07	0.71	1.34	1	0.72	1.26	0.55	1.98	0.75	1.34	0.92
Bis(2-chloroethyl) phosphate**	1.18	1	1	1	0.82	1.09	0.91	1	1	1.18	1.09	1	0.73	1.09	1.09	0.91
Bromodichloromethane	1	1.4		1.1	1.2	1	1	1	1.5	1.2		1.1	1.2	0.8	1.5	0.9
Bromoform	0.64	0.61	1.04	1.05	1.2	1.12	1.27	0.93	0.64	0.64	1.05	0.95	1.2	0.87	1.38	0.8
Cadmium	0.85	0.97	1.15	1.24	1.26	0.97	0.88	1.12	0.94	0.94	1.12	1.24	1.26	1.06	0.91	1
Chloroform	1.16	0.65	0.88	0.74	1.12	0.65	1.09	1	0.95	0.81	0.86	0.91	0.81	0.91	1.09	1.02
Cobalt	1.48	1.4	0.85	1.11	0.82	1.22	0.98	0.89	1.65	1.22	1.14	0.92	1.36	0.87	0.86	0.99
Crotonaldehyde	0.89	0.77	0.84	0.75	2.73	0.78	0.86	1.23	0.66	1.13	0.84	0.83	0.82	2.43	0.95	1.16
Dieldrin	1.28	1.33	1.06	1.11	0.67	1	0.89	1.06	1.72	1.22	1.39	0.89	0.56	0.72	0.94	1.06
Diisooxyl phthalate	0.74	0.59	0.86	0.98	0.4	0.73	0.94	1.82	0.58	1	0.71	0.89	0.47	0.73	0.8	1.52
Dimethyl dithiophosphate**	1.45	0.91	0.85	1.72	0.61	0.56	0.99	0.98	1.32	0.89	1.14	1.36	0.61	0.63	1.33	0.95
Dimethyl phosphate**	0.76	0.66	0.88	1.38	0.36	0.61	0.84	1.07	0.64	0.73	0.86	1.13	0.48	0.58	1.07	1.03
Dimethyl thiophosphate**	1.27	1.09	1.28	0.93	1.76	1.56	0.86	0.72	1.06	1.37	1.06	0.89	1.17	1.86	0.74	0.75
Dimethylarsinic acid																
Dem. Subgroup:	Hsp	Hsp	Blk	Blk	Oth	Oth	Wht	Wht	Hsp	Hsp	Blk	Blk	Oth	Oth	Wht	>2
Inc/PL ratio:	F	M	F	M	F	M	F	M	Hsp	Hsp	Blk	Blk	Oth	Oth	Wht	>2
Race/eth:																
Sex:																

Continued, next page...

Table A11: NHANES P95 biomarker concentration ratios: demographic interactions

Chemical	0.47	0.73	0.91	1.29	0.91	1.27	1.08	1.12	0.6	0.6	1.02	1.01	1.25	0.97	1.31	0.61
Ethylbenzene	0.45	0.71	1.04	1.52	0.84	0.57	0.91	1.12	0.64	0.6	1.37	1.19	1.28	0.27	1.26	1.1
Ethylene oxide	1.05	1.16	1.19	1.93	3.09	2.77		0.84	1.35	0.74	1.53	1.19	3.09	0.88	0.79	
Ethylene thiourea	1.03	1.03	0.99	0.99	0.99	0.98	1	1	1.03	1.05	0.99	0.99	0.99	0.97	0.95	1
Formaldehyde	0.43	0.5	0.97	1.31	1	1.49	1.15	0.97	0.43	0.42	1.06	1.2	1.49	1.27	1.84	0.4
Furan																
Heptachlor epoxide B	1.04	0.67	1.11	0.76	1.26	0.88	1.04	1.1	0.82	0.79	1.12	1	0.84	0.95	1.18	1.04
Lead	0.68	1.1	1.05	1.29	0.99	1.16	0.83	1.14	1.02	0.76	1.22	0.92	1.44	1.04	1.04	0.96
Malathion dicarboxylic acid**	0.63	0.71	0.72	0.54	0.8	0.4	1.26	1.09	0.73	0.54	0.73	0.45	0.35	2.28	1.4	1.11
Mandelic acid**	0.91	1.09	1.26	1.51	0.92	0.83	0.95	0.92	1.05	0.91	1.4	1.14	0.92	0.78	1.19	0.82
MeA-alpha-C	0.23	0.58	1.29	2.1	0.6	0.27	0.86	1.22	0.56	0.29	1.76	1.06	0.73	0.23	1.55	0.64
Melamine	1.17	2.15	0.92	1.83	0.69	0.39	0.96	0.84	1.17	2.15	1.24	1	0.69	0.39	1	0.96
Methylarsonic acid	1.12	1.23	0.94	1.02	1.1	1.11	0.93	0.96	1.12	0.96	0.96	1.02	1.01	1.07	0.96	0.93
Mono-(2-ethyl-5-hydroxyhexyl) phthalate**	0.96	1.21	1.14	1.31	1.08	1.04	0.71	1.11	1.21	0.85	1.23	1.11	2.13	0.87	1	0.81
Mono-(2-ethyl-5-oxohexyl) phthalate**	1.04	1.07	1.07	1.26	1.16	0.96	0.72	1.09	1.15	0.8	1.23	0.93	2.01	0.87	1.01	0.9
Mono-(2-ethylhexyl) phthalate**	1.26	1.4	1.4	1.04	1.18	1.1	0.66	1	1.38	0.84	1.32	1.26	1.24	1	0.7	0.84
Mono-2-ethyl-5-carboxypentyl phthalate**	1.15	1.03	1.06	1.28	1.04	1.21	0.78	1.04	1.13	0.81	1.18	0.94	1.73	0.93	0.99	0.67
N'-Nitrosornicotine	0.2	0.37	0.97	1.22	0.42	1.43	0.87	1.51	0.38	0.17	1.31	0.76	0.95	0.7	1.95	0.78
N-Acetyl-S-(1-cyano-2-hydroxyethyl)-L-cysteine (CYHA)**	0.35	0.92	1.31	2.17	0.39	0.78	0.89	1.04	0.75	0.5	1.68	1.2	1.94	0.14	1.54	0.74
N-Acetyl-S-(n-propyl)-L-cysteine**	1.17	1.32	1.1	1.03	1.69	1.19	1.01	0.95	1	1.99	0.87	1.14	1.2	1.28	0.77	0.98
N-Acetyl-S-(phenyl)-L-cysteine**	0.92	1.33	0.82	0.83	0.94	1.41	0.93	1.03	1.35	0.92	0.81	0.82	1.05	1.31	0.88	1.03
N-acetyl-S-(2-Carboxyethyl)-L-cysteine**	0.56	0.86	1.21	1.57	0.59	0.88	1	1.11	0.81	0.62	1.45	1.33	0.83	0.58	1.18	0.99
N-acetyl-S-(2-Hydroxyethyl)-L-cysteine**	0.85	0.9	1.6	1.58	1.28	1.01	0.94	0.81	0.95	0.69	2.08	1.33	1.51	1.01	1.05	0.73
N-acetyl-S-(2-carbamoyl-ethyl)-L-cysteine**	0.68	1.19	1.07	1.89	0.83	1.09	0.77	1.07	0.85	0.99	1.77	1.27	1.64	0.95	1.27	0.86
N-acetyl-S-(2-cyanoethyl)-L-cysteine**	0.36	0.92	1.47	2.06	0.42	0.77	0.84	1.22	0.62	0.47	2.06	1.52	1.22	0.12	1.66	0.82
N-acetyl-S-(2-hydroxy-3-butenyl)-L-cysteine**	0.37	1.16	0.7	2.26	0.47	0.59	0.99	1.02	0.94	0.51	1.6	1.15	1.09		1.46	0.75
N-acetyl-S-(2-hydroxypropyl)-L-cysteine**	0.66	0.8	0.89	1.43	0.89	0.69	1.16	1.14	0.89	0.57	0.89	1.04	1.43	0.79	1.31	0.88
N-acetyl-S-(3,4-dihydroxy butyl)-L-cysteine**	0.83	0.99	0.91	1.07	0.79	0.99	1.06	1	0.96	0.85	1.26	0.88	0.91	0.99	1.08	0.99
N-acetyl-S-(3-Hydroxypropyl)-L-cysteine**	0.56	0.74	1.17	1.75	0.79	0.79	1.12	1	0.72	0.68	1.67	1.17	0.98	0.67	1.28	1.05
N-acetyl-S-(N-methylcarbamoyl)-L-cysteine**	0.64	0.81	1.16	1.35	0.57	1.14	1.01	1.16	0.71	0.64	1.21	1	0.78	0.89	1.32	0.93
N-acetyl-s-(3-hydroxypropyl-1-methyl)-L-cysteine**	0.5	0.72	1.02	1.79	0.93	0.83	1.11	1.16	0.79	0.47	1.7	0.87	1	0.59	1.43	0.93
NNAL**	0.13	0.42	1.06	1.18	0.32	1.07	0.81	1.39	0.36	0.16	1.18	0.99	1	0.54	1.75	0.76
Nickel	0.83	0.94	1.13	1.03	0.97	1.08	0.96	1.04	1.03	0.78	1.19	1	1.1	0.97	1.13	0.8
Nitromethane	0.95	0.95	1.29	1.12	1.08	1.04	0.95	0.93	1.01	0.87	1.25	1.13	1.21	0.96	0.96	0.94
Oxypyrimidine**	0.96	1.06	1.67	1.91	0.86	0.94	0.82	0.89	0.96	1.16	1.98	1.55	0.88	0.86	0.9	0.79
Pentachlorophenol	0.64	0.81	1.23	1.07	0.77	1.54	0.89	1.36	0.81	0.68	1.15	1.12	0.73	0.34	0.97	1.04
Dem. Subgroup:																
Inc/PL ratio:	Hsp	Hsp	Blk	Blk	Oth	Oth	Wht	Wht	Hsp	Hsp	Blk	Blk	Oth	Oth	Wht	Wht
Race/eth:	F	M	F	M	F	M	F	M	F	M	F	M	F	M	F	M
Sex:																

Continued, next page...

Table A12: NHANES P95 biomarker concentration ratios: age-differentiated results

Chemical																									
1,4-Dichlorobenzene	1.13	1	<u>1.56</u>	1.29	1.63	3.86	0.28	0.72	0.45	0.6															
1-Naphthylamine	0.22	1.09	0.2	0.52	0.31	<u>1.57</u>	0.45	0.66	0.22	1.14															
1-naphthol**	0.32	1.24	0.37	0.85	0.71	<u>1.54</u>	0.52	0.67	0.22	1.49															
2,4,5-trichlorophenol**	0.67	1	0.67	0.67	0.67	0.67	0.67	1	1	1															
2,4,6-Trichlorophenol	1.18	1	1.18	1.09	1.27	1.36	0.73	0.91	1.27	0.82															
2,4-Dichlorophenoxyacetic acid	1.19	0.92	1.21	0.8	0.94		0.99	0.82	1.25	0.97															
2,4-dichlorophenol**	1.26	0.96	2.1	2.36	4.95	10.51	0.58	0.73	0.7	0.48															
2,5-dichlorophenol**	1.43	0.94	<u>3.28</u>	<u>3.62</u>	<u>12.5</u>	<u>13.67</u>	0.22	0.39	0.5	0.18															
2,6-Dimethylaniline	0.78	1.04	0.3	1	0.28	1.14	0.21	0.83	3.04	1.12															
2-Amino-1-methyl-6-phenylimidazo [4,5-b]pyridine	0.49	1.09	0.52	0.99	0.72	1.23	0.49	0.91	0.39	1.16															
2-Amino-alpha-carboline	0.14	1.12	0.09	0.55	0.14	<u>1.51</u>	0.52	0.45	0.34	1.12															
2-Anisidine	0.72	1.05	0.65	0.99	0.64		0.66	1.04	0.89	1.07															
2-Methylaniline	1.08	0.93	0.83	0.88	1.26	1.2	0.88	0.93	1.29	0.87															
2-Naphthylamine	0.56	1.14	0.52	0.56	0.75	<u>1.52</u>	0.46	0.9	0.51	1.21															
2-Phenylphenol	1.2	1	1	1	<u>1.6</u>	1.2	0.6	0.4	1.2	1															
2-naphthol**	0.9	1.04	0.99	1.15	1.12	1.47	0.93	0.68	0.72	0.96															
4,4'-Diaminobiphenyl methane	1.17	0.97	1.25	0.9	1.45		1.26	0.89	0.99	0.97															
4-Biphenylamine	0.52	1.09	0.47	0.65	0.68	<u>1.7</u>	0.55	1	0.41	1.16															
Acrylamide	0.49	1.08	0.5	0.93	0.67	<u>1.73</u>	0.53	0.79	0.43	1.05															
Arsenic	0.64	1.12	0.56	1.24	1.3	1.43	1.27	1.82	0.45	0.85															
Arsenic acid	1.12	0.99	1.24	1.06			0.95	1.05	1.05	0.94															
Benzene	0.18	1.07	0.17	0.51	0.22	1.38	0.14	1.45	0.18	1.12															
Bis(2-chloroethyl) phosphate**	1.35	0.92	1.27	0.75	1.23	0.73	1.25	1.16	1.88	0.97															
Bromodichloromethane	1	1	0.73	1.09	0.91	1.09	1	1.09	1.27	0.91															
Bromoform	1	1	0.9	1.3		0.8	1	1	1	1															
Cadmium	0.21	1.12	0.2	0.73	0.22	1.29	0.29	1.3	0.2	1.16															
Chloroform	0.94	1	0.71	0.94	1.06	1.24	1	1.26	1.26	1															
Cobalt		1		0.95		0.86		0.86		1.05															
Crotonaldehyde	0.84	1.02	1.19	1.46	0.76	1.04	1.27	1.22	0.8	0.98															
Dieldrin	0.48	1.03	0.47	0.89	0.42	0.84	0.45	1.37	0.54	1.05															
Diisononyl phthalate	1	1	1.33	1.33	1.22	1.06	0.78	0.72	1	0.94															
Dimethyl dithiophosphate**	1.24	0.88	0.89	0.58	<u>3.11</u>	0.62	1.19	0.43	1.42	1.27															
Dimethyl phosphate**	1.45	0.93	1.32	0.89	<u>1.72</u>	1.14	1.77	0.47	1.33	0.96															
Dimethyl thiophosphate**	1.28	0.81	0.85	0.73	<u>1.86</u>	0.67	1.15	0.44	1.19	1.03															
Dimethylarsinic acid	0.9	1.05	0.96	1.16	0.69	1.28	1.16	<u>1.76</u>	0.77	0.79															
Ethylbenzene	0.31	1.08	0.2	0.63	0.3	1.14	0.38	1.25	0.33	1.1															
Dem. Subgroup:																									
Age:	<19	20+	Hispanic	<19	NH Black	20+	<19	NH Other	20+	<19	NH Other	20+	<19	NH White	<19	NH White	20+	<19	NH White	20+	<19	NH White	20+	<19	NH White
Race/eth:																									

Continued, next page...

Table A12: NHANES P95 biomarker concentration ratios: age-differentiated results

Chemical	0.17	1.12	0.17	0.71	0.27	1.36	0.18	0.84	0.14	1.12
Ethylene oxide	1.02	0.95	0.88	1.16	1.42	1.81	0.97	3.09	1.02	
Ethylene thiourea	0.99	1.01	1.01	1.03	0.99		0.29	0.99	0.99	1
Formaldehyde		1.13		0.51		1.2		1.49		1.15
Furan										
Heptachlor epoxide B		1.09		0.84		1.01		0.88		1.13
Lead	0.47	1.07	0.47	1	0.49	1.24	0.56	1.28	0.46	1.02
Malathion dicarboxylic acid**	0.83	1.07	0.62	0.67	0.65	0.65	1.64	0.8	0.9	1.14
Mandelic acid**	0.71	1.06	0.64	1.26	1	1.42	0.81	0.92	0.68	1.01
MeA-alpha-C	0.17	1.13		0.55	0.25	1.76	0.48	0.55	0.19	1.15
Melamine	1.11	0.97	2.15	1.17	1	1.39	0.69	0.39	1.11	0.96
Methylarsonic acid	1.01	0.99	0.9	1.14	0.96	0.99	1.07	1.17	1.1	0.93
Mono-(2-ethyl-5-hydroxyhexyl) phthalate**	1.24	0.94	1.43	0.95	1.39	1.11	1.55	0.88	1.15	0.81
Mono-(2-ethyl-5-oxohexyl) phthalate**	1.34	0.91	1.44	1.04	1.23	1.07	1.64	0.9	1.28	0.8
Mono-(2-ethylhexyl) phthalate**	1.04	0.98	1.3	1.3	1.42	1.26	1.42	1.1	0.82	0.84
Mono-2-ethyl-5-carboxypentyl phthalate**	1.3	0.85	1.34	0.97	1.21	1.15	1.72	1.01	1.23	0.63
N'-Nitrosomnicotine	0.16	1.13		0.4	0.16	1.27		1.14	0.22	1.26
N-Acetyl-S-(1-cyano-2-hydroxyethyl)-L-cysteine (CYHA)**		1.23		0.94	0.35	1.98		0.78		1.16
N-Acetyl-S-(n-propyl)-L-cysteine**	0.66	1.12	0.78	1.41	0.7	1.14	0.85	1.87	0.55	0.98
N-Acetyl-S-(phenyl)-L-cysteine**	1.21	0.93	1.3	1.2	0.96	0.81	1.21	1.05	1.25	0.91
N-acetyl-S-(2-Carboxyethyl)-L-cysteine**	0.58	1.07	0.52	0.85	0.95	1.58	0.6	0.73	0.5	1.11
N-acetyl-S-(2-Hydroxyethyl)-L-cysteine**	0.78	1.11	0.78	0.92	0.94	1.9	0.66	1.28	0.69	0.94
N-acetyl-S-(2-carbamoyl-ethyl)-L-cysteine**	0.76	1.05	0.64	1.04	1.07	1.65	0.75	1.09	0.67	0.98
N-acetyl-S-(2-cyanoethyl)-L-cysteine**	0.07	1.23	0.05	0.93	0.24	2.06	0.07	0.94	0.03	1.19
N-acetyl-S-(2-hydroxy-3-butenyl)-L-cysteine**		1.28		1.16		1.98		0.73		1.28
N-acetyl-S-(2-hydroxypropyl)-L-cysteine**	0.47	1.16	0.63	0.8	0.73	1.01	0.44	1.19	0.33	1.29
N-acetyl-S-(3,4-dihydroxybutyl)-L-cysteine**	0.93	1.01	0.83	0.93	1.11	1	0.87	0.92	0.92	1.05
N-acetyl-S-(3-Hydroxypropyl)-L-cysteine**	0.55	1.13	0.47	0.74	0.64	1.96	0.78	0.92	0.49	1.24
N-acetyl-S-(N-methylcarbamoyl)-L-cysteine**	0.27	1.18	0.24	0.81	0.34	1.33	0.29	0.89	0.27	1.26
N-acetyl-s-(3-hydroxypropyl-1-methyl)-L-cysteine**	0.35	1.16	0.33	0.8	0.39	1.8	1.21	0.83	0.3	1.27
NNAL**	0.11	1.15	0.03	0.43	0.13	1.23	0.06	1.02	0.17	1.31
Nickel	1.17	0.93	1.22	0.78	1.41	1.06	1.24	0.95	1.11	0.95
Nitromethane	0.7	1.02	0.77	1.01	0.76	1.26	0.72	1.07	0.6	0.95
Oxypyrimidine**	0.71	1.06	0.6	1.17	1.12	1.97	0.86	0.86	0.51	0.89
Pentachlorophenol	1.23	0.94	0.81	0.8	1.14	1.12	1.28	0.77	1.42	0.95
Perfluorooctanesulfonic acid	0.5	1.03	0.42	0.71	0.49	1.54	0.5	1.42	0.65	0.98
Perfluorooctanoic acid	0.63	1.03	0.76	0.79	0.44	0.93	0.76	1.53	0.63	1.05
Phenylglyoxylic acid**	0.75	1.08	0.65	0.93	0.94	1.35	0.8	0.82	0.7	1.08
Dem. Subgroup:										
Age:	<19	20+	Hispanic	20+	<19	20+	<19	20+	<19	20+
Race/eth:			Hispanic	Hispanic	NH Black	NH Black	NH Other	NH Other	NH White	NH White

Continued, next page...

Table A12: NHANES P95 biomarker concentration ratios: age-differentiated results

Chemical										
Styrene Tetrachloroethylene	0.61	1.02	0.49	0.65	0.36	1.33	0.41	0.73	0.72	1.16
	0.78	1.03		1.07		1.13			1.15	1.07
Dem. Subgroup:										
Age:	<19	20+		Hispanic	20+	<19	NH Black	20+	<19	20+
Race/eth:			Hispanic		NH Black		NH Other		NH White	NH White

Notes: Age-based analysis showing 95th percentile ratios for the carcinogens and associated metabolites (N = 42) for which censoring by detection limits did not prevent statistical calculations. Demographic subgroups presented by age, including age subgroups differentiated by race/ethnicity. **Bold** values denote concentration ratios ≥ 1.25 . Underlined values denote concentration ratios ≥ 1.5 . Statistics based on individual-level NHANES biomonitoring samples.

Demographic subgroups: Hispanic, non-Hispanic Black, non-Hispanic White, Other race including multi-racial

Parent Chems: {1-naphthol: *1-Naphthalenol*, *1-(N-methylcarbamate)***}; {1-naphthol, 2-naphthol: *Naphthalene***}; {2,4-dichlorophenol: *2,4-Dichlorophenoxyacetic acid***}; {2,4-dichlorophenol: *Nitrofen***}; {2,5-dichlorophenol: *1,4-Dichlorobenzene***}; {Bis(2-chloroethyl) phosphate: *Tris(2-chloroethyl) phosphate***}; {Dimethyl phosphate: *Dichlorvos***}; {Dimethyl phosphate: *Malathion***}; {Dimethyl phosphate: *Z-Tetrachlorvinphos***}; {Dimethyl dithiophosphate: *Malathion***}; {Dimethyl phosphate: *Malathion***}; {Malathion dicarboxylic acid: *Malathion***}; {Mandelic acid: *Styrene***}; {Mandelic acid: *Styrene***}; {Mono-(2-ethyl-5-hydroxyhexyl) phthalate, Mono-(2-ethyl-5-oxohexyl) phthalate, Mono-(2-ethyl-5-carboxypentyl) phthalate: *Di(2-ethylhexyl) phthalate***}; {N-Acetyl-S-(1-cyano-2-hydroxyethyl)-L-cysteine (CYHA), N-acetyl-S-(2-Hydroxyethyl)-L-cysteine, N-acetyl-S-(2-cyanoethyl)-L-cysteine, N-acetyl-S-(3-Hydroxypropyl)-L-cysteine: *Acrylonitrile***}; {N-Acetyl-S-(n-propyl)-L-cysteine: *1-Bromopropane***}; {N-Acetyl-S-(phenyl)-L-cysteine: *Benzene***}; {N-Acetyl-S-(2-Carboxyethyl)-L-cysteine, N-acetyl-S-(2-Hydroxyethyl)-L-cysteine: *Ethylene Oxide***}; {N-Acetyl-S-(2-carbamoyl)-L-cysteine: *Acrylamide***}; {N-Acetyl-S-(2-hydroxypropyl)-L-cysteine: *1,2-Propylene oxide***}; {N-Acetyl-S-(2-hydroxy-3-butenyl)-L-cysteine, N-acetyl-S-(3,4-dihydroxybutyl)-L-cysteine: *1,3-Butadiene***}; {N-Acetyl-S-(N-methylcarbamoyl)-L-cysteine: *1,2-Propylene oxide***}; {N-Acetyl-S-(3-hydroxypropyl-1-methyl)-L-cysteine: *Crotonaldehyde***}; {NNAL: *4-(N-Methyl-N-nitrosamino)-1-(3-pyridyl)-1-butanone***}; {Oxypyrimidine: *Diazepam***}; {Phenylglyoxylic acid: *Ethylbenzene***}; {Phenylglyoxylic acid: *Ethylbenzene***}

Return to: Section 3.1.3.

Table A13: NHANES detected sample ratios: primary results

Chemical		0.41	1.26	0.37	1.16	1.27	1.12	1.18	0.81
1,1,1-Trichloroethane		0.41	1.26	0.37	1.16	1.27	1.12	1.18	0.81
1,1,2,2-Tetrachloroethane		0.72	1.49	0	1.24	0.98	1.87	0.67	1.33
1,4-Dichlorobenzene		1.1	0.87	1	1.53	0.74	0.85	0.99	1.01
1-Naphthylamine		1.11	0.89	0.95	1.22	0.98	0.84	0.96	1.04
1-naphthol**		1	1	1	1	1	1	1	1
2,4,5-trichlorophenol**		0.99	0.96	0.74	1.24	1.06	1.16	1	1
2,4,6-Trichlorophenol		1.1	0.83	1.02	1.41	0.83	0.83	1	1
2,4-Dichlorophenoxyacetic acid		1	1.01	0.98	1.06	1.02	0.9	0.95	1.05
2,4-dichlorophenol**		1.01	0.99	1.02	1.05	0.96	0.95	1	1
2,5-dichlorophenol**		1.01	0.99	1.01	1.02	0.99	0.97	0.99	1.01
2,6-Dimethylaniline		1.03	0.97	1	1.1	1	0.84	0.96	1.05
PhIP		0.99	1.02	1.1	1.21	0.89	0.79	0.96	1.04
IQ (heterocyclic amine)		1.05	0.85	0.99	0.96	1.03	1	1.02	0.98
2-Amino-alpha-carboline		1.06	0.93	0.98	1.04	1.06	0.84	0.91	1.09
2-Anisidine		1	1.01	1	1	1	1	1	1
2-Methylaniline		1.02	0.98	1	1.08	0.98	0.93	0.97	1.03
2-Naphthylamine		1.05	0.95	0.95	1.18	0.98	0.89	0.96	1.04
2-Phenylphenol		1.08	0.92	0.94	1.17	1.01	0.68	1.01	0.99
2-naphthol**		1	1	1	1	1	1	1	1
4,4'-Diaminobiphenyl methane		1.02	0.96	1.04	1.12	0.99	0.78	0.97	1.03
4-Biphenylamine		1.03	0.97	0.95	1.07	1.02	0.93	0.98	1.02
4-Methyl-2-pentanone		0.96	1.22	0.84	0.9	1.62	0.25	0.88	1.13
Acetochlor mercapturate**		1.01	0.94	0.86	1.71	0.64	1.5	1.04	0.95
Acrylamide		1	1	1	1	1	1	1	1
Alachlor mercapturate**		1.03	1.34	0.73	0.79	1.7	0	1.17	0.82
Arsenic		1	1	1	1	1	1	1	1
Arsenic acid		1.02	1.07	1.38	0.67	1.01	0.91	1	1
Benzene		1.18	0.82	0.81	1.31	1.04	0.8	0.89	1.12
Beta-Hexachlorocyclohexan†				1.09	1.42	0.97	0.97	1	1
Dem. Subgroup:									
Inc/PL ratio:		<2x P.L.	>2x P.L.	Hispanic	NH Asian	NH Black	NH White	Other	
Race/eth:									
Sex:								Female	Male

Continued, next page...

Table A13: NHANES detected sample ratios: primary results

Chemical									
Bis(2-chloroethyl) phosphate**	1.02	0.98	1	1.04	0.99	0.96	0.99	1.01	1.01
Bromodichloromethane	0.9	1.11	0.89	1.13	0.92	1.13	1	0.99	0.99
Bromoform	1.1	0.93	1.43	0.57	1.02	0.93	0.95	1.06	1.06
Cadmium	0.95	1.04	0.94	1	1	1.08	1.02	0.97	0.97
Carbon tetrachloride	0.8	1.48	1.4	1.51	0.98	0	0.65	1.38	1.38
Chloroethane	1.58	0.74	0	1.51	1.93	0	0	2.05	2.05
Chloroform	0.94	1.04	0.9	1.21	0.89	1.06	1.03	0.97	0.97
Cobalt	1	1	1	1	1	1	1	1	1
Crotonaldehyde	1.07	0.9	1.13	1.05	0.85	1.15	1	1	1
Cumene	0.89	0.84	1.03	0	1.83	0.64	0.49	1.55	1.55
DDT†			0.92	0.87	1.05	0.98	1.02	0.98	0.98
				0.99					
Dichloromethane	0.8	1.48	1.4	0	1.47	0.84	0.97	1.03	1.03
Dieldrin	0.85	1.13	0.85	0.8	1.19	1.16	0.95	1.05	1.05
Diisononyl phthalate	1.05	0.83	1.15	1.2	0.86	0.82	1	1	1
Dimethyl dithiophosphate**	0.99	1.04	1.04	0.98	1.03	0.93	0.98	1.02	1.02
Dimethyl phosphate**	0.99	1.01	1.01	1	0.99	1.01	1	1	1
Dimethyl thiophosphate**	0.99	1.01	1	1	1.01	0.99	0.99	1.01	1.01
Dimethylarsinic acid	0.99	0.99	1.07	1.02	0.89	1.07	0.96	1.04	1.04
Ethylbenzene	1.19	0.85	0.61	1.4	1.11	0.84	0.83	1.18	1.18
Ethylene oxide	1.03	0.95	1	1.06	0.93	1.07	0.99	1.01	1.01
Ethylene thiourea	1.1	0.9	1.22	1.17	0.69	1.48	0.96	1.04	1.04
Formaldehyde	1	1	1	1	1	1	1	1	1
Furan	1.41	0.61	0.44	1.56	1.2	0.71	0.87	1.14	1.14
Glu-P-1	1.13	0.98	1.03	1.35	0.99	0.48	1.09	0.91	0.91
Glu-P-2	1.29	0.56	1.88	1.74	0	0.83	1.75	0.25	0.25
Heptachlor epoxide B	0.95	1.03	0.92	0.77	1.2	0.79	1.04	0.96	0.96
Lead	1	1	1	1	1	1	1	1	1
Malathion dicarboxylic acid**	0.94	1.1	0.92	0.84	1.16	0.68	0.97	1.03	1.03
Mandelic acid**	1	1	0.99	1	1	1.01	1	1	1
Dem. Subgroup:									
Inc/PL ratio:	<2x P.L.			>2x P.L.					
Race/eth:				Hispanic			NH Asian NH Black NH White Other		
Sex:							Female Male		

Continued, next page...

Table A13: NHANES detected sample ratios: primary results

Chemical		1.2	0.78	0.58	1.35	1.24	0.64	0.84	1.16
MeA-alpha-C		1.01	1.01	0.93	0.92	1.08	1.25	0.99	1.01
Melamine		0.99	1	1.11	0.92	0.85	1.19	0.92	1.09
Methylarsonic acid				0.82	1.05	1.05	0.97	0.99	1.01
Mirex†									
Mono-(2-ethyl-5-hydroxyhexyl) phthalate**		1	1	1	1	1	1	1	1
Mono-(2-ethyl-5-oxohexyl) phthalate**		1	1	1	1	1	1	1	1
Mono-(2-ethylhexyl) phthalate**		1.05	0.95	1.05	1.13	0.83	1.07	0.94	1.07
Mono-2-ethyl-5-carboxypentyl phthalate**		1	1	1	1	1	1	1	1
Mono-n-octyl phthalate**		0.9	0.86	1.12	1.24	0.84	0.82	0.95	1.05
N'-Nitrosornicotine		1.26	0.73	0.47	1.26	1.33	0.71	0.81	1.19
NAS-(1-cyano-2-hydroxyethyl)-LC (CYHA)**		1.23	0.77	0.58	1.7	1.13	0.63	0.78	1.22
NAS-(n-propyl)-LC**		0.99	1.01	1.02	1.05	0.94	1.01	0.97	1.03
NAS-(phenyl)-LC**		0.97	1.03	1.03	1.05	1.03	0.8	0.91	1.09
N-Nitroso-N-methylethylamine		1.27	0.62	1.03	1.03	0.61	1.81	0.78	1.21
N-Nitrosodiethylamine		0.84	0.89	1.09	0.95	0.83	1.29	1	1
N-Nitrosomorpholine		1.25	0.77	0.6	1.54	0.98	1	1.13	0.87
N-Nitrosopiperidine		1.24	0.79	1.27	1.25	0.66	1	0.92	1.08
N-Nitrosopyrrolidine		0.86	1.12	1.56	0.77	0.91	0.55	1.83	0.17
NAS-(2-Carboxyethyl)-LC**		1	1	1	1	1	1	1	1
NAS-(2-Hydroxyethyl)-LC**		1.03	0.96	0.96	1.25	0.96	0.81	1	1
NAS-(2-carbamoylethyl)-LC**		1	1	1	1	1	1	1	1
NAS-(2-cyanoethyl)-LC**		0.99	1.01	0.96	1.13	0.98	0.94	0.94	1.06
NAS-(2-hydroxy-3-butenyl)-LC**		1.21	0.79	0.71	1.43	1.21	0.59	0.81	1.19
NAS-(2-hydroxypropyl)-LC**		1	1.01	1	1.02	1	0.97	0.98	1.02
NAS-(3,4-dihydroxybutyl)-LC**		1	1	1	1	1	1	1	1
NAS-(3-Hydroxypropyl)-LC**		1	1	1	1	1	1	1	1
NAS-(N-methylcarbamoyl)-LC**		1	1	1	1	1	0.99	1	1
NAS-(3-hydroxypropyl-1-methyl)-LC**		1	1	1	1	1	1	1	1
NNAL**		1.17	0.81	0.95	1.21	0.96	0.86	0.94	1.06
Dem. Subgroup:									
Inc/PL ratio:		<2x P.L.	>2x P.L.	Hispanic	NH Asian	NH Black	NH White	Other	
Race/eth:									
Sex:								Female	Male

Continued, next page...

Table A13: NHANES detected sample ratios: primary results

Chemical										
Nickel	1	1	1	1	1.02	1	0.98	0.99	1.01	
Nitromethane	1	1	1	1	1	1	1	1	1	
Oxypyrimidine**	1.03	0.92	0.85		1.56	0.83	0.85	0.98	1.02	
PBB153†			0.97	0.82	1.01	1.03	0.93	1	1	
PCB028†			0.78	1.33	0.7	1.1	0.78	1.13	0.86	
PCB066†			0.49	2.11	0.9	1.06	0.93	1	1	
PCB074†			0.96	1.04	0.99	1.01	0.98	0.98	1.02	
PCB081†			1.08	3.99	0	0.97	0	1.66	0.3	
PCB105†			0.97	1.03	0.97	1.01	0.93	0.99	1.01	
PCB114†			0.68	1.06	0.9	1.09	1.16	0.97	1.03	
PCB126†			1	1.28	0.89	1.02	0.68	1	1	
PCB146†			0.87	1.06	1.03	1.02	1.06	0.98	1.02	
PCB156†			0.93	1.04	1	1.01	1.04	0.97	1.03	
PCB157†			0.7	1.21	0.9	1.07	1.05	0.92	1.08	
PCB167†			0.87	1.41	0.92	1.03	0.75	0.98	1.02	
PCB169†			0.82	1.21	0.84	1.08	0.65	0.94	1.06	
PCB170†			1	1.01	0.99	1.01	0.96	1	1	
PCB178†			0.79	1.18	0.95	1.05	0.92	0.94	1.06	
PCB183†			0.94	1.04	0.99	1.02	0.89	0.98	1.02	
PCB187†			0.98	1.04	1.04	0.99	1.04	0.98	1.02	
PCB189†			0.54	1.04	0.9	1.12	1.24	1.05	0.94	
PCB194†			0.95	1.05	0.98	1.02	0.95	0.98	1.02	
PCB196†			0.96	1.03	0.99	1.01	1.03	0.97	1.03	
PCB199†			0.88	1.04	0.97	1.03	1.06	0.99	1.02	
PCB206†			0.82	1.15	1.03	1.03	1.02	0.92	1.08	
PCB209†			0.89	1.27	1.01	1	1.03	0.87	1.14	
PeCDF (furan)†			0.99	1	0.86	1.05	0.72	0.92	1.08	
Pentachlorophenol	1.03	0.97	0.75		1.22	1.04	0.85	0.98	1.02	
Perfluorooctanesulfonic acid	1	1	1		0.99	1	1	1	1	
Dem. Subgroup:										
Inc/PL ratio:	<2x P.L.			>2x P.L.						
Race/eth:				Hispanic	NH Asian	NH Black	NH White	Other	Female	Male
Sex:										

Continued, next page...

Table A14: NHANES detected sample ratios: demographic interactions

Chemical		0.69	0	1.46	0.81	0.85	1.4	1.55	1.01	0	0.53	0	1.95	1.36	0	3.44	0.59	2.05
1,1,1-Trichloroethane		0.69	0	1.46	0.81	0.85	1.4	1.55	1.01	0	0.53	0	1.95	1.36	0	3.44	0.59	2.05
1,1,2,2-Tetrachloroethane		0	0	0	2.57	3.9	0	0	1.93	0	0	0	3.23	0	0	2.25	0	0
1,4-Dichlorobenzene		0.95	1.05	1.57	1.49	0.87	0.83	0.71	0.78	1.16	0.75	1.68	1.35	0.91	0.8	0.76	0.71	0.86
1-Naphthylamine		0.93	0.96	1.22	1.22	0.82	0.87	0.91	1.04	0.96	0.89	1.29	1.1	1.03	0.76	1.14	0.86	0.86
1-naphthol**		1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1
2,4,5-trichlorophenol**		0.8	0.69	1.23	1.25	0.97	1.3	1.04	1.08	0.7	0.64	1.25	1.22	1.34	0.76	1.07	1.01	1.01
2,4,6-Trichlorophenol		0.97	1.07	1.49	1.34	0.7	0.93	0.86	0.81	1	0.91	1.61	1.11	0.97	0.51	0.93	0.75	0.75
2,4-Dichlorophenoxyacetic acid		0.94	1.01	1.02	1.09	0.79	1.01	0.97	1.07	0.98	0.98	1.05	1.06	0.92	0.89	1	1.05	1.05
2,4-dichlorophenol**		1.02	1.02	1.04	1.06	0.95	0.96	0.95	0.98	1.03	0.99	1.06	1.03	0.94	1	0.97	0.96	0.96
2,5-dichlorophenol**		1	1.02	1.02	1.02	0.98	0.97	0.97	1	1.02	0.99	1.03	1.02	0.99	0.96	1	0.98	0.98
2,6-Dimethylaniline		0.92	1.1	1.09	1.11	0.89	0.79	0.94	1.07	0.98	1.1	1.11	1.06	0.84	0.82	1.07	0.96	0.96
PhIP		1.07	1.13	1.23	1.19	0.79	0.79	0.81	0.98	1.1	1.12	1.15	1.31	0.86	0.76	0.81	0.98	0.98
IQ (heterocyclic amine)		1.2	0.76	1.27	0.67	0.74	1.28	0.85	1.19	1.1	0.46	1.06	0.71	0.84	1.19	1.05	0.89	0.89
2-Amino-alpha-carboline		0.83	1.14	0.94	1.14	0.78	0.91	1.01	1.1	0.97	0.94	1.07	0.99	1.02	0.77	1.16	0.97	0.97
2-Anisidine		1	1	1	1	1.01	0.98	1	1.01	1	1	0.99	1.01	0.99	1	1	1.01	1.01
2-Methylaniline		0.97	1.03	1.07	1.1	0.93	0.93	0.95	1.02	0.98	1.01	1.1	1.05	1.01	0.87	1	0.98	0.98
2-Naphthylamine		0.93	0.96	1.14	1.21	0.84	0.95	0.93	1.02	0.95	0.93	1.23	1.1	0.9	0.89	1.06	0.92	0.92
2-Phenylphenol		1	0.9	1.17	1.16	0.48	0.82	1.01	1.02	0.95	0.96	1.32	1.06	0.75	0.47	1.13	0.9	0.9
2-naphthol**		1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1
4,4'-Diaminobiphenyl methane		1	1.09	1.12	1.11	0.73	0.83	0.94	1.03	1.03	1.03	1.17	1.07	0.82	0.72	0.96	1	1
4-Biphenylamine		0.93	0.97	1.05	1.09	0.96	0.9	0.99	1.05	0.93	0.96	1.11	1.04	0.91	0.95	1.1	0.96	0.96
4-Methyl-2-pentaone		0.8	0.89	0.43	1.43	0	0.52	1.78	1.45	1.33	0.6	0.52	1.7	0.79	0	1.03	2.07	2.07
Acetochlor mercaptate**		0.83	0.88	2.05	1.37	0.67	2.54	0.67	0.61	0.8	1.12	1.85	1.13	2.14	0.9	0.12	0.82	0.82
Acrylamide		1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1
Alachlor mercaptate**		0.48	1.01	1.48	0	0	0	2.13	1.3	0.82	1.06	0.77	1.34	0	0	2.27	1.62	1.62
Arsenic		1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1
Arsenic acid		1.27	1.51	0.64	0.7	0.99	0.82	1.03	0.98	1.59	1.13	0.7	0.76	0.77	1.07	0.89	1.19	1.19
Benzene		0.74	0.89	1.12	1.53	0.68	0.93	0.96	1.12	0.87	0.74	1.41	1.14	1.11	0.62	1.29	0.82	0.82
Beta-Hexachlorocyclohexanef		1.16	1.02	0.79	0.86	1.14	0.8	0.95	1	1	1	1	1	1	1	1	1	1
Bis(2-chloroethyl) phosphate**		0.99	1.01	1.04	1.05	0.91	1.02	0.99	0.98	1.01	0.98	1.03	1.07	0.99	0.97	1.04	0.96	0.96
Bromodichloromethane		0.85	0.92	1.2	1.06	1.15	1.11	0.89	0.94	0.88	1.04	0.98	1.4	0.91	1.19	0.86	0.93	0.93
Bromoform		1.48	1.37	0.52	0.63	0.78	1.09	0.94	1.1	1.46	1.59	0.51	0.65	0.91	0.81	1.29	0.83	0.83
Cadmium		0.96	0.91	1.03	0.97	1.12	1.04	1.01	0.99	0.9	0.99	0.94	1.05	1.05	1.09	0.96	1.03	1.03
Carbon tetrachloride		0	3	2.84	0	0	0	0	1.94	0	4.02	3.51	0	0	0	0	1.96	1.96
Chloroethane		0	0	0	3.19	0	0	0	3.82	0	0	0	3.83	0	0	4.47	0	0
Dem. Subgroup:																		
Inc/PL ratio:																		
Race/eth:																		
Sex:																		

Continued, next page...

Table A14: NHANES detected sample ratios: demographic interactions

Chemical																
Chloroform	0.89	0.92	1.3	1.12	1.07	1.06	0.91	0.87	0.87	0.93	1.25	1.2	0.96	1.13	0.8	0.96
Cobalt	1.01	1	1	1	1.01	1	1	0.99	1.01	1	1.01	0.99	1	1	0.99	1
Crotonaldehyde	1.15	1.11	0.99	1.1	1.11	1.2	0.87	0.84	1.19	1.03	1.1	0.95	1.37	1.07	0.93	0.77
Cumene	0.97	1.09	0	0	0	1.33	0.74	2.89	0	1.46	0	0	1.97	0	1.71	1.47
DDT†	0.89	0.96	1.03	0.95	1.03	0.94	1.09	1								
Dichloromethane	1.33	1.48	0	0	0	1.72	1.98	0.97	2.96	0	0	0	0	1.53	0	2.95
Dieldrin	0.81	0.89	0.86	0.74	1.16	1.15	1.07	1.32	0.83	0.93	0.58	1.05	0.64	1.37	1.11	1.2
Diisononyl phthalate	1.19	1.09	1.29	1.12	0.69	0.96	0.84	0.88	1.23	0.71	1.22	0.97	0.71	0.87	0.94	0.77
Dimethyl dithiophosphate**	1.07	1	1.03	0.94	0.87	0.99	0.93	1.11	1.03	1.01	0.99	1.02	0.97	0.95	0.95	1.12
Dimethyl phosphate**	1	1.01	1	0.99	1.02	1.01	0.99	0.99	1.01	1.02	0.98	1.01	1.01	1.01	0.97	1.01
Dimethyl thiophosphate**	1	1.01	0.98	1.02	0.98	0.99	0.98	1.03	1	0.99	0.98	1.01	0.98	1	1	1.02
Dimethylarsinic acid	1.03	1.12	1.01	1.03	1.04	1.1	0.82	0.97	1.06	1.06	1.02	0.99	0.98	1.1	0.9	0.88
Ethylbenzene	0.54	0.68	1.12	1.71	0.68	1	0.95	1.27	0.68	0.53	1.44	1.34	1.27	0.63	1.37	0.88
Ethylene oxide	1	0.99	1.05	1.08	1.06	1.07	0.9	0.95	1.03	0.92	1.09	1.04	1.07	1.07	0.98	0.88
Ethylene thiourea	1.06	1.38	1.27	1.08	1.56	1.4	0.65	0.73	1.43	0.82	1.24	1.15	1.84	1.1	0.56	0.81
Formaldehyde	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1
Furan	0.37	0.51	1.3	1.86	0.52	0.9	1.16	1.24	0.45	0.4	1.81	1.11	1.34	0.41	1.93	0.58
Glu-P-1	1.16	0.88	1.4	1.3	0.71	0.25	1.02	0.96	1.42	0.45	1.61	1.15	0	0.76	0.79	1.25
Glu-P-2	2.66	1	3.61	0	1.62	0	0	0	0.81	3.08	3.01	0	2.77	0	0	0
Heptachlor epoxide B	0.96	0.87	1.04	0.5	0.87	0.69	1.09	1.31	0.97	0.85	0.48	1.03	0.49	0.93	1.35	1.09
Lead	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1
Lindane†	0	0	0	0	0	0	2.99	0								
Malathion dicarboxylic acid**	0.89	0.96	0.9	0.77	0.86	0.47	1.08	1.25	1.03	0.85	0.99	0.64	0.48	0.89	0.9	1.32
Mandelic acid**	0.99	1	1	1.01	1.01	1	1	1	0.99	0.99	1	1.01	1.01	1.01	1	1
MeA-alpha-C	0.41	0.77	1.13	1.56	0.59	0.69	1.12	1.36	0.64	0.41	1.55	1.02	0.67	0.59	1.63	0.94
Melamine	0.89	0.96	0.98	0.87	1.19	1.32	1.05	1.11	0.94	0.93	0.94	0.92	1.32	1.17	1.1	1.06
Methylarsonic acid	0.98	1.26	0.81	1.03	1.15	1.24	0.79	0.91	1.08	1.15	0.95	0.91	1.1	1.21	0.9	0.82
Mirex†	0.83	0.81	0.92	1.2	1.1	0.83	1.04	1.06								
Mono-(2-ethyl-5-hydroxyhexyl) phthalate**	1	1	1	1	0.99	1	1	1	1	0.99	1	1.01	1	0.99	1	1
Mono-(2-ethyl-5-oxohexyl) phthalate**	1	0.99	1	1	0.99	1	1	1	1	0.99	0.99	1.01	1	0.99	1	1
Mono-(2-ethylhexyl) phthalate**	0.99	1.12	1.08	1.19	1	1.15	0.75	0.9	1.08	0.94	1.2	1.07	1.11	1.06	0.88	0.81
Mono-2-ethyl-5-carboxypentyl phthalate**	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1
Mono-n-octyl phthalate**	1.43	0.83	0.71	1.77	0	1.43	0.83	0.84	0.92	0.79	0.83	1.42	0.46	0.82	1.01	0.72
N ⁺ -Nitrosornicotine	0.35	0.6	1.03	1.48	0.47	0.98	1.18	1.49	0.49	0.33	1.43	1.01	0.96	0.6	1.97	0.83
NAS-(1-cyano-2-hydroxyethyl)-LC**	0.43	0.76	1.26	2.15	0.38	0.87	1.08	1.19	0.61	0.54	2.05	1.29	0.92	0.36	1.6	0.86
Dem. Subgroup:																
Inc/PL ratio:	Hsp	Hsp	Asn	Asn	Blk	Blk	Oth	Oth	Whit	Hsp	Blk	Blk	Oth	Oth	Whit	Whit
Race/eth:	F	M	F	M	F	M	F	M	F	Hsp	Blk	Blk	Oth	Oth	Whit	Whit
Sex:																

Continued, next page...

Table A14: NHANES detected sample ratios: demographic interactions

Chemical																
NAS-(n-propyl)-LC**	1	1.05	1.03	1.07	0.92	1.09	0.9	0.97	1.01	1.05	1.02	1.12	0.99	1.02	0.92	0.94
NAS-(phenyl)-LC**	0.97	1.1	0.96	1.15	0.64	0.96	0.95	1.11	1	1.07	1.01	1.14	0.76	0.86	1	1.05
N-Nitroso-N-methylethylamine	0.39	1.76	1.63	0.49	0.71	2.97	0.64	0.59	1.43	0.67	0.9	0.7	2.47	1.71	1.05	0
N-Nitrosodiethylamine	0	2.34	1	0.9	2.55	0	1.15	0.54	0.63	1.18	0.83	0	0	2.02	1.3	0.53
N-Nitrosomorpholine	0.84	0.33	1.57	1.51	1.35	0.64	1.01	0.95	0.57	0.66	1.64	1.31	1.99	0.5	1.4	0.73
N-Nitrosopiperidine	1.1	1.46	1.33	1.17	0.84	1.17	0.57	0.74	1.36	1.13	1.69	0.74	0.79	1.14	0.9	0.47
N-Nitrosopyrrolidine	2.93	0	1.6	0	1.08	0	1.41	0.44	0.54	3.03	0.67	1.05	1.84	0	1.03	0.89
NAS-(2-Carboxyethyl)-LC**	0.99	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1
NAS-(2-Hydroxyethyl)-LC**	0.99	0.93	1.23	1.26	0.76	0.86	0.97	0.95	0.98	0.92	1.3	1.2	0.73	0.85	1	0.92
NAS-(2-carbamoylethyl)-LC**	1	1	1	1	1	1	1	1	1	1	1	1	1	1	0.99	1
NAS-(2-cyanoethyl)-LC**	0.91	1.02	1.09	1.17	0.84	1.03	0.91	1.04	0.93	1.02	1.13	1.14	0.94	0.94	0.97	0.98
NAS-(2-hydroxy-3-butenyl)-LC**	0.51	0.96	1.02	1.84	0.28	0.88	1.32	1.11	0.72	0.85	1.63	1.19	0.97	0.22	1.76	0.86
NAS-(2-hydroxypropyl)-LC**	0.98	1.02	1.01	1.03	0.93	1.02	0.99	1.01	1	1	1.02	1.03	0.94	0.99	1	1.01
NAS-(3,4-dihydroxybutyl)-LC**	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1
NAS-(3-Hydroxypropyl)-LC**	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1
NAS-(N-methylcarbamoyl)-LC**	1	1	1	1.01	0.99	1	1	1	1	1	1.01	1	0.98	1	1	1.01
NAS-(3-hydroxypropyl-1-methyl)-LC**	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1
NNAL**	0.9	1.01	1.18	1.24	0.8	0.93	0.88	1.04	1.02	0.83	1.36	0.97	1.09	0.72	1.2	0.78
Nickel	0.98	1.02	1.02	1.02	0.98	0.99	0.97	1.02	1.01	1	1	1.02	0.98	0.98	1.01	0.99
Nitromethane	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1
Oxypyrimidine**	0.95	0.74	1.51	1.61	0.58	1.11	0.84	0.83	0.81	0.75	1.69	1.29	0.84	0.85	0.81	0.86
PBB153†	0.97	0.97	0.75	0.88	0.75	0.92	0.95	1.02	1.03							
PCB028†	0.89	0.67	1.33	1.32	0.84	0.53	1.03	0.53	1.25	0.95						
PCB066†	0.38	0.6	2.34	1.86	0.95	0.84	1.13	0.72	1.04	1.07						
PCB074†	0.91	1.01	1.04	1.04	1	0.98	1.04	0.93	0.99	1.04						
PCB081†	2.16	0	6.16	1.61	0	0	0	1.57	0.33							
PCB105†	0.97	0.98	1.03	1.03	1	0.95	1.03	0.83	0.99	1.03						
PCB114†	0.75	0.61	0.98	1.15	0.93	0.86	1.33	0.99	1.01	1.17						
PCB126†	1.02	0.97	1.23	1.33	0.94	0.83	0.84	0.53	1	1.03						
PCB146†	0.87	0.86	1.06	1.06	1.06	1.06	1.06	0.99	1.05							
PCB156†	0.9	0.97	1.04	1.04	1.04	1.04	0.99	1.04								
PCB157†	0.7	0.71	1.07	1.36	0.98	0.8	1.3	0.8	0.93	1.22						
PCB167†	0.76	0.97	1.37	1.45	0.94	0.9	0.75	1.03	1.03							
PCB169†	0.76	0.89	1.22	1.21	0.75	0.95	0.68	1.02	1.15							
PCB170†	0.99	1.01	1.01	0.97	1.01	0.9	1.01	1.01								
Dem. Subgroup:																
Inc/PL ratio:	Hsp	Hsp	Asn	Asn	Blk	Blk	Oth	Oth	Wht	Wht	<2	>2	<2	>2	<2	>2
Race/eth:	F	M	F	M	F	M	F	M	F	M	Hsp	Hsp	Oth	Oth	Wht	Wht
Sex:																

Continued, next page...

Table A14: NHANES detected sample ratios: demographic interactions

Chemical																	
PCB178†	0.66	0.92	1.12	1.24	0.83	1.09	1.13	0.7	1	1.11							
PCB183†	0.94	0.95	0.99	1.08	0.95	1.03	1.08	0.69	0.99	1.05							
PCB187†	0.97	0.99	1.04	1.04	1.04	1.04	1.04	1.04	0.97	1.02							
PCB189†	0.46	0.62	1.2	0.87	0.95	0.84	1.42	1.07	1.19	1.04							
PCB194†	0.94	0.96	1.05	1.05	0.93	1.05	0.96	0.95	0.99	1.04							
PCB196†	0.93	1	1.03	1.03	0.95	1.03	1.03	1.03	0.98	1.03							
PCB199†	0.89	0.86	1.02	1.06	0.97	0.98	1.06	1.06	1	1.06							
PCB206†	0.7	0.95	1.24	1.06	0.97	1.11	1.04	0.99	0.93	1.12							
PCB209†	0.72	1.05	1.24	1.29	0.93	1.12	1.05	1	0.84	1.17							
PeCDF (furan)†	0.99	1	0.91	1.09	0.78	0.95	0.58	0.87	0.96	1.14							
Pentachlorophenol	0.74	0.75			1.27	1.16	0.96	0.75	0.95	1.15	0.71	0.76	1.28	1.15	0.94	0.69	1.16
Perfluorooctanesulfonic acid	1	1			1	0.99	1	1	1	1	1	1	0.99	1	1	1	1
Perfluorooctanoic acid	1.01	1			0.99	0.99	1	1.01	1.01	1	1.01	1.01	0.99	1	1.01	1	1.01
Phenylglyoxylic acid**	1	1			1.01	1.01	0.99	1.01	1	1	1	1	1.01	1.01	1	1	0.99
Styrene	0.83	0.95			0.89	1.14	0.61	0.97	1.08	1.22	0.88	0.81	1.07	0.94	0.79	0.91	1.22
TCDD (dioxin)†	0.91	0.86	1	1.64	0.88	0.76	1.19	0.38	1.19	0.9							
Tetrachloroethylene	0.62	1.15			1.23	1.12	0.83	0.63	0.86	1.38	0.65	1.45	0.76	1.71	0.84	0.74	0.8
Tetrahydrofuran	0.66	0.74			0.71	0.79	2.44	0	1	1.46	1.47	0	0	0.94	1.32	1.54	1.14
Trichloroethylene	0	0.99			1.9	1.06	1.09	1.15	0.99	0.97	0.49	0	2.33	0.63	2.62	0.51	0.38
Trp-P-1	1.16	0.37			0.9	1.45	1.51	0.94	0.92	0.98	1.06	0.57	0.93	1.17	1.55	0.98	1.01
Dem. Subgroup:																	
Inc/PL ratio:	Hsp	Hsp	Asn	Asn	Blk	Blk	Oth	Oth	Whit	Whit	Hsp	Hsp	Blk	Blk	Oth	Oth	>2
Race/eth:	F	M	F	M	F	M	F	M	F	M	F	M	F	M	F	M	Whit
Sex:																	>2

Notes: Demographic interaction analysis of sample detection ratios for (N = 125) matched carcinogens and carcinogen-associated metabolites. Subgroups in this table include interactions of race/ethnicity and sex, and interactions of race/ethnicity and household income. Statistics based on both individual-level (N = 96) and pooled (N = 29) NHANES samples. **Bold** values denote concentration ratios >=1.25. Underlined values denote concentration ratios >=1.5. **: Carcinogen metabolite. †: Result based on pooled sampling data.

Demographic subgroups: Household income below 2 times the poverty line, Household income above 2 times the poverty line, Hispanic, non-Hispanic Asian, non-Hispanic Black, non-Hispanic White, Other race including multi-racial, Female sex, Male sex. Non-Hispanic Asian results presented separately only for chemicals with pooled sampling.

Parent Chems: {1-naphthol: *1-Naphthalenol*, *1-(N-methylcarbamate)***}; {1-naphthol, 2-naphthol: *Naphthalene***}; {2,4-dichlorophenol: *2,4-Dichlorophenoxyacetic acid***}; {2,4-dichlorophenol: *Nitrofen***}; {2,4,5-trichlorophenol: *beta-Hexachlorocyclohexane***}; {2,4,5-trichlorophenol: *Lindane***}; {2,4,5-trichlorophenol: *1,2,3,4,5,6-Hexachlorocyclohexane***}; {2,4,5-trichlorophenol: *Pentachlorophenol***}; {2,5-dichlorophenol: *1,4-Dichlorobenzene***}; {Bis(2-chloroethyl) phosphate: *Tris(2-chloroethyl) phosphate***}; {Dimethyl phosphate: *Dichlorvos***}; {Dimethyl phosphate: *Malathion***}; {Dimethyl phosphate: *Z-Tetrachlorophos***}; {Dimethyl dithiophosphate: *Malathion***}; {Dimethyl thiophosphate: *Malathion***}; {Malathion dicarboxylic acid: *Malathion***}; {Mandelic acid: *Styrene***}; {Mandelic acid: *Ethylbenzene***}; {Mono-(2-ethyl-5-hydroxyhexyl) phthalate, Mono-(2-ethyl-5-oxohexyl) phthalate, Mono-(2-ethyl-5-carboxypentyl) phthalate: *Di(2-ethylhexyl) phthalate***}; {N-Acetyl-S-(1-cyano-2-hydroxyethyl)-L-cysteine, N-Acetyl-S-(2-Hydroxyethyl)-L-cysteine, N-Acetyl-S-(2-cyanoethyl)-L-cysteine: *Acrylonitrile***}; {N-Acetyl-S-(n-propyl)-L-cysteine: *1-Bromopropane***}; {N-Acetyl-S-(phenyl)-L-cysteine: *Benzene***}; {N-Acetyl-S-(2-Hydroxyethyl)-L-cysteine, N-Acetyl-S-(3-Hydroxypropyl)-L-cysteine: *Acrolein***}; {N-Acetyl-S-(2-Hydroxyethyl)-L-cysteine: *Vinyl Chloride***}; {N-Acetyl-S-(2-Hydroxy-3-butenyl)-L-cysteine, N-Acetyl-S-(2-carbamoyl-ethyl)-L-cysteine: *Acrylamide***}; {N-Acetyl-S-(2-Hydroxyethyl)-L-cysteine: *1,2-Propylene oxide***}; {N-Acetyl-S-(2-Hydroxy-3-butenyl)-L-cysteine, N-Acetyl-S-(3,4-dihydroxybutyl)-L-cysteine: *1,3-Butadiene***}; {N-Acetyl-S-(N-methylcarbamoyl)-L-cysteine: *N,N-Dimethylformamide***}; {N-Acetyl-S-(3-hydroxypropyl-1-methyl)-L-cysteine: *Crotonaldehyde***}; {NNAL: *4-(N-Methyl-N-nitrosamino)-1-(3-pyridyl)-1-butanone***}; {Oxypyrimidine: *Diazonon***}; {Phenylglyoxylic acid: *Styrene***}; {Phenylglyoxylic acid: *Ethylbenzene***}

Return to: Section 3.1.4.

Table A15: NHANES detected sample ratios: age-differentiated results

Chemical												
1,1,1-Trichloroethane	0.27	1.14	0	0.46		1.34	1.12	0	1.35	0	1.48	
1,1,2,2-Tetrachloroethane	0	1.23	0	0		0	1.59	0	2.33	0	1.12	
1,4-Dichlorobenzene	0.88	1.02	0.88	1.03		1.45	1.55	0.72	0.88	0.63	0.76	
1-Naphthylamine	1.01	0.99	0.97	0.93		1.24	1.21	0.89	0.82	0.93	0.99	
1-naphthol**	1	1	1	1		1	1	0.99	1	1	1	
2,4,5-trichlorophenol**	0.99	1	0.81	0.7		1.33	1.19	1.02	1.24	1	1.08	
2,4,6-Trichlorophenol	1.09	0.96	1.07	0.99		1.4	1.42	0.94	0.77	0.94	0.8	
2,4-Dichlorophenoxyacetic acid	1.08	0.96	1.04	0.93		1.09	1.03	1.08	0.82	1.13	0.99	
2,4-dichlorophenol**	1.02	0.99	1.02	1.02		1.07	1.04	0.99	0.93	1	0.95	
2,5-dichlorophenol**	1	1	1.01	1.01		1.02	1.02	0.97	0.97	0.99	0.99	
2,6-Dimethylaniline	0.97	1.01	0.98	1.02		1.03	1.14	0.89	0.81	0.94	1.02	
PhIP	0.97	1.01	1.05	1.13		1.1	1.27	0.77	0.8	0.84	0.91	
IQ (heterocyclic amine)	0.89	1.05	0.54	1.29		0.92	0.99	0.69	1.18	1.52	0.9	
2-Amino-alpha-carboline	0.84	1.07	0.86	1.05		0.83	1.18	0.83	0.85	0.86	1.11	
2-Anisidine	1	1	1.01	0.99		1	1.01	1	0.99	1	1	
2-Methylaniline	1.02	0.99	1.02	0.98		1.06	1.09	1	0.88	0.97	0.99	
2-Naphthylamine	0.99	1	0.95	0.94		1.15	1.19	0.93	0.87	0.93	0.99	
2-Phenylphenol	1.26	0.89	1.06	0.87		1.51	0.99	1.07	0.45	1.38	0.91	
2-naphthol**	1	1	1	1		1	1	1	1	1	1	
4,4'-Diaminobiphenyl methane	1.15	0.9	1.19	0.94		1.3	0.97	0.92	0.69	1.12	0.92	
4-Biphenylamine	1.03	0.99	0.99	0.92		1.07	1.07	0.98	0.91	1.08	1.01	
4-Methyl-2-pentanone	0	1.2	0	1.05		0	1.05	0	0.31	0	1.89	
Acetochlor mercapturate**	0.93	1.06	0.55	1.22		1.91	1.41	0	2.94	0.44	0.73	
Acrylamide	1	1	1	1		1	1	1	1	1	1	
Alachlor mercapturate**	0.86	1.19	0.75	0.7		0.59	1.2	0	0	1.69	1.7	
Arsenic	1	1	1	1		1	1	1	1	1	1	
Arsenic acid	1.06	0.96	1.57	1.24		0.65	0.68	0.93	0.89	1.01	1	
Benzene	0.6	1.08	0.58	0.87		0.74	1.41	0.4	0.89	0.68	1.1	
Beta-Hexachlorocyclohexanef	0.3	1.1	0.4	1.23	1.42	0.2	0.93	0.39	1.06	0.18	1.07	
Bis(2-chloroethyl) phosphate**	1.04	0.97	1.04	0.97		1.06	1.03	1.03	0.92	1.04	0.96	
Bromodichloromethane	0.94	1.01	0.61	0.95		0.83	1.19	1.16	1.13	1.17	0.87	
Bromoform	0.82	1.04	0.93	1.56		0.54	0.58	0.61	1	1.06	1.01	
Cadmium	0.7	1.15	0.66	1.1		0.67	1.16	0.87	1.19	0.66	1.15	
Carbon tetrachloride	0	1.2	0	1.76		0	1.76	0	0	0	1.14	
Chloroethane	0	1.2	0	0		0	1.77	0	0	0	2.26	
Chloroform	0.89	1.02	0.69	0.96		1.2	1.21	1.05	1.07	0.78	0.91	
Dem. Subgroup:												
Age:	<19	20+	<19	20+	<19	<19	20+	<19	20+	<19	20+	20+
Race/eth:	Hispanic	Hispanic	NH Asian	NH Asian	NH Black	NH Black	NH Black	NH Other	NH Other	NH White	NH White	NH White

Continued, next page...

Table A15: NHANES detected sample ratios: age-differentiated results

Chemical																				
N-Nitrosomorpholine	1.04	0.98	0.45	0.7									1.86	1.34	1.29	0.84	0.8	1.02		
N-Nitrosopiperidine	1.45	0.79	1.63	1.03									1.85	0.87	1.42	0.75	0.78	0.63		
N-Nitrosopyrrolidine	1.05	0.98	0.78	2.08									2.02	0	0	0.87	1.13	0.85		
NAS-(2-Carboxyethyl)-LC**	1	1	0.99	1									1	1	1	1	1	1		
NAS-(2-Hydroxyethyl)-LC**	1.06	0.96	1.04	0.9									1.27	1.23	0.86	0.78	1.01	0.94		
NAS-(2-carbamoylethyl)-LC**	1	1	1	1									1	1	1	1	1	1		
NAS-(2-cyanoethyl)-LC**	0.95	1.04	0.89	1.01									1.05	1.19	0.94	0.94	0.93	1.01		
NAS-(2-hydroxy-3-butenyl)-LC**	0.19	1.55	0.1	1.17									0.41	2.25	0.27	0.81	0.07	1.84		
NAS-(2-hydroxypropyl)-LC**	0.99	1.01	1	1									1	1.04	0.98	0.97	0.99	1.01		
NAS-(3,4-dihydroxybutyl)-LC**	1	1	1	1									1	1	1	1	1	1		
NAS-(3-Hydroxypropyl)-LC**	1	1	1	1									1	1	1	1	1	1		
NAS-(N-methylcarbamoyl)-LC**	1	1	0.99	1									1	1.01	0.99	1	1	1		
NAS-(3-hydroxypropyl-1-methyl)-LC**	1	1	1	1									1	1	1	1	1	1		
NNAL**	1.01	0.99	0.93	0.97									1.23	1.2	0.89	0.84	0.98	0.96		
Nickel	1.01	0.99	1	1									1.02	1.01	1.02	0.96	1.01	0.99		
Nitromethane	1	1	1	1									1	1	1	1	1	1		
Oxypyrimidine**	0.96	1.02	0.84	0.85									1.43	1.64	0.82	0.86	0.7	0.87		
PBB153†	0.15	1.12	0.17	1.14				0.31	0.88				0.39	1.11	0	1.07	0.07	1.14		
PCB028†	0.02	1.14	0	0.95				0	1.49				0.12	0.8	0	0.89	0	1.23		
PCB066†	0.06	1.13	0	0.59				0.33	2.33				0.29	1.01	0	1.06	0	1.19		
PCB074†	0.79	1.03	0.73	1				1.04	1.04				0.75	1.03	0.64	1.04	0.81	1.04		
PCB081†	0	1.14	0	1.32				0	4.48				0	0	0	0	0	1.08		
PCB105†	0.8	1.03	0.77	1.01				1.03	1.03				0.65	1.03	0.64	0.98	0.84	1.03		
PCB114†	0.13	1.12	0.1	0.81				0.2	1.16				0	1.05	0	1.33	0.18	1.2		
PCB126†	0.22	1.11	0.29	1.15				0.66	1.35				0.31	0.99	0	0.78	0.14	1.12		
PCB146†	0.65	1.05	0.17	1.01				1.06	1.06				0.91	1.06	1.06	1.06	0.73	1.06		
PCB156†	0.76	1.03	0.41	1.04				1.04	1.04				0.72	1.04	1.04	1.04	0.88	1.03		
PCB157†	0.13	1.12	0.09	0.84				0.19	1.33				0.1	1.04	0	1.21	0.17	1.18		
PCB167†	0.18	1.12	0.09	1.03				0.56	1.51				0.09	1.07	0	0.86	0.22	1.13		
PCB169†	0.11	1.13	0	1				0.65	1.28				0	0.99	0	0.75	0.14	1.2		
PCB170†	0.99	1	1.01	1				1.01	1.01				0.87	1.01	1.01	0.95	1.01	1.01		
PCB178†	0.26	1.11	0.12	0.93				0.69	1.24				0.27	1.07	0	1.05	0.31	1.15		
PCB183†	0.6	1.06	0.41	1.06				1.08	1.03				0.6	1.06	0.67	0.92	0.64	1.07		
PCB187†	0.94	1.01	0.88	1				1.04	1.04				1.04	1.04	1.04	1.04	0.92	1		
PCB189†	0	1.14	0	0.66				0	1.17				0	1.06	0	1.42	0	1.25		
PCB194†	0.69	1.04	0.45	1.05				1.05	1.05				0.59	1.05	0.29	1.05	0.81	1.04		
PCB196†	0.79	1.03	0.72	1.01				1.03	1.03				0.71	1.03	1.03	1.03	0.79	1.03		
Dem. Subgroup:																				
Age:	<19	20+	Hispanic	<19	Hispanic	20+	Hispanic	<19	NH Asian	20+	NH Asian	<19	NH Black	20+	<19	NH Other	20+	<19	NH White	20+
Race/eth:	<19	20+	Hispanic	<19	Hispanic	20+	Hispanic	<19	NH Asian	20+	NH Asian	<19	NH Black	20+	<19	NH Other	20+	<19	NH White	20+

Continued, next page...

Table A16: RSEI average ambient concentration ratios: complete results

Chemical	Hispanic	Amer. Ind.	Asian	Black	Pac. Isl.	White	Inc: <2xPL	Inc: >2xPL
1,1,1,2-Tetrachloroethane	0.36	0.24	0.42	1.24	0.39	1.23	1.31	0.93
1,1,1-Trichloroethane	0.29	0.25	0.32	1.26	0.05	1.26	1.25	0.94
1,1,2,2-Tetrachloroethane	0.88	0.24	0.35	1.78	0.13	0.96	1.22	0.92
1,1,2-Trichloroethane	0.62	0.23	0.37	1.81	0.22	1.03	1.28	0.92
1,1-Dichloroethane	0.37	0.25	0.34	1.5	0.04	1.18	1.27	0.94
1,1-Dichloroethylene	0.41	0.19	0.3	1.38	0.16	1.2	1.26	0.93
1,1-Dimethylhydrazine	1.08	0.4	0.51	2.18	0.33	0.82	1.19	0.95
1,2,3-Trichloropropane	3.09	0.18	0.06	1.35	0.23	0.42	1.01	0.76
1,2-(Methylenedioxy)-4-propylbenzene	0.1	0.11	0.23	0.38	1.18	1.49	0.99	0.99
1,2-Benzenediol	1.02	0.4	3.13	1.97	4.15	0.57	1.01	1.03
1,2-Dibromo-3-chloropropane	0.17	0.21	0.22	1.38	0.59	1.28	1.18	0.95
1,2-Dibromoethane	0.65	0.32	0.63	4.1	0.31	0.52	1.83	0.82
1,2-Dichloroethane	1.14	0.2	0.34	2.29	0.16	0.78	1.29	0.9
1,2-Dichloropropane	1.74	0.24	0.14	1.96	0.41	0.68	1.19	0.81
1,2-Diphenylhydrazine	1.12	0.47	0.46	2.16	0.28	0.82	1.09	0.98
1,2-Epoxybutane	2.12	0.15	0.92	1.3	0.23	0.63	1.02	0.97
1,2-Phenylenediamine	0.52	0.56	0.1	4.93	0.13	0.46	1.14	0.88
1,2-Propylene oxide	1.65	0.42	0.64	1.43	0.63	0.76	1.22	0.93
1,3-Butadiene	2.38	0.27	0.57	1.34	0.39	0.57	1.36	0.89
1,3-Dichloropropene	2.08	0.29	0.37	1.6	0.51	0.63	1.23	0.84
1,3-Propane sultone	0.87	0.13	0.54	1.05	0.08	1.09	1.15	0.94
1,4-Dichloro-2-butene	0.36	0.38	0.11	5.45	0.08	0.4	1.34	0.87
1,4-Dichlorobenzene	0.78	0.64	0.2	1.38	0.37	1.05	1.03	0.95
1,4-Dioxane	1.08	0.27	0.93	1.66	0.25	0.87	1.18	0.96
1-Bromopropane	1.02	0.29	1.66	0.94	5.02	0.93	0.97	1.03
1-Ethyl-1-nitrosourea	0.11	0.1	0.27	0.39	0.9	1.49	0.93	1.01
1-Naphthalenol, 1-(N-methylcarbamate)	0.4	0.24	0.41	2.94	0.19	0.82	1.26	0.91
1-Naphthylamine	0.82	0.67	0.3	2.5	0.23	0.86	1.18	0.95
2,3,7,8-Tetrachlorodibenzo-p-dioxin	1.05	0.91	0.55	0.9	0.51	1.06	1.19	0.92
2,4,6-Trichlorophenol	1.01	0.51	0.61	1.51	0.29	0.96	1.06	0.97
2,4-Diaminotoluene	1.01	0.15	0.16	2.19	0.13	0.88	1.47	0.89
2,4-Dichlorophenoxyacetic acid	0.42	0.27	0.39	1.79	0.15	1.1	1.43	0.89
2,4-Dinitrotoluene	0.13	0.54	0.13	0.21	0.52	1.53	1.4	0.87
2,6-Dimethylaniline	0.14	0.14	0.63	0.27	0.07	1.5	0.59	1.09
2,6-Dinitrotoluene	0.06	0.12	0.1	0.23	0.13	1.56	1.39	0.88
2-Anisidine	0.28	1.59	0.05	0.42	0.05	1.32	0.8	0.93
2-Mercaptobenzothiazole	0.42	0.24	0.25	0.72	0.37	1.3	1.48	0.92
2-Methoxy-5-methylaniline	0.15	0.25	0.37	4.9	0.38	0.54	1.76	0.8
2-Methylaniline	2.66	0.58	0.41	1	0.32	0.55	1.48	0.82
2-Methylaniline hydrochloride	0.09	0.13	0.18	0.36	1.67	1.5	1.1	0.95
2-Nitropropane	0.16	0.1	0.23	1.72	0.07	1.23	1.6	0.89
2-Phenylphenol	2.04	0.29	1.39	1.33	0.24	0.59	0.99	0.91
3,3',5,5'-Tetrabromobisphenol A	1.8	0.2	2.07	1.07	1.1	0.65	1.09	0.98
3,3'-Dichlorobenzidine	1.11	0.44	0.49	2.2	0.3	0.81	1.15	0.96
3,3'-Dichlorobenzidine dihydrochloride	0.36	1.13	0.14	0.68	0.1	1.32	0.91	0.94
3,3'-Dimethylbenzidine	1.13	0.49	0.37	2	0.23	0.86	0.97	1.01
3-Chloro-2-methylpropene	0.28	0.62	0.54	1.63	0.25	1.13	1.07	0.99
4,4'-Diaminobiphenyl methane	0.58	0.15	0.18	2.22	0.14	0.98	0.87	1.08
4,4'-Methylenebis(2-chloroaniline)	0.58	0.15	0.86	1.36	0.14	1.09	0.85	1.03
4,4'-Oxydianiline	0.5	0.15	0.49	1.52	0.16	1.12	1.15	0.98
4-Biphenylamine	0.55	0.6	0.08	5.08	0.15	0.43	1.12	0.88
4-Chloroaniline	0.51	0.21	0.37	1.76	0.42	1.06	0.77	1.05
4-Methyl-2-pentanone	1.14	0.56	0.73	1.69	0.59	0.85	1.21	0.94

Continued, next page...

Table A16: RSEI average ambient concentration ratios: complete results

Chemical	Hispanic	Amer. Ind.	Asian	Black	Pac. Isl.	White	Inc: <2xPL	Inc: >2xPL
5,5-Diphenylhydantoin	0.32	0.38	0.32	0.82	0.27	1.3	1.03	0.97
Acetaldehyde	1.06	0.36	0.59	1.43	0.34	0.95	1.25	0.92
Acetamide	0.26	0.11	0.29	1.5	0.24	1.2	1.24	0.94
Acifluorfen sodium	0.32	0.18	0.15	4.68	0.29	0.58	1.93	0.76
Acrolein	0.92	0.41	0.47	0.99	0.37	1.1	1.14	0.94
Acrylamide	0.71	0.49	1.3	3.09	0.57	0.63	1.4	0.89
Acrylonitrile	0.56	0.37	0.38	1.67	0.22	1.07	1.2	0.93
Alachlor	0.74	0.19	0.17	0.32	0.04	1.33	1	0.99
Aldrin	0.35	0.31	0.44	3.73	0.22	0.7	1.73	0.83
Amitrole	1.05	0.4	0.49	2.16	0.33	0.83	1.19	0.95
Aniline	0.61	0.34	0.48	2.49	0.11	0.89	0.82	1.03
Anthracene	1.64	1.27	1.26	1.43	1.66	0.68	1.25	0.94
Arsenic	1.72	4.58	0.42	0.38	3.87	0.92	1.03	0.99
Asbestos	1.46	0.84	0.73	1.18	0.42	0.85	1.07	0.97
Auramine	1.13	0.49	0.36	1.97	0.22	0.87	0.95	1.01
Benzene	1.58	0.5	0.79	1.56	1.18	0.74	1.31	0.91
Benzotrichloride	0.07	0.32	0.14	0.44	0.6	1.46	1.48	0.83
Benzyl chloride	0.65	0.25	1.18	1.83	0.17	0.93	1	0.93
Beryllium	0.36	0.39	0.4	0.69	0.26	1.34	1.05	0.98
Bis(chloroethyl) ether	1.36	7.17	0.5	0.86	0.45	0.87	1.19	0.91
Bromodichloromethane	0.15	0.2	0.11	0.43	0.34	1.49	1.29	0.92
Bromoform	0.15	0.2	0.11	0.43	0.34	1.49	1.29	0.92
C.I. Azoic Diazo Component 112	1.14	0.49	0.4	2.06	0.23	0.84	1	1
C.I. Direct Blue 14	1.16	0.41	0.5	2.13	0.33	0.81	1.18	0.95
C.I. Direct Blue 218	0.4	0.26	0.37	1.85	0.48	1.07	0.78	1.06
C.I. Disperse Black 6	1.13	0.48	0.42	2.09	0.25	0.84	1.03	0.99
Cadmium	1.04	0.93	0.39	1.48	0.69	0.96	1.33	0.92
Captan	0.46	1.22	0.04	1.78	0.04	1.11	1.23	0.86
Carbon tetrachloride	0.77	0.32	0.52	2.18	0.59	0.89	1.14	0.96
Chlordane	0.44	0.46	0.42	3.22	0.22	0.78	1.53	0.87
Chlorobenzilate	0.4	0.59	0.3	3.02	0.29	0.87	1.29	0.9
Chloroethane	0.58	0.2	0.4	3.18	0.25	0.75	1.61	0.8
Chloroform	0.62	0.24	0.45	2.09	0.22	0.97	1.16	0.95
Chloromethyl methyl ether	0.26	0.48	0.46	0.27	0.04	1.47	0.96	0.99
Chloroprene	0.41	0.41	0.2	4.41	0.09	0.58	1.29	0.89
Chlorothalonil	0.37	0.23	0.42	3.03	0.19	0.81	1.29	0.91
Cobalt	1.54	0.5	0.79	1.28	0.72	0.81	1.15	0.95
Creosote	0.36	0.36	0.21	3.03	0.15	0.88	1.58	0.81
Crotonaldehyde	3.75	0.37	0.17	0.41	0.24	0.37	2.24	0.79
Cumene	1.62	0.27	0.96	1.53	0.43	0.72	1.38	0.9
Cupferron	0.15	0.59	0.23	3.47	0.18	0.86	1.31	0.88
Di(2-ethylhexyl) phthalate	1.34	0.32	2.11	0.72	1.75	0.83	0.94	1.02
Diazinon	0.15	0.16	0.13	5.36	0.09	0.5	2.15	0.71
Dichloromethane	0.7	0.36	0.9	1.67	0.63	0.98	1.12	0.98
Dichlorvos	0.11	1.83	0.18	2.29	0.09	1.1	1.28	0.9
Diethanolamine	2.68	0.6	0.87	1.56	1.08	0.4	1.69	0.8
Diethyl sulfate	0.28	0.31	0.1	0.64	0.35	1.42	1.29	0.87
Diglycidyl resorcinol ether	0.17	0.05	0.62	0.14	0.06	1.52	0.31	1.24
Dimethyl sulfate	0.7	0.22	1.35	1.33	0.24	1.01	0.96	0.96
Dimethylcarbamoyl chloride	1.08	0.41	0.5	2.16	0.33	0.83	1.17	0.95
Diuron	1.39	0.3	1.06	1.13	1.06	0.85	0.97	1.01
Epichlorohydrin	1.62	0.65	0.75	1.47	0.84	0.75	1.19	0.9
Ethoprop	0.13	1.34	0.11	3.38	0.07	0.9	1.79	0.76

Continued, next page...

Table A16: RSEI average ambient concentration ratios: complete results

Chemical	Hispanic	Amer. Ind.	Asian	Black	Pac. Isl.	White	Inc: <2xPL	Inc: >2xPL
Ethyl acrylate	0.8	0.21	0.67	<u>1.5</u>	0.25	1	<u>1.73</u>	0.85
Ethylbenzene	1.23	0.5	0.9	1.39	1.15	0.86	1.19	0.94
Ethylene oxide	1.3	0.54	0.74	<u>1.81</u>	0.58	0.78	1.16	0.95
Ethylene thiourea	0.4	0.28	0.25	0.66	0.38	1.31	<u>1.62</u>	0.9
Ethyleneimine	0.22	0.43	0.1	<u>1.58</u>	0.51	1.24	1.32	0.9
Formaldehyde	0.82	1.02	0.55	1.24	0.66	1.05	1.21	0.93
Furan	0.54	0.41	0.27	<u>4.69</u>	0.04	0.45	2.12	0.69
Glycidol	<u>1.55</u>	0.51	0.44	<u>1.59</u>	0.17	0.77	1.31	0.9
Heptachlor	1.23	0.57	0.34	<u>2.17</u>	0.22	0.77	1.42	0.87
Hexachloro-1,3-butadiene	0.31	0.23	0.36	<u>2.66</u>	0.11	0.96	1.09	1.01
Hexachlorobenzene	0.39	0.18	0.46	0.95	0.1	1.29	1.21	0.99
Hexachloroethane	0.85	0.17	0.28	<u>2.73</u>	0.19	0.78	1.21	0.88
Hydrazine	0.45	0.36	0.23	<u>1.96</u>	0.14	1.08	<u>1.67</u>	0.86
Hydrazine sulfate	0.62	0.11	0.88	<u>2.19</u>	0.25	0.89	0.75	1.07
Isoprene	<u>1.9</u>	0.25	0.56	<u>2.07</u>	0.26	0.57	1.31	0.91
Lead	0.87	0.77	0.72	1.32	<u>1.55</u>	1	1.06	0.98
Lindane	1.14	0.61	0.44	<u>1.77</u>	0.29	0.89	1.16	0.93
Malathion	0.66	0.09	0.06	<u>5.23</u>	0.03	0.39	<u>1.96</u>	0.76
Maneb	0.33	0.48	0.12	<u>3.66</u>	0.01	0.78	<u>1.74</u>	0.82
Metam-sodium	<u>2.5</u>	0.56	0.36	0.19	0.77	0.77	1.09	0.89
Methyl acrylate	1.25	0.74	0.72	<u>1.78</u>	0.19	0.8	1.25	0.94
Methyl iodide	0.79	0.17	0.25	0.6	0.14	1.26	1.45	0.86
Methylhydrazine	1.12	0.46	0.47	<u>2.19</u>	0.29	0.81	1.12	0.97
Molybdenum trioxide	<u>2.17</u>	0.19	0.81	<u>1.88</u>	0.1	0.5	1.27	0.91
N,N-Dimethylformamide	1.15	0.39	1.47	1.11	0.66	0.89	0.98	1.02
N-Methylolacrylamide	0.79	0.17	0.71	0.89	0.16	1.13	<u>2.27</u>	0.73
N-Nitroso-N-methylurea	0.11	0.1	0.27	0.39	0.88	1.49	0.93	1.01
N-Nitrosodi-n-propylamine	1.04	0.25	0.14	0.2	0.02	1.25	1.26	0.86
N-Nitrosodibutylamine	0.11	0.11	0.25	0.38	1.02	1.49	0.96	1
N-Nitrosodiethylamine	0.11	0.11	0.26	0.38	0.97	1.49	0.95	1.01
N-Nitrosodiphenylamine	0.37	0.43	0.12	<u>5.28</u>	0.1	0.43	1.33	0.87
N-Nitrosomorpholine	0.2	0.06	0.3	1.2	<u>1.56</u>	1.28	<u>1.96</u>	0.73
N-Nitrosopiperidine	0.11	0.11	0.25	0.38	1.02	1.49	0.96	1
Naphthalene	<u>1.89</u>	0.47	0.98	1.43	0.86	0.65	1.22	0.93
Nickel	0.88	0.72	1.16	0.83	1.45	1.04	1.07	1
Nitrapyrin	<u>1.96</u>	0.37	<u>2.39</u>	1.19	<u>4.92</u>	0.49	1	1.01
Nitrioltri-acetic acid	<u>2</u>	0.31	1.08	1.26	0.16	0.65	0.82	1.04
Nitrobenzene	0.64	0.19	0.69	<u>1.83</u>	0.23	1	0.77	1.07
Nitromethane	0.15	0.36	0.2	<u>2.01</u>	0.09	1.17	<u>1.63</u>	0.87
Oxadiazon	1	0.14	0.19	0.73	0.05	1.16	<u>1.56</u>	0.84
Parathion	0.39	0.32	0.05	<u>4.5</u>	0	0.61	<u>1.64</u>	0.86
Pentachlorophenol	0.6	1.01	1.15	0.77	1.05	1.14	0.88	1
Pentaerythritol dibromide	0.89	0.2	1.23	<u>2.07</u>	0.1	0.8	1.43	0.91
Perfluorooctanesulfonic acid	0.34	0.63	0.1	<u>2.93</u>	0.02	0.93	<u>1.56</u>	0.83
Perfluorooctanoic acid	0.4	0.48	0.83	<u>3.18</u>	0.24	0.76	<u>1.71</u>	0.84
Phenolphthalein	0.74	1	1.1	1.19	0.08	1.01	0.81	1.08
Polychlorinated biphenyls	0.86	1.02	0.45	<u>1.77</u>	0.28	0.95	1.17	0.95
Polycyclic aromatic hydrocarbons	1.24	0.42	0.83	1.48	0.73	0.86	1.2	0.94
Potassium N-methyldithiocarbamate	0.09	2.2	0.16	<u>2.24</u>	0.08	1.12	1.32	0.9
Potassium bromate	1.1	0.15	<u>3.22</u>	1.07	0.13	0.75	0.74	1.12
Propachlor	0.35	0.27	0.31	0.57	0.08	1.37	0.8	1.06
Propargite	0.35	0.09	0.22	<u>5.11</u>	0.7	0.47	<u>2.2</u>	0.57
Propoxur	0.38	0.34	0.05	<u>4.47</u>	0	0.62	<u>1.66</u>	0.85
Propyleneimine	<u>1.98</u>	0.35	0.71	0.79	0.15	0.79	0.92	1.02

Continued, next page...

Table A16: RSEI average ambient concentration ratios: complete results

Chemical	Hispanic	Amer. Ind.	Asian	Black	Pac. Isl.	White	Inc: <2xPL	Inc: >2xPL
Propyzamide	0.11	0.11	0.26	0.38	0.95	1.49	0.94	1.01
Pyridine	0.67	0.18	0.67	1.55	0.23	1.03	1.32	0.9
Quinoline	2.3	0.13	0.88	1.57	0.09	0.52	1.27	0.91
Rhodamine B	4.34	0.08	0.22	0.53	0.11	0.18	1.87	0.64
Safrole	0.1	0.12	0.22	0.37	1.32	1.5	1.03	0.98
Sodium 2-phenylphenate	0.14	0.33	0.2	3.02	0.11	0.96	1.62	0.87
Styrene	1.27	0.86	0.79	1.09	0.66	0.93	1.21	0.92
Sulfuric acid	1.16	0.46	1.06	1.22	1.73	0.9	1.07	0.99
Tetrachloroethylene	1.73	0.57	0.85	0.97	1.11	0.79	1.15	0.93
Tetrafluoroethylene	0.49	0.45	0.38	1.58	0.16	1.12	1.09	0.97
Thioacetamide	1.1	0.42	0.5	2.2	0.32	0.81	1.17	0.95
Thiodicarb	0.39	0.24	0.42	2.95	0.19	0.82	1.27	0.91
Thiourea	0.4	0.39	1.34	0.94	2.02	1.14	1.2	0.98
Toluene 2,4-diisocyanate	0.37	0.29	0.46	3.33	0.65	0.79	1.17	0.95
Toluene diisocyanate	1.58	0.63	0.86	1.84	0.69	0.66	1.02	0.98
Toluene-2,6-diisocyanate	0.54	0.22	0.5	3.15	0.58	0.77	1.1	0.97
Toxaphene	1.16	0.6	0.42	1.85	0.28	0.86	1.2	0.93
Tribufos	0.1	2.12	0.17	2.27	0.09	1.11	1.3	0.9
Trichloroethylene	0.69	0.25	0.65	1.22	0.18	1.09	1.19	0.94
Tris(2,3-dibromopropyl) phosphate	0.11	0.11	0.25	0.38	1.03	1.49	0.96	1
Urethane	2.1	0.11	0.27	3.06	0.07	0.35	1.61	0.84
Vinyl acetate	1.57	0.31	0.94	1.31	0.36	0.78	1.16	0.95
Vinyl chloride	1.34	0.21	0.44	2.55	0.19	0.66	1.43	0.88
Vinyl fluoride	0.39	2.7	0.43	2.53	0.36	0.92	1.6	0.85
alpha-1,2,3,4,5,6-Hexachlorocyclohexane	1.09	0.4	0.51	2.19	0.32	0.82	1.19	0.95
bis(2-Chloro-1-methylethyl) ether	2.2	0.31	0.11	2.28	0.7	0.49	1.24	0.74

Notes: Presentation of the complete set of *average* ambient concentration ratios for the 186 carcinogens matched to RSEI data for our analysis. Ratios calculated as the arithmetic mean ambient concentration for a given demographic subgroup divided by the national arithmetic mean ambient concentration. **Bold** values denote concentration ratios ≥ 1.25 . Underlined values denote concentration ratios ≥ 1.5 .

Demographic subgroups: Hispanic, non-Hispanic American Indian or Alaskan Native, non-Hispanic Asian, non-Hispanic Black, non-Hispanic Pacific Islander, non-Hispanic White, household income below 2 times the poverty line, household income above 2 times the poverty line.

Return to: Section 3.2.1.

Table A17: RSEI 95th percentile ambient concentration ratios: complete results

Chemical	Hispanic	Amer. Ind.	Asian	Black	Pac. Isl.	White	Inc: <2xPL	Inc: >2xPL
1,1,1,2-Tetrachloroethane	*	*	*	*	*	*	*	*
1,1,1-Trichloroethane	*	*	*	*	*	*	*	*
1,1,2,2-Tetrachloroethane	*	*	*	*	*	*	*	*
1,1,2-Trichloroethane	*	*	*	*	*	*	*	*
1,1-Dichloroethane	*	*	*	*	*	*	*	*
1,1-Dichloroethylene	*	*	*	*	*	*	*	*
1,1-Dimethylhydrazine	*	*	*	*	*	*	*	*
1,2,3-Trichloropropane	*	*	*	*	*	*	*	*
1,2-(Methylenedioxy)-4-propylbenzene	*	*	*	*	*	*	*	*
1,2-Benzenediol	*	*	*	*	*	*	*	*
1,2-Dibromo-3-chloropropane	*	*	*	*	*	*	*	*
1,2-Dibromoethane	*	*	*	*	*	*	*	*
1,2-Dichloroethane	15.65	0	0	1076.99	0	0.04	48.61	0.1
1,2-Dichloropropane	*	*	*	*	*	*	*	*
1,2-Diphenylhydrazine	*	*	*	*	*	*	*	*
1,2-Epoxybutane	3.53	0	2.4	1.92	0.5	0.13	1.08	0.97
1,2-Phenylenediamine	*	*	*	*	*	*	*	*
1,2-Propylene oxide	1.32	0.31	0.37	3.67	0.02	0.72	1.4	0.91
1,3-Butadiene	1.67	0.06	0.67	2.18	0.68	0.74	1.38	0.9
1,3-Dichloropropene	*	*	*	*	*	*	*	*
1,3-Propane sultone	*	*	*	*	*	*	*	*
1,4-Dichloro-2-butene	*	*	*	*	*	*	*	*
1,4-Dichlorobenzene	*	*	*	*	*	*	*	*
1,4-Dioxane	2.08	0.05	1.04	3.74	0.04	0.69	1.09	0.98
1-Bromopropane	0.91	0.07	2.55	1.22	0.89	0.83	0.79	1.12
1-Ethyl-1-nitrosourea	*	*	*	*	*	*	*	*
1-Naphthalenol, 1-(N-methylcarbamate)	*	*	*	*	*	*	*	*
1-Naphthylamine	*	*	*	*	*	*	*	*
2,3,7,8-Tetrachlorodibenzo-p-dioxin	0.97	1.1	0.5	1.13	0.45	1.05	1.15	0.95
2,4,6-Trichlorophenol	*	*	*	*	*	*	*	*
2,4-Diaminotoluene	*	*	*	*	*	*	*	*
2,4-Dichlorophenoxyacetic acid	0.58	0	0	79.94	0	0.72	1.83	0.94
2,4-Dinitrotoluene	*	*	*	*	*	*	*	*
2,6-Dimethylaniline	*	*	*	*	*	*	*	*
2,6-Dinitrotoluene	*	*	*	*	*	*	*	*
2-Anisidine	*	*	*	*	*	*	*	*
2-Mercaptobenzothiazole	*	*	*	*	*	*	*	*
2-Methoxy-5-methylaniline	*	*	*	*	*	*	*	*
2-Methylaniline	*	*	*	*	*	*	*	*
2-Methylaniline hydrochloride	*	*	*	*	*	*	*	*
2-Nitropropane	*	*	*	*	*	*	*	*
2-Phenylphenol	*	*	*	*	*	*	*	*
3,3',5,5'-Tetrabromobisphenol A	2.6	0	3.8	1.11	0.15	0.2	0.96	1.03
3,3'-Dichlorobenzidine	*	*	*	*	*	*	*	*
3,3'-Dichlorobenzidine dihydrochloride	*	*	*	*	*	*	*	*
3,3'-Dimethylbenzidine	*	*	*	*	*	*	*	*
3-Chloro-2-methylpropene	*	*	*	*	*	*	*	*
4,4'-Diaminobiphenyl methane	*	*	*	*	*	*	*	*
4,4'-Methylenebis(2-chloroaniline)	*	*	*	*	*	*	*	*
4,4'-Oxydianiline	*	*	*	*	*	*	*	*
4-Biphenylamine	*	*	*	*	*	*	*	*
4-Chloroaniline	*	*	*	*	*	*	*	*
4-Methyl-2-pentanone	1.28	0.18	0.92	1.54	0.5	0.82	1.19	0.95

Continued, next page...

Table A17: RSEI 95th percentile ambient concentration ratios: complete results

Chemical	Hispanic	Amer. Ind.	Asian	Black	Pac. Isl.	White	Inc: <2xPL	Inc: >2xPL
5,5-Diphenylhydantoin	*	*	*	*	*	*	*	*
Acetaldehyde	0.98	0.28	0.66	<u>1.97</u>	0.19	0.92	1.25	0.94
Acetamide	*	*	*	*	*	*	*	*
Acifluorfen sodium	*	*	*	*	*	*	*	*
Acrolein	0.7	0	0.34	1.27	0	1.17	1.16	0.94
Acrylamide	0.41	0	88.88	35.31	47.07	0.75	0.85	1.1
Acrylonitrile	0.45	0	0.6	<u>3.39</u>	0.32	1.05	1.21	0.95
Alachlor	*	*	*	*	*	*	*	*
Aldrin	*	*	*	*	*	*	*	*
Amitrole	*	*	*	*	*	*	*	*
Aniline	0.03	0.13	0.01	82.18	0	0.82	1.46	0.92
Anthracene	1.17	0.39	<u>2.83</u>	<u>2.35</u>	<u>1.71</u>	0.64	1.24	0.99
Arsenic	1	1.58	0.7	1.15	2.2	0.98	1.07	0.96
Asbestos	*	*	*	*	*	*	*	*
Auramine	*	*	*	*	*	*	*	*
Benzene	<u>1.75</u>	0.49	0.82	<u>1.57</u>	0.94	0.74	1.31	0.92
Benzotrichloride	*	*	*	*	*	*	*	*
Benzyl chloride	3.12	0	2.29	9.01	0	0.24	1.11	1.01
Beryllium	0.23	0	0.82	<u>1.72</u>	0	1.11	1.02	1.01
Bis(chloroethyl) ether	*	*	*	*	*	*	*	*
Bromodichloromethane	*	*	*	*	*	*	*	*
Bromoform	*	*	*	*	*	*	*	*
C.I. Azoic Diazo Component 112	*	*	*	*	*	*	*	*
C.I. Direct Blue 14	*	*	*	*	*	*	*	*
C.I. Direct Blue 218	*	*	*	*	*	*	*	*
C.I. Disperse Black 6	*	*	*	*	*	*	*	*
Cadmium	1.21	0.04	0.58	2.19	1.3	0.88	1.18	0.97
Captan	*	*	*	*	*	*	*	*
Carbon tetrachloride	65.56	0	74.97	14.72	0	0.17	1.85	0.9
Chlordane	*	*	*	*	*	*	*	*
Chlorobenzilate	*	*	*	*	*	*	*	*
Chloroethane	4.22	0	0	44.77	0	0	4.34	0.57
Chloroform	0.73	0	0.86	<u>2.71</u>	0	0.98	1.09	1
Chloromethyl methyl ether	*	*	*	*	*	*	*	*
Chloroprene	*	*	*	*	*	*	*	*
Chlorothalonil	*	*	*	*	*	*	*	*
Cobalt	1.33	0.18	0.9	<u>1.59</u>	0.57	0.87	1.18	0.97
Creosote	0.28	0	0	1275.01	0	6.29	2.83	0.83
Crotonaldehyde	*	*	*	*	*	*	*	*
Cumene	<u>1.58</u>	0.18	<u>1.53</u>	3.03	0.47	0.65	1.36	0.94
Cupferron	*	*	*	*	*	*	*	*
Di(2-ethylhexyl) phthalate	<u>1.71</u>	0.02	2.92	0.55	1.32	0.68	0.92	1.04
Diazinon	*	*	*	*	*	*	*	*
Dichloromethane	0.81	0.18	1.11	1.23	0.6	1.04	0.96	1.02
Dichlorvos	*	*	*	*	*	*	*	*
Diethanolamine	<u>3.55</u>	0.27	1.41	<u>2.19</u>	0.52	0.38	1.41	0.88
Diethyl sulfate	*	*	*	*	*	*	*	*
Diglycidyl resorcinol ether	*	*	*	*	*	*	*	*
Dimethyl sulfate	*	*	*	*	*	*	*	*
Dimethylcarbamoyl chloride	*	*	*	*	*	*	*	*
Diuron	<u>1.71</u>	0	<u>1.64</u>	<u>1.58</u>	0	0	1.29	0.94
Epichlorohydrin	<u>1.71</u>	0	0.96	<u>3.32</u>	0	0.65	<u>1.61</u>	0.89
Ethoprop	*	*	*	*	*	*	*	*

Continued, next page...

Table A17: RSEI 95th percentile ambient concentration ratios: complete results

Chemical	Hispanic	Amer. Ind.	Asian	Black	Pac. Isl.	White	Inc: <2xPL	Inc: >2xPL
Ethyl acrylate	0.98	0.03	0.97	<u>1.88</u>	0.11	0.85	1.14	0.98
Ethylbenzene	1.32	0.54	0.91	<u>1.55</u>	1.33	0.83	1.25	0.93
Ethylene oxide	<u>1.6</u>	0.47	0.62	<u>3.16</u>	0.82	0.72	1.2	0.94
Ethylene thiourea	*	*	*	*	*	*	*	*
Ethyleneimine	*	*	*	*	*	*	*	*
Formaldehyde	0.92	0.81	0.54	1.28	0.59	1.02	1.13	0.95
Furan	*	*	*	*	*	*	*	*
Glycidol	*	*	*	*	*	*	*	*
Heptachlor	*	*	*	*	*	*	*	*
Hexachloro-1,3-butadiene	*	*	*	*	*	*	*	*
Hexachlorobenzene	0.85	0	0.61	<u>6.14</u>	0.22	0.81	1.36	0.96
Hexachloroethane	*	*	*	*	*	*	*	*
Hydrazine	<u>1.51</u>	0	1.41	<u>2.32</u>	0.25	0.56	0.91	1.04
Hydrazine sulfate	*	*	*	*	*	*	*	*
Isoprene	7.09	0	0.92	<u>4.99</u>	0.33	0.22	<u>1.84</u>	0.83
Lead	0.78	0.69	0.78	1.41	1.24	1.01	1.11	0.98
Lindane	*	*	*	*	*	*	*	*
Malathion	*	*	*	*	*	*	*	*
Maneb	*	*	*	*	*	*	*	*
Metam-sodium	*	*	*	*	*	*	*	*
Methyl acrylate	1.47	0	1.08	<u>1.83</u>	0	0.58	1.33	0.92
Methyl iodide	*	*	*	*	*	*	*	*
Methylhydrazine	*	*	*	*	*	*	*	*
Molybdenum trioxide	<u>5.6</u>	0.03	1	<u>5.08</u>	0.12	0.42	<u>1.62</u>	0.91
N,N-Dimethylformamide	1.4	0.17	1.38	1.01	0.98	0.8	0.98	1.01
N-Methylolacrylamide	0.33	0	1.44	<u>1.97</u>	0	0.78	0.57	1.1
N-Nitroso-N-methylurea	*	*	*	*	*	*	*	*
N-Nitrosodi-n-propylamine	*	*	*	*	*	*	*	*
N-Nitrosodibutylamine	*	*	*	*	*	*	*	*
N-Nitrosodiethylamine	*	*	*	*	*	*	*	*
N-Nitrosodiphenylamine	*	*	*	*	*	*	*	*
N-Nitrosomorpholine	*	*	*	*	*	*	*	*
N-Nitrosopiperidine	*	*	*	*	*	*	*	*
Naphthalene	<u>1.6</u>	0.36	1.15	<u>1.93</u>	0.58	0.69	1.26	0.94
Nickel	1.11	0.36	0.88	1.27	0.71	0.93	1.18	0.96
Nitrapyrin	*	*	*	*	*	*	*	*
Nitrilotriacetic acid	*	*	*	*	*	*	*	*
Nitrobenzene	*	*	*	*	*	*	*	*
Nitromethane	*	*	*	*	*	*	*	*
Oxadiazon	*	*	*	*	*	*	*	*
Parathion	*	*	*	*	*	*	*	*
Pentachlorophenol	*	*	*	*	*	*	*	*
Pentaerythritol dibromide	*	*	*	*	*	*	*	*
Perfluorooctanesulfonic acid	*	*	*	*	*	*	*	*
Perfluorooctanoic acid	*	*	*	*	*	*	*	*
Phenolphthalein	*	*	*	*	*	*	*	*
Polychlorinated biphenyls	*	*	*	*	*	*	*	*
Polycyclic aromatic hydrocarbons	1.22	0.3	0.88	<u>2.26</u>	0.45	0.74	1.35	0.92
Potassium N-methyldithiocarbamate	*	*	*	*	*	*	*	*
Potassium bromate	*	*	*	*	*	*	*	*
Propachlor	*	*	*	*	*	*	*	*
Propargite	*	*	*	*	*	*	*	*
Propoxur	*	*	*	*	*	*	*	*
Propyleneimine	*	*	*	*	*	*	*	*

Continued, next page...

Table A17: RSEI 95th percentile ambient concentration ratios: complete results

Chemical	Hispanic	Amer. Ind.	Asian	Black	Pac. Isl.	White	Inc: <2xPL	Inc: >2xPL
Propyzamide	*	*	*	*	*	*	*	*
Pyridine	0.21	0	<u>2.51</u>	0.68	0.43	1.38	0.72	1.14
Quinoline	*	*	*	*	*	*	*	*
Rhodamine B	*	*	*	*	*	*	*	*
Safrole	*	*	*	*	*	*	*	*
Sodium 2-phenylphenate	*	*	*	*	*	*	*	*
Styrene	1.45	0.53	1.04	1.04	0.7	0.84	1.2	0.93
Sulfuric acid	1.18	0.41	0.93	1.26	<u>1.82</u>	0.88	1.09	0.98
Tetrachloroethylene	<u>1.66</u>	0.13	1	1.07	0.65	0.84	0.97	1
Tetrafluoroethylene	*	*	*	*	*	*	*	*
Thioacetamide	*	*	*	*	*	*	*	*
Thiodicarb	*	*	*	*	*	*	*	*
Thiourea	*	*	*	*	*	*	*	*
Toluene 2,4-diisocyanate	0.17	0	0.81	<u>11.88</u>	0	0.91	0.98	1.03
Toluene diisocyanate	<u>2.39</u>	0.16	0.99	<u>1.51</u>	<u>1.69</u>	0.78	0.94	1.01
Toluene-2,6-diisocyanate	0.01	0	0.32	<u>16.28</u>	0	0.75	0.96	1.03
Toxaphene	*	*	*	*	*	*	*	*
Tribufos	*	*	*	*	*	*	*	*
Trichloroethylene	0.51	0.01	0.78	<u>1.58</u>	0.01	1.07	1.05	1.01
Tris(2,3-dibromopropyl) phosphate	*	*	*	*	*	*	*	*
Urethane	*	*	*	*	*	*	*	*
Vinyl acetate	<u>1.67</u>	0.07	1.18	<u>2.2</u>	0.33	0.7	1.21	0.95
Vinyl chloride	*	*	*	*	*	*	*	*
Vinyl fluoride	*	*	*	*	*	*	*	*
alpha-1,2,3,4,5,6-Hexachlorocyclohexane	*	*	*	*	*	*	*	*
bis(2-Chloro-1-methylethyl) ether	*	*	*	*	*	*	*	*

Notes: Presentation of the complete set of *95th percentile* ambient concentration ratios for the 186 carcinogens matched to RSEI data for our analysis. Ratios calculated as the 95th percentile ambient concentration for a given subgroup divided by the national 95th percentile ambient concentration. Values marked as * denote 95th percentile ratios that were not calculable due to the national 95th percentile concentration falling at an effective value of zero. This occurs with high frequency for chemicals that are only released from a small number of facilities with modest population density in proximity. **Bold** values denote concentration ratios ≥ 1.25 . Underlined values denote concentration ratios ≥ 1.5 .

Demographic subgroups: Hispanic, non-Hispanic American Indian or Alaskan Native, non-Hispanic Asian, non-Hispanic Black, non-Hispanic Pacific Islander, non-Hispanic White, household income below 2 times the poverty line, household income above 2 times the poverty line.

Return to: Section 3.2.1.

Table A18: Drinking water 95th percentile concentration ratios: complete results

Chemical	Hispanic	Amer. Ind.	Asian	Black	Pac. Isl.	White	<2x PL	>2x PL
1,4-DIOXANE	1.02	0.69	0.97	1.01	0.64	1.02	1.02	0.99
ARSENIC	1.11	<u>2.15</u>	0.81	0.69	0.92	0.98	1.01	0.98
BROMATE	0.73	0.96	0.84	1.04	0.74	1.04	1	0.96
CHROMIUM-6	1.37	<u>1.7</u>	1.18	0.55	1.18	0.95	1.05	0.99
DIBROMOACETIC ACID	<u>1.62</u>	0.96	0.95	0.88	0.95	0.95	1.09	0.99
DICHLOROACETIC ACID	0.94	1.15	0.95	1.08	0.98	1.01	1.03	0.99
DICHLOROMONOBROMOMETHANE	1.01	1.15	0.91	1.01	0.92	1	1.01	1
HAA5	1.1	0.87	1.1	1.1	0.75	0.89	1.02	0.96
HAA6BR	1.2	1.18	0.94	0.86	1.08	0.97	1.04	0.98
HAA9	1.03	1.01	1.03	1.03	0.74	0.93	1.03	0.97
MICROCYSTIN-LR	<u>5.59</u>	1	1	1	1	1	<u>5.59</u>	1
NDMA	1.2	0.88	0.84	1.2	0.65	0.93	1.2	1
TETRACHLOROETHYLENE	1	1.04	1	0.92	0.98	0.95	1	1
BROMOFORM	1.43	0.95	0.99	0.8	1	0.88	1.02	0.99
TRICHLOROACETIC ACID	1.01	0.92	1.01	1.01	0.77	0.85	1.01	1
TRICHLOROETHYLENE	<u>1.92</u>	1.13	1.48	0.94	0.98	0.94	1.04	0.98
CHLOROFORM	0.91	1.15	0.96	1.03	0.97	1.02	1.02	0.99

Notes: Presentation of the complete set of *95th percentile* relative drinking water concentration ratios for the 17 carcinogens that were monitored and detectable. Ratios calculated as the 95th percentile concentration for a given subgroup divided by the national 95th percentile concentration. **Bold** values denote concentration ratios ≥ 1.25 . Underlined values denote concentration ratios ≥ 1.5 .

Demographic subgroups: Hispanic, non-Hispanic American Indian or Alaskan Native, non-Hispanic Asian, non-Hispanic Black, non-Hispanic Pacific Islander, non-Hispanic White, household income below 2 times the poverty line, household income above 2 times the poverty line.

Return to: Section 3.2.2.

Table A19: Drinking water sample detection ratios: complete results

Chemical	Hispanic	Amer. Ind.	Asian	Black	Pac. Isl.	White	<2x PL	>2x PL
1,1,1-TRICHLOROETHANE	<u>2.27</u>	0.82	0.73	0.85	0.37	0.67	1.04	0.98
1,1,2-TRICHLOROETHANE	<u>3.67</u>	1.25	0.29	0.96	0.1	0.25	1.25	0.88
1,1-DICHLOROETHANE	1.18	1.13	1.23	0.56	0.52	1.02	0.88	1.06
1,1-DICHLOROETHYLENE	<u>2.91</u>	1.07	0.95	0.71	0.51	0.48	1.15	0.93
1,2,3-TRICHLOROPROPANE	<u>1.56</u>	0.63	<u>1.76</u>	0.48	<u>6.25</u>	0.79	0.95	1.02
1,2-DIBROMO-3-CHLOROPROPANE	<u>2.23</u>	0.8	<u>1.5</u>	0.55	<u>1.73</u>	0.61	1.12	0.94
1,2-DIBROMOMETHANE	1.25	<u>1.61</u>	1.4	1.41	<u>1.73</u>	0.77	1.1	0.95
1,2-DICHLOROETHANE	<u>2.74</u>	0.93	0.58	0.78	0.17	0.56	1.1	0.95
1,2-DICHLOROPROPANE	<u>3.19</u>	1.12	0.6	0.91	0.25	0.38	1.18	0.92
1,3-BUTADIENE	<u>1.55</u>	0.26	0.63	<u>2.02</u>	0.4	0.58	1.36	0.83
1,4-DIOXANE	1.06	0.62	0.94	1.11	0.54	0.97	1.03	0.99
17-ALPHA-ETHYNYLESTRADIOL	0.65	0.49	0.66	<u>1.77</u>	0.15	0.98	1.21	0.9
17-BETA-ESTRADIOL	0.9	1.01	0.85	0.61	<u>2.19</u>	1.13	1.31	0.85
2,3,7,8-TCDD	0.26	<u>2.25</u>	0.17	0.18	0.41	<u>1.68</u>	1.02	0.99
2,4-D	0.6	0.64	0.8	1.42	0.28	1.06	1	1
4-ANDROSTENE-3,17-DIONE	0.91	0.78	0.96	1.37	0.58	0.93	1.09	0.96
ALACHOR	0.74	1.46	0.98	<u>1.91</u>	0.25	0.88	1.18	0.91
ALPHA-HEXACHLOROCYCLOHEXANE	0.8	0.81	0.68	1.09	0.65	1.09	1.04	0.98
ARSENIC	<u>1.54</u>	1.42	0.92	0.75	0.98	0.88	1.04	0.98
ASBESTOS	0.76	<u>1.76</u>	0.64	1.15	0.33	1.1	0.94	1.03
BENZENE	<u>3.03</u>	1.4	0.34	0.99	0.23	0.44	1.21	0.9
BERYLLIUM, TOTAL	1.01	0.81	0.97	1.02	0.39	1	0.98	1
BHC-GAMMA	1.3	1.08	1.01	2.35	0.13	0.61	1.29	0.86
BIS(2-ETHYLHEXYL) PHTHALATE	0.74	0.7	0.81	<u>1.78</u>	0.3	0.94	0.98	1.01
BROMATE	1.22	1.27	1.23	0.78	1.13	0.9	1.02	0.99
CADMIUM	0.8	1.34	0.62	1	0.32	1.1	1.02	0.99
CARBON TETRACHLORIDE	<u>1.88</u>	0.95	0.68	1.45	0.32	0.66	1.18	0.91
CHLORDANE	0.47	<u>2.32</u>	4	0.53	<u>13.64</u>	0.86	0.87	1.06
CHROMIUM-6	1.02	0.97	1.06	1.01	1.09	0.98	1	1
COBALT	0.98	1.27	0.83	0.89	0.21	1.06	0.79	1.1
DECACHLOROBIPHENYL	0.41	<u>6.38</u>	0.4	0.45	0.16	1.31	0.9	1.05
DIBROMOACETIC ACID	1.2	0.88	0.98	0.94	0.76	0.95	1.01	0.99
DICHLOROACETIC ACID	0.94	0.92	1.01	1.09	0.71	1	1	1
DICHLOROMONOBROMOMETHANE	0.96	0.95	0.99	1.05	0.75	1.01	0.99	1
ETHOPROP	0.3	0.38	0.22	1.38	0.37	1.26	1.08	0.96
ETHYLBENZENE	<u>1.72</u>	1.17	0.63	0.89	0.37	0.84	1.03	0.98
GLYPHOSATE	<u>1.72</u>	1.44	<u>1.61</u>	0.37	0.71	0.86	0.8	1.1
HAA5	1	1	1	1	1	1	1	1
HAA6BR	1	1	1	1	1	1	1	1
HAA9	1	1	1	1	1	1	1	1
HEPTACHLOR	1.04	<u>1.62</u>	0.99	1.06	0.46	0.97	1.22	0.89
HEPTACHLOR EPOXIDE	0.58	1.41	0.67	1.06	0.75	1.15	0.99	1.01
HEXACHLOROBENZENE	0.82	<u>2.29</u>	0.47	0.6	0.24	1.19	0.82	1.09
METHYLENE CHLORIDE	<u>1.79</u>	0.96	0.77	1.05	0.91	0.76	1.08	0.96
MICROCYSTIN-LR	<u>4.01</u>	0	0	0.01	0	0.03	<u>2.5</u>	0.54
NDBA	0.93	0.78	0.7	0.7	0.64	1.18	0.93	1.03
NDEA	0.57	0.71	1.01	<u>1.87</u>	0.37	0.91	0.97	1.01
NDMA	1.11	0.79	0.81	0.95	0.61	1	1.02	0.99
NMEA	0.43	<u>17.15</u>	0.14	0.04	0.73	<u>1.57</u>	0.98	1.01
NPYR	0.72	0.53	0.85	1.11	0.28	1.13	1.03	0.99
O-TOLUIDINE	1.39	0.8	0.87	0.84	0.55	0.92	1.14	0.93
P-DICHLOROBENZENE	<u>2.34</u>	1.38	0.31	0.77	0.38	0.7	1.19	0.91

Continued, next page...

Table A19: Drinking water sample detection ratios: complete results

Chemical	Hispanic	Amer. Ind.	Asian	Black	Pac. Isl.	White	<2x PL	>2x PL
PFOA	0.9	0.71	1.22	1.06	0.63	1	0.95	1.03
PFOS	<u>1.74</u>	1.2	0.94	1.27	0.57	0.67	1.2	0.91
PAHS	1	<u>3.3</u>	0.87	0.91	0.8	1.01	0.93	1.04
QUINOLINE	0.7	0.75	<u>2.44</u>	0.57	1.14	1.04	0.79	1.1
STYRENE	<u>2.55</u>	1.29	1	0.9	0.16	0.54	1.18	0.91
TESTOSTERONE	0.88	0.5	0.85	<u>1.51</u>	0.57	0.92	1.09	0.96
TETRACHLOROETHYLENE	<u>1.62</u>	0.57	1.45	0.87	0.77	0.8	1.04	0.98
TOXAPHENE	0.3	<u>8.13</u>	0.13	0.28	0.02	1.43	0.81	1.09
TRIBROMOMETHANE	1.32	0.85	1.11	0.72	1.34	0.94	1	1
TRIBUFOS	0.13	0.23	0.07	0.38	0.08	<u>1.63</u>	0.98	1.01
TRICHLOROACETIC ACID	0.92	0.92	1.02	1.14	0.71	1	1	1
TRICHLOROETHYLENE	<u>2.1</u>	0.51	<u>1.79</u>	0.74	0.66	0.65	1.1	0.95
TRICHLOROMETHANE	0.93	0.95	0.99	1.07	0.74	1.01	0.99	1
VINYL CHLORIDE	<u>2.53</u>	0.94	0.47	1.07	0.11	0.57	1.15	0.93

Notes: Presentation of the complete set of drinking water sample detection concentration ratios for the 66 carcinogens that were monitored and had at least one detectable sample. The sample detection ratio for a given chemical and subgroup is given as the subgroup's survey-weighted fraction of samples with concentrations above the detection limit divided by the total population's survey-weighted fraction of samples with concentrations above the detection limit. Drinking water detection limits can vary both within and across PWSs; interpretation of these results should bear in mind the potential for correlations between a PWS's detection limit for a chemical, its size, and its customers' demographics. **Bold** values denote concentration ratios ≥ 1.25 . Underlined values denote concentration ratios ≥ 1.5 .

Demographic subgroups: Hispanic, non-Hispanic American Indian or Alaskan Native, non-Hispanic Asian, non-Hispanic Black, non-Hispanic Pacific Islander, non-Hispanic White, household income below 2 times the poverty line, household income above 2 times the poverty line.

Return to: Section 3.2.2.

Table A20: Household products consumption prevalence ratios: complete results

Chemical	Hispanic	Asian	Black	White	<\$30k	>\$30k	Child <18	Female, CBA
1,2-Propylene oxide	<u>1.9</u>	1.38	0.36	1.22		1.01	0.68	0.80
1,4-Dichlorobenzene	0.95	1.05	<u>2.7</u>	0.76	1.25	0.96	0.62	0.55
1,4-Dioxane	1.39	1.24	0.79	0.83	0.49	0.95	1.24	1.24
1-Chloro-4-(trifluoromethyl)benzene				<u>1.97</u>		1.01	1.27	
1-Naphthalenol, 1-(N-methylcarbamate)	0.59	0.6	0.78	1.04	1.06	1.03	0.61	0.63
2,4-Dichlorophenoxyacetic acid	0.98	0.98	0.98	1.02	0.91	1.04	0.73	0.67
4-Methyl-2-pentanone	<u>1.74</u>	0.54	0.99	0.8	0.7	0.77	0.73	0.63
Acetamide	0.56	0.61		0.94	0.87	0.89	0.3	0.09
Amides, coco, N,N-bis(hydroxyethyl)	<u>1.63</u>	0.47	0.31	1.07	0.98	0.95	0.95	0.93
Arsenic	1.42	0.91	0.86	0.98	0.64	1.01	1.47	1.60
Asphalt, oxidized	0.86	<u>2.48</u>	0.82	1.2	0.56	1.28	0.3	0.18
Benzophenone		<u>7.74</u>		1.47			0.95	
Butylated hydroxyanisole	0.9	1.24	0.97	1.02	0.83	1.02	0.78	0.76
C.I. Solvent Yellow 14	1.06		1.21	0.87	<u>1.79</u>	0.68	0.56	
Cadmium	<u>1.51</u>	0.76	0.71	0.96	0.69	0.95	1.41	1.55
Chlorothalonil	0.43	0.47	0.99	0.97	0.73	0.87	0.63	0.48
Coal tar			<u>2.72</u>	0.66				
Cobalt				0.98				
Cumene	0.87	0.97	1.15	0.92	1.16	0.96	0.74	0.81
Di(2-ethylhexyl) phthalate	<u>3.57</u>			1.23		<u>1.52</u>		
Dichloromethane	1	0.43	0.91	1.09	1.21	1.22	1.01	0.85
Diethanolamine	0.73	<u>1.76</u>	1.19	0.88	1.17	0.9	0.86	0.57
Diuron	<u>1.6</u>	0.77	0.61	0.84	0.9	0.7	<u>1.51</u>	1.23
Ethanol	1.03	0.94	1.01	0.99	0.92	1.01	1.06	1.03
Ethylbenzene	0.92	0.83	1.05	0.94	1.07	0.96	0.72	0.73
Ethylene oxide	<u>1.58</u>	1.21	0.8	0.8	0.42	0.89	1.26	1.35
Formaldehyde	0.67	0.26	0.91	0.92	1.02	0.97	0.77	0.57
Furan	<u>1.9</u>	1.38	0.36	1.22		1.01	0.68	0.80
Lead	0.94	0.6	1.25	0.9	0.77	0.97	1.05	0.98
Melamine			<u>1.86</u>	0.62	1.38	0.43	0.43	
Mineral oil - includes paraffin oil	1.25	1.11	<u>1.83</u>	0.88	0.92	1.02	1.27	1.28
Myrcene	0.72	0.31	0.5	1.17	1.47	0.8	1.23	1.28
Naphthalene	0.7	0.81	1.2	0.94	1.23	0.97	0.66	0.66
Nickel	0.81	0.36	0.91	0.96	1.11	0.93	0.74	0.68
Nickel (II) oxide	0.49	1.06	<u>1.81</u>	0.82	0.9	1.06	0.48	0.46
Nitrofurazone	1.46			0.35		0.16	0.39	0.46
Phenolphthalein	<u>1.6</u>	<u>2.25</u>	0.73	0.96	1.06	0.92	<u>2.19</u>	1.69
Phenyl glycidyl ether	1.38		0.2	1.21	1.07	1.13	<u>1.56</u>	0.98
Quartz (SiO2)	0.98	0.68	1.44	0.95	0.98	1.01	0.87	0.79
Talc	1.18	0.62	1.03	0.91	0.73	0.96	<u>1.56</u>	1.45
Tetrachloroethylene	0.61	1.33	0.53	1.04	0.95	1.11	0.74	1.16
Thiourea	0.68	0.34	0.94	1.07	0.73	1.06	0.7	0.64
Titanium dioxide	1.1	1.17	0.98	0.98	0.87	1	1.08	1.07
Vinyl acetate	1.01	0.35	0.12	1.02	0.52	0.88	0.92	0.94

Notes: Presentation of the complete set of chemical purchase prevalence ratios for the 44 carcinogens matched to the Stanfield et al. (2021) household purchasing and products data. Due to the original paper's data construction, racial subgroups (Asian, Black, White) are not exclusively non-Hispanic ethnicity. All racial/ethnic subgroup assignments are based on the female head of household. Household-income subgroups differentiated by those earning above and below \$30,000 in the year of data collection. *Child <18* denotes whether there is a child younger than 18 in the household. *Female, CBA* denotes whether there is a female of child-bearing age (<45 years old) in the household.

Return to: Section 3.2.3.

Table A21: Extended synthesis: carcinogens with evidence of potential exposure disparities

Chemical	NHANES Avg			NHANES P95			HHP	DW	RSEI	NHANES Cycle
	Prim.	Int.	Age	Prim.	Int.	Age				
Arsenic	✓	✓	✓	✓	✓	✓	✓	✓	✓	2015-2016
1-Naphthylamine	✓	✓	✓	✓	✓	✓			✓	2013-2014
Acrylonitrile**	✓	✓	✓	✓	✓	✓			✓	2015-2016
1,4-Dichlorobenzene*	✓	✓	✓	✓	✓	✓	✓		✗	2017-2018
2,4-Dichlorophenoxyacetic acid*	✓	✓	✓	✓	✓	✓	✗		✓	2015-2016
Ethylene oxide*	✗	✓	✓	✓	✓	✓	✓		✓	2015-2016
Nitrofen**	✓	✓	✓	✓	✓	✓				2015-2016
Acrylamide*	✗	✓	✓	✓	✓	✓			✓	2015-2016
Melamine	✓	✓	✓	✗	✓	✓	✓			2003-2004
1-Naphthalenol, 1-(N-methylcarbamate)**	✓	✓	✓	✗	✓	✓	✗		✓	2015-2016
Naphthalene**	✓	✓	✓	✗	✓	✓	✗		✓	2015-2016
2-Naphthylamine	✓	✓	✓	✗	✓	✓				2013-2014
4-Biphenylamine	✗	✓	✓	✗	✓	✓			✓	2013-2014
Acrolein**	✓	✓	✓	✗	✓	✓			✗	2015-2016
Crotonaldehyde*	✗	✓	✓	✗	✓	✓			✓	2015-2016
Di(2-ethylhexyl) phthalate**	✗	✗	✓	✗	✓	✓	✓		✓	2017-2018
Diazinon**				✓	✓	✓			✓	2013-2014
Ethylene thiourea				✓	✓	✓			✓	2007-2008
PCBs†	✓	✓	✓						✓	2015-2016
Vinyl chloride**				✓	✓	✓			✓	2015-2016
Furan				✓	✓	✗	✓		✓	2017-2018
4-(N-Methyl-N-nitrosamino)-1-(3-pyridyl)-1-butanone**	✓	✓	✓	✗	✓	✗				2013-2014
2-Methylaniline	✓	✓	✓	✗	✗	✗			✓	2013-2014
Dichlorvos**	✗	✗	✓	✗	✓	✓			✓	2017-2018
Malathion**	✗	✗	✓	✗	✓	✓			✓	2017-2018
Perfluorooctanesulfonic acid	✗	✗	✗	✓	✓	✓			✓	2017-2018
Beta-Hexachlorocyclohexane†	✓	✓	✓							2015-2016
DDT†	✓	✓	✓							2015-2016
Dimethylarsinic acid	✗	✗	✗	✓	✓	✓				2017-2018
Z-Tetrachlorvinphos**	✗	✗	✓	✗	✓	✓				2017-2018
1,3-Butadiene**	✗	✗	✗	✗	✓	✓			✓	2015-2016
1-Bromopropane**	✗	✗	✗	✗	✓	✓			✓	2015-2016
Perfluorooctanoic acid	✗	✗	✗	✗	✓	✓			✓	2017-2018
4-Methyl-2-pentanone							✓		✓	2017-2018
2-Amino-alpha-carboline				✗	✓	✓				2013-2014
MeA-alpha-C				✗	✓	✓				2013-2014
2,4,6-Trichlorophenol				✗	✓	✗			✓	2009-2010
2-Phenylphenol				✗	✗	✓			✓	2009-2010
Benzene*				✗	✓	✗			✓	2017-2018
Tetrachloroethylene*				✗	✓	✗	✗		✓	2017-2018
Styrene*	✗	✓	✗	✗	✓	✗			✗	2015-2016
1,2-Propylene oxide**	✗	✗	✗	✗	✗	✗	✓		✓	2015-2016
Cadmium	✗	✗	✓	✗	✗	✗	✓		✗	2017-2018
Ethylbenzene*	✗	✓	✗	✗	✓	✗	✗		✗	2017-2018
Diethanolamine							✓		✓	
1,2,3,4,5,6-Hexachlorocyclohexane**				✗	✓	✗				2009-2010
Dieldrin				✗	✓	✗				2003-2004
Diisononyl phthalate				✗	✓	✗				2017-2018
N'-Nitrosornicotine				✗	✓	✗				2013-2014
Pentachlorophenol*				✗	✓	✗			✗	2009-2010
Bromoform				✗	✓	✗		✗	✗	2017-2018
Tris(2-chloroethyl) phosphate**	✗	✗	✗	✗	✓	✗				2015-2016

Continued, next page...

Table A21: Extended synthesis: carcinogens with evidence of potential exposure disparities

Chemical	NHANES Avg			NHANES P95			HHP	DW	RSEI	NHANES Cycle
	Prim.	Int.	Age	Prim.	Int.	Age				
2,6-Dimethylaniline	×	×	×	×	✓	×			×	2013-2014

Notes: Cross-analysis summary of chemical-specific exposure disparities, including extended set of NHANES demographic subgroups and based on ratio inclusion threshold value of 1.5. ✓: evidence of potential exposure disparities for at least of the analysis's primary subgroups. ×: no evidence of potential disparities in exposure. Chemicals listed in first column are parent carcinogens. NHANES results may be based on metabolite or pooled concentrations. **: Carcinogens for which only metabolite biomonitoring results were available. *: Carcinogens for which both direct and metabolite biomonitoring results were available. †: NHANES average analysis based on pooled data. For NHANES biomonitoring subcolumns, *Prim.* denotes results from the primary analyses (Tables A1, A4, A7), *Int.* denotes results from the demographic interaction analyses (Tables A2, A5, A8), and *Age* denotes results from the age-differentiated analyses (Tables A3, A6, A9). *HHP* denotes results from the household purchasing analysis. *DW* denotes results from the drinking water analysis. *RSEI* denotes results from the air toxics analysis. *NHANES Cycle* notes the biomonitoring data cycle years used to calculate chemical-specific statistics.

Parent Chems: {N-Acetyl-S-(1-cyano-2-hydroxyethyl)-L-cysteine, N-acetyl-S-(2-cyanoethyl)-L-cysteine, N-acetyl-S-(2-Hydroxyethyl)-L-cysteine: *Acrylonitrile***}; {2,5-dichlorophenol: *1,4-Dichlorobenzene**}; {2,4-dichlorophenol: *2,4-Dichlorophenoxyacetic acid**}; {N-acetyl-S-(2-Hydroxyethyl)-L-cysteine: *Ethylene oxide**}; {2,4-dichlorophenol: *Nitrofen***}; {N-acetyl-S-(2-carbamoyl-ethyl)-L-cysteine: *Acrylamide**}; {1-naphthol: *1-Naphthalenol*, *1-(N-methylcarbamate)***}; {1-naphthol: *Naphthalene***}; {N-acetyl-S-(2-Carboxyethyl)-L-cysteine, N-acetyl-S-(3-Hydroxypropyl)-L-cysteine: *Acrolein***}; {N-acetyl-s-(3-hydroxypropyl-1-methyl)-L-cysteine: *Crotonaldehyde**}; {Mono-(2-ethyl-5-hydroxyhexyl) phthalate, Mono-(2-ethyl-5-oxohexyl) phthalate, Mono-2-ethyl-5-carboxypentyl phthalate: *Di(2-ethylhexyl) phthalate***}; {Oxypyrimidine: *Diazinon***}; {N-acetyl-S-(2-Hydroxyethyl)-L-cysteine: *Vinyl chloride***}; {NNAL: *4-(N-Methyl-N-nitrosamino)-1-(3-pyridyl)-1-butanone***}; {Dimethyl phosphate: *Dichlorvos***}; {Dimethyl phosphate: *Malathion***}; {Dimethyl phosphate: *Z-Tetrachlorvinphos***}; {N-acetyl-S-(2-hydroxy-3-butenyl)-L-cysteine: *1,3-Butadiene***}; {N-Acetyl-S-(n-propyl)-L-cysteine: *1-Bromopropane***}; {N-Acetyl-S-(phenyl)-L-cysteine: *Benzene**}; {N-acetyl-s-(trichlorovinyl)-L-cysteine: *Tetrachloroethylene**}; {Mandelic acid: *Styrene**}; {N-acetyl-S-(2-hydroxypropyl)-L-cysteine: *1,2-Propylene oxide***}; {Mandelic acid: *Ethylbenzene**}; {2,4,5-trichlorophenol: *1,2,3,4,5,6-Hexachlorocyclohexane***}; {2,4,5-trichlorophenol: *Pentachlorophenol**}; {Bis(2-chloroethyl) phosphate: *Tris(2-chloroethyl) phosphate***}

Return to: Section 3.3, Section 3.4.

Table A22: Extended synthesis: alternative inclusion ratio threshold of 1.25

Chemical	NHANES Avg			NHANES P95			HHP	DW	RSEI	NHANES Cycle
	Prim.	Int.	Age	Prim.	Int.	Age				
Arsenic	✓	✓	✓	✓	✓	✓	✓	✓	✓	2015-2016
1,4-Dichlorobenzene*	✓	✓	✓	✓	✓	✓	✓		✓	2017-2018
Di(2-ethylhexyl) phthalate**	✓	✓	✓	✓	✓	✓	✓		✓	2017-2018
Ethylene oxide*	✓	✓	✓	✓	✓	✓	✓		✓	2015-2016
1-Bromopropane**	✓	✓	✓	✓	✓	✓			✓	2015-2016
1-Naphthylamine	✓	✓	✓	✓	✓	✓			✓	2013-2014
4-Biphenylamine	✓	✓	✓	✓	✓	✓			✓	2013-2014
Acrylamide*	✓	✓	✓	✓	✓	✓			✓	2015-2016
Acrylonitrile**	✓	✓	✓	✓	✓	✓			✓	2015-2016
Crotonaldehyde*	✓	✓	✓	✓	✓	✓			✓	2015-2016
Melamine	✓	✓	✓	✓	✓	✓	✓			2003-2004
Styrene*	✓	✓	✓	✓	✓	✓			✓	2015-2016
1-Naphthalenol, 1-(N-methylcarbamate)**	✓	✓	✓	✓	✓	✓	✗		✓	2015-2016
2,4-Dichlorophenoxyacetic acid*	✓	✓	✓	✓	✓	✓	✗		✓	2015-2016
Ethylbenzene*	✓	✓	✓	✓	✓	✓	✗		✓	2017-2018
Naphthalene**	✓	✓	✓	✓	✓	✓	✗		✓	2015-2016
2-Naphthylamine	✓	✓	✓	✓	✓	✓				2013-2014
Dimethylarsinic acid	✓	✓	✓	✓	✓	✓				2017-2018
Nitrofen**	✓	✓	✓	✓	✓	✓				2015-2016
1,3-Butadiene**	✗	✓	✓	✓	✓	✓			✓	2015-2016
2-Methylaniline	✓	✓	✓	✗	✓	✓			✓	2013-2014
Acrolein**	✓	✓	✓	✓	✓	✓			✗	2015-2016
Dichlorvos**	✗	✓	✓	✓	✓	✓			✓	2017-2018
Malathion**	✗	✓	✓	✓	✓	✓			✓	2017-2018
Perfluorooctanesulfonic acid	✓	✓	✗	✓	✓	✓			✓	2017-2018
1,2-Propylene oxide**	✓	✓	✓	✗	✓	✗	✓		✓	2015-2016
Cadmium	✗	✓	✓	✗	✓	✓	✓		✓	2017-2018
Bromoform				✓	✓	✓		✓	✓	2017-2018
Furan				✓	✓	✓	✓		✓	2017-2018
4-(N-Methyl-N-nitrosamino)-1-(3-pyridyl)-1-butanone**	✓	✓	✓	✓	✓	✗				2013-2014
Tris(2-chloroethyl) phosphate**	✗	✓	✓	✓	✓	✓				2015-2016
Z-Tetrachlorvinphos**	✗	✓	✓	✓	✓	✓				2017-2018
2,6-Dimethylaniline	✓	✓	✓	✗	✓	✗			✓	2013-2014
Lead	✗	✓	✓	✗	✓	✓	✗		✓	2017-2018
2,4,6-Trichlorophenol				✓	✓	✓			✓	2009-2010
2-Phenylphenol				✓	✓	✓			✓	2009-2010
Benzene*				✓	✓	✓			✓	2017-2018
Diazinon**				✓	✓	✓			✓	2013-2014
Ethylene thiourea				✓	✓	✓			✓	2007-2008
PCBs†	✓	✓	✓						✓	2015-2016
Vinyl chloride**				✓	✓	✓			✓	2015-2016
Chloroform				✓	✓	✓		✗	✓	2017-2018
N,N-Dimethylformamide**	✗	✗	✓	✗	✓	✓			✓	2015-2016
Perfluorooctanoic acid	✗	✗	✗	✓	✓	✓			✓	2017-2018
Nickel	✗	✓	✓	✗	✗	✓	✗		✓	2017-2018
2-Amino-alpha-carboline				✓	✓	✓				2013-2014
Beta-Hexachlorocyclohexane†	✓	✓	✓							2015-2016
DDT†	✓	✓	✓							2015-2016
Dieldrin				✓	✓	✓				2003-2004
Diisononyl phthalate				✓	✓	✓				2017-2018
MeA-alpha-C				✓	✓	✓				2013-2014
N'-Nitrosonornicotine				✓	✓	✓				2013-2014

Continued, next page...

Table A22: Extended synthesis: alternative inclusion ratio threshold of 1.25

Chemical	NHANES Avg			NHANES P95			HHP	DW	RSEI	NHANES Cycle
	Prim.	Int.	Age	Prim.	Int.	Age				
PBB153†	✓	✓	✓							2015-2016
Pentachlorophenol*				✓	✓	✓			×	2009-2010
Tetrachloroethylene*				×	✓	×	✓		✓	2017-2018
Nitromethane	×	×	×	×	✓	✓			✓	2011-2012
4-Methyl-2-pentanone							✓		✓	2017-2018
1,4-Dioxane							✓	×	✓	2015-2016
4,4'-Diaminobiphenyl methane				×	×	✓			✓	2015-2016
Diethanolamine							✓		✓	
Diuron							✓		✓	
1,2,3,4,5,6-Hexachlorocyclohexane**				×	✓	×				2009-2010
2-Amino-1-methyl-6-phenylimidazo [4,5-b]pyridine				×	✓	×				2013-2014
Heptachlor epoxide B				×	✓	×				2003-2004

Notes: Cross-analysis summary of chemical-specific exposure disparities, including extended set of NHANES demographic subgroups and based on alternative ratio inclusion threshold value of 1.25. ✓: evidence of potential exposure disparities for at least of the analysis's primary subgroups. ×: no evidence of potential disparities in exposure. Chemicals listed in first column are parent carcinogens. NHANES results may be based on metabolite or pooled concentrations. **: Carcinogens for which only metabolite biomonitoring results were available. *: Carcinogens for which both direct and metabolite biomonitoring results were available. †: NHANES average analysis based on pooled data. For NHANES biomonitoring subcolumns, *Prim.* denotes results from the primary analyses (Tables A1, A4, A7), *Int.* denotes results from the demographic interaction analyses (Tables A2, A5, A8), and *Age* denotes results from the age-differentiated analyses (Tables A3, A6, A9). *HHP* denotes results from the household purchasing analysis. *DW* denotes results from the drinking water analysis. *RSEI* denotes results from the air toxics analysis. *NHANES Cycle* notes the biomonitoring data cycle years used to calculate chemical-specific statistics.

Parent Chems: {2,5-dichlorophenol: *1,4-Dichlorobenzene**}; {Mono-(2-ethyl-5-hydroxyhexyl) phthalate, Mono-(2-ethyl-5-oxohexyl) phthalate, Mono-2-ethyl-5-carboxypentyl phthalate: *Di(2-ethylhexyl) phthalate***}; {N-acetyl-S-(2-Hydroxyethyl)-L-cysteine: *Ethylene oxide**}; {N-Acetyl-S-(n-propyl)-L-cysteine: *1-Bromopropane***}; {N-acetyl-S-(2-carbamoyl-ethyl)-L-cysteine: *Acrylamide**}; {N-Acetyl-S-(1-cyano-2-hydroxyethyl)-L-cysteine, N-acetyl-S-(2-cyanoethyl)-L-cysteine, N-acetyl-S-(2-Hydroxyethyl)-L-cysteine: *Acrylonitrile***}; {N-acetyl-s-(3-hydroxypropyl-1-methyl)-L-cysteine: *Crotonaldehyde**}; {Mandelic acid: *Styrene**}; {1-naphthol: *1-Naphthalenol, 1-(N-methylcarbamate)***}; {2,4-dichlorophenol: *2,4-Dichlorophenoxyacetic acid**}; {Mandelic acid: *Ethylbenzene**}; {1-naphthol: *Naphthalene***}; {2,4-dichlorophenol: *Nitrofen***}; {N-acetyl-S-(2-hydroxy-3-butenyl)-L-cysteine: *1,3-Butadiene***}; {N-acetyl-S-(2-Carboxyethyl)-L-cysteine, N-acetyl-S-(3-Hydroxypropyl)-L-cysteine: *Acrolein***}; {Dimethyl phosphate: *Dichlorvos***}; {Dimethyl phosphate: *Malathion***}; {N-acetyl-S-(2-hydroxypropyl)-L-cysteine: *1,2-Propylene oxide***}; {NNAL: *4-(N-Methyl-N-nitrosamino)-1-(3-pyridyl)-1-butanone***}; {Bis(2-chloroethyl) phosphate: *Tris(2-chloroethyl) phosphate***}; {Dimethyl phosphate: *Z-Tetrachlorvinphos***}; {N-Acetyl-S-(phenyl)-L-cysteine: *Benzene**}; {Oxypyrimidine: *Diazinon***}; {N-acetyl-S-(2-Hydroxyethyl)-L-cysteine: *Vinyl chloride***}; {N-acetyl-S-(N-methylcarbamoyl)-L-cysteine: *N,N-Dimethylformamide***}; {2,4,5-trichlorophenol: *Pentachlorophenol**}; {N-acetyl-s-(trichlorovinyl)-L-cysteine: *Tetrachloroethylene**}; {2,4,5-trichlorophenol: *1,2,3,4,5,6-Hexachlorocyclohexane***}

Return to: Section 3.3, Section 3.4.

Table A23: Extended synthesis: chemicals with evidence of no exposure disparities

Chemical	NHANES Avg			NHANES P95			HHP	DW	RSEI	NHANES Cycle
	Prim.	Int.	Age	Prim.	Int.	Age				
Formaldehyde	×	×	×	×	×	×	×		×	2015-2016
Arsenic acid				×	×	×				2017-2018
Methylarsonic acid				×	×	×				2017-2018
PeCDF (furan)†	×	×	×							2015-2016
1,3-Propane sultone									×	
2,3,7,8-Tetrachlorodibenzo-p-dioxin									×	2015-2016
Bromate								×		
Butylated hydroxyanisole							×			
Dichloroacetic acid								×		
Ethanol							×			
N-Nitrosodimethylamine								×		
Titanium dioxide							×			
Trichloroethylene									×	2017-2018

Notes: This table lists the carcinogens which, across all applicable analyses, provide suggestive evidence showing limited demographic heterogeneity in exposure. ×: denotes an analysis showing no substantive evidence of exposure disparities, based on the *more lenient, alternative inclusion ratio threshold value of 1.25*. For NHANES biomonitoring subcolumns, *Prim.* denotes results from the primary analyses (Tables A1, A4, A7), *Int.* denotes results from the demographic interaction analyses (Tables A2, A5, A8), and *Age* denotes results from the age-differentiated analyses (Tables A3, A6, A9). *HHP* denotes results from the household purchasing analysis. *DW* denotes results from the drinking water analysis. *RSEI* denotes results from the air toxics analysis. *NHANES Cycle* notes the biomonitoring data cycle years used to calculate chemical-specific statistics.

Return to: Section 3.3.

Table A24: National cancer mortality by race/ethnicity and sex

Cancer Type													
		Female						Male					
		All	Amer Ind	Asian/PI	Female Black	Female Hisp	Female White	All	Amer Ind	Asian/PI	Male Black	Male Hisp	Male White
Brain and Other Nervous System		3.6	2.1	1.9	2.3	2.7	4.1	5.4	2.9	2.8	3.3	3.7	6.2
Breast		19.6	14.9	11.9	27.2	13.7	19.7	0.3		0.1	0.5	0.1	0.3
Cervix Uteri		2.2	2.6	1.7	3.3	2.4	2.1		16				
Colon and Rectum		10.9	11.2	7.8	13.9	8.6	11	15.6		10.9	21.9	13.6	15.4
Corpus Uteri		3.0	2.1	2	5.1	2.3	2.8						
Esophagus		1.4	1.3	0.7	1.5	0.7	1.5	6.6	5.1	2.6	4.6	3.5	7.6
Gallbladder		0.6	0.9	0.7	1	1	0.5	0.4	0.6	0.5	0.7	0.5	0.3
Kidney and Renal Pelvis		2.2	3	1	2.1	2.1	2.2	5.1	6.8	2.3	5	4.7	5.3
Larynx		0.3	0.4	0.1	0.5	0.1	0.4	1.6	1.3	0.6	2.8	1.2	1.6
Leukemias		4.5	3.2	2.6	4.2	3.5	4.7	8.0	4.7	4.5	6.6	5.3	8.6
Liver		2.4	4.6	3.3	3	4.1	2.1	7.4	11.3	9.3	10.4	10.7	6.4
Lung and Bronchus		28.6	26.3	15.3	26.9	11.3	32.1	40.6	34.3	24.7	48.9	20.3	43.2
Melanoma of the Skin		1.3	0.4	0.3	0.3	0.5	1.7	3.0	1	0.3	0.4	0.9	3.8
Myeloma		2.4	2.1	1.2	5	2.1	2.2	3.9	3.2	1.9	7.4	3.1	3.7
Non-Hodgkin Lymphoma		3.8	2.7	2.7	2.9	3.6	4	6.7	5	4.7	5.1	5.5	7.1
Oral Cavity and Pharynx		1.4	1.1	1.2	1.2	0.8	1.5	4.0	2.7	3.1	4.3	2.4	4.2
Ovary		6.2	5.1	4.4	5.7	4.9	6.5						
Pancreas		9.8	7.6	7.2	12.4	8.2	9.8	12.9	9.3	8.4	15.4	9.6	13.3
Prostate								19.2	15.2	8.8	37.5	15.6	18.2
Stomach		2.1	2.8	3.5	3.4	3.8	1.5	3.7	5.3	5.6	6.9	5.9	2.9
Thyroid		0.5	0.4	0.6	0.5	0.7	0.4	0.5	0.4	0.5	0.4	0.5	0.5
Urinary Bladder		2.0	1.4	0.9	2.2	1.3	2.2	7.2	3.8	2.8	5.3	3.9	8
Demographic subgroup:													
Sex:		Female	Female	Female	Female	Female	Female	Male	Male	Male	Male	Male	Male
Race/Ethnicity:		All	Amer Ind	Asian/PI	Female Black	Female Hisp	Female White	All	Amer Ind	Asian/PI	Male Black	Male Hisp	Male White

Notes: Table presents national cancer mortality rates (per 100,000 population). Data obtained from CDC, 2021b in October 2024, and are based on mortality during the years 2017-2021. Cancer sites presented in this table include all leading and invasive cancer types. All statistics are age-adjusted rates, based on 2000 U.S. standard population. Race/ethnicity subgroups are: non-Hispanic American Indian and Alaska Native, non-Hispanic Asian and Pacific Islander, non-Hispanic Black, Hispanic, non-Hispanic White. **Bold font** designates subgroup-specific incidence rates that are 25% or more above the corresponding national average. Underlined values denote concentration ratios subgroup-specific incidence rates that are 50% or more above the corresponding national average.

Return to: Section 3.5.

B Exposure pathways for known carcinogens with scarce biomonitoring

In this appendix, we conducted a supplement assessment of common exposure pathways and product types for *known carcinogens*. Though a chemical’s status as an IARC- or ROC-listed known carcinogen may influence the creation of a health-protective regulation to address exposure (Mehta et al., 2023), to our knowledge, previous or current national-level exposure surveillance is only available for 18 of the 112 known carcinogens.¹ To explore a path forward in identifying potential exposure disparities for the many known carcinogens lacking biomonitoring or indirect measures of exposure, we summarized what is known regarding exposure pathways and major product categories for the subset of chemicals listed as known carcinogens. More specifically, using identical data and methods to the review exercise conducted in section 3.4, we individually researched the 112 known carcinogens to ascertain readily-available information on their common exposure pathways, product categories, exposure types, and associated cancer sites. In what follows, we highlighted exposure pathways and product categories that were most common for the broader set of known carcinogens, and noted the carcinogens known to be associated with cancer types that differentially impact demographic subgroups.

Figure B1 and Figure B2 visually summarize general exposure pathway and product category trends for this set of chemicals. The two most common exposure pathways across chemicals with known carcinogenicity were via manufacturing (78 of 112 chemicals) and materials processing (71/112). As one would expect, these two exposure pathways are largely occupational. The set of known carcinogens with industrial exposure pathways included several dozen heavy metals and asbestos varieties; some of these carcinogens are still actively used, while others are now legacy contaminants. Other common occupational exposure pathways included construction (22/112), farming/agriculture (22/112), medical/dental (21/112), and mining (20/112).

Our review found that the general population was exposed to 53 of 112 known carcinogen via environmental releases (either ongoing or former) in the soil, air or surface water. And perhaps logically, we observed a strong correlation between a known carcinogen having potential for general population exposure via environmental releases and potential for occupational exposure via manufacturing and/or processing. Other common general exposure pathways included food/food additives (42/112), medical/dental (36/112), drinking water (34/112), and art/office supplies (33/112).

We next considered the set of key product categories in which known carcinogens were found to occur. The most commonly occurring product category (53 of 112 known carcinogens) was metals and electronics. As described above, many known carcinogens are heavy metal compounds, which were used directly or as an intermediate in larger industrial or electronic processes. The second most commonly occurring product category was pharmaceuticals; 43 of 112 known carcinogens were used in pharmaceutical products. The reason for this is fairly intuitive: several hormones and/or chemicals are carcinogenic, but have potential benefits via therapeutic use in treating other conditions. These pharmaceutical uses included as a chemotherapy, an immunosuppressant, a hormone replacement therapies, or as a birth control. Other common product categories included paints/adhesives (39/112), mining (32/112), and plastics/rubber (32/112).

We next considered the subset of 16 known carcinogens that our review found to be associated with liver cancer. Recall from Section 3.5 that national statistics showed elevated incidence of liver cancer among both females and males of several racial and ethnic subgroups. These 16 known carcinogens vary widely in

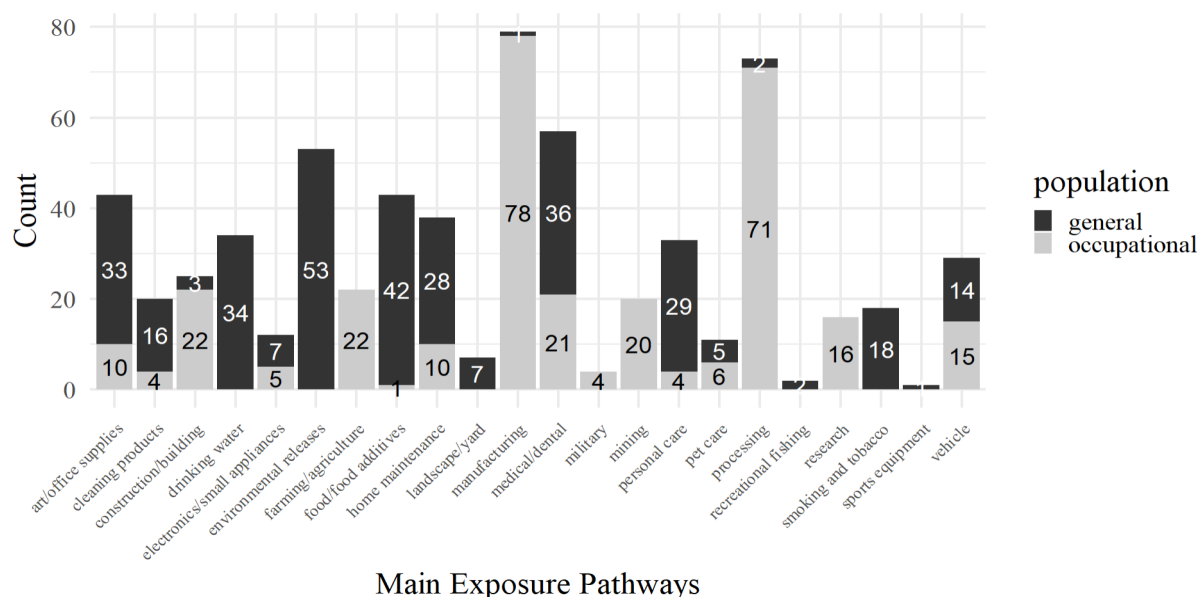
¹See Table B1 for more information on exposure pathways, associated cancer types, and biomonitoring availability for each of the known carcinogens.

chemical structure, though arsenic is a component of nearly half of them. Common exposure pathways among these 16 chemicals included food and food additives (general population; 10/16), drinking water (general population; 10/16), and manufacturing and processing (occupational; 12/16). Though we've strictly noted correlations here, these 16 chemicals and the noted pathways might prove promising starting points for further exploration of potential exposure disparities.

We performed a similar logical extrapolation for two other cancer types: kidney and the renal system (for which national incidence was especially elevated for American Indian individuals), and prostate (for which Black males had highly elevated incidence). Thirteen known carcinogens were found to be associated with renal cancers; again a large fraction of them (6/13) are chemicals containing arsenic. The most common exposure pathways were manufacturing and processing (occupational; 11 of 13 carcinogens), food and food additives (general; 6/13), pet care (general/occupational; 6/13), and art/office supplies (general/occupational; 6/13). Five known carcinogens were found to be associated with prostate cancer: cadmium, coal tar, diethylstilbestrol, lindane, and cadmium chloride. Four chemicals shared a common medical/dental general exposure pathway, and three chemicals shared drinking water and food/food additive general exposure pathways. While this information is fairly limited and again, strictly built from correlations, these chemicals and exposure pathways might merit further exploration as research on exposure disparities continues.

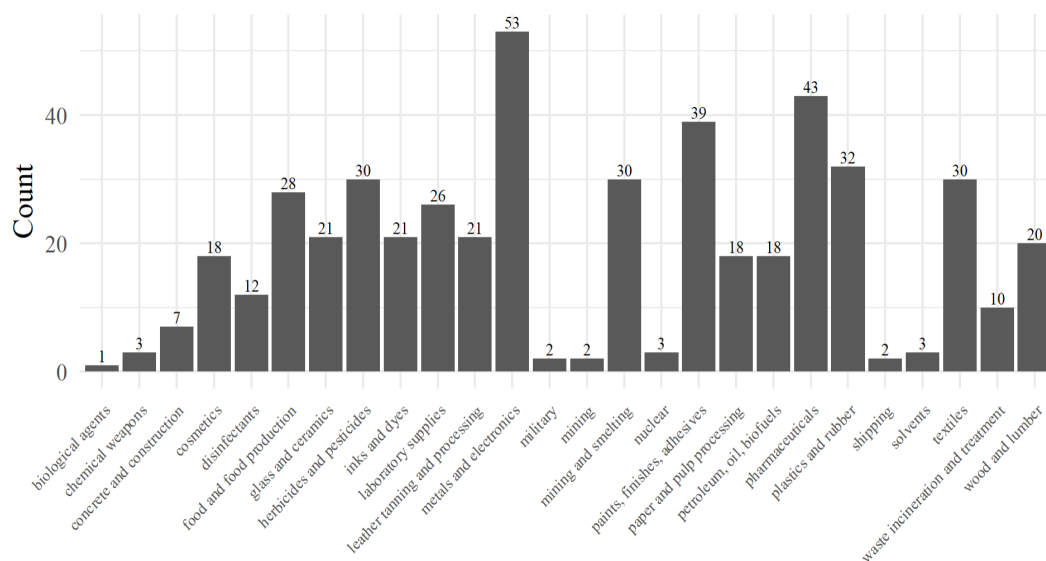
Figure B1: Known carcinogens: likely pathways and product categories

(a) Frequency of identified uses (general/occupational) amongst known carcinogens



Main Exposure Pathways

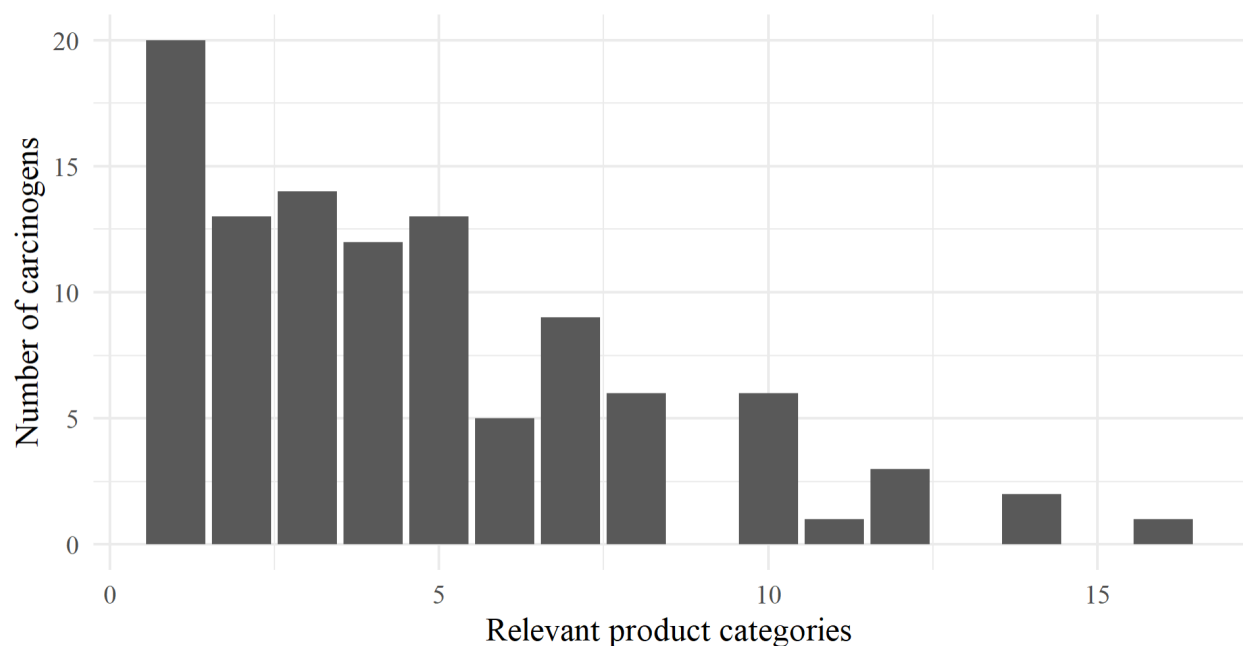
(b) Product category frequency amongst known carcinogens



Major Product Categories

Notes: Panel (a) shows the prevalence of exposure pathways across chemicals known to be carcinogenic, distinguishing between general population and occupational pathways. Panel (b) presents the count of known carcinogens (out of N=112) which were found to be associated with the given product category. See Section 2.4 and Section 3.4 for description of exposure pathway information collection and product categorization methods.

Figure B2: Exposure sourcing for known carcinogens: number of identified product categories in which each chemical is used



Notes: Histogram of the count of associated product categories for each known carcinogen ($N = 112$). For example, twenty known carcinogens are associated with a single product category, thirteen known carcinogens are associated with two different product categories, ..., and one carcinogen is associated with sixteen different product categories. See Section 2.4 and Section 3.4 for description of exposure pathway information collection and product categorization methods.

Table B1: Exposure pathways and cancer types for known carcinogens

Chemical	Major Prod. Categories	Exp. route	Associated cancers	Occupational exposure	General exposure	NHANES data
1,3-Butadiene	Plastics and Rubber;	Inhalation;	Leukemia; Non-Hodgkin Lymphoma;	Manufacturing; Vehicle	Environmental Releases; Smoking and Tobacco; Vehicle	✓
2,3,7,8-Tetrachlorodibenzo-P-Dioxin	Waste Incineration and Treatment; Paper and Pulp Processing; Herbicides and Pesticides;	Dermal; Inhalation; Ingestion;	Lung; Non-Hodgkin Lymphoma; Thyroid;	Processing; Research	Environmental Releases	
2-Methylaniline	Textiles; Pharmaceuticals; Herbicides and Pesticides;	Dermal; Inhalation; Ingestion;	Bladder;	Manufacturing; Research	Personal Care; Smoking and Tobacco	✓
2-Naphthylamine	Plastics and Rubber; Textiles;	Dermal; Inhalation; Ingestion;	Bladder;	Manufacturing	Cleaning Products; Smoking and Tobacco; Environmental Releases	✓
4-Biphenylamine	Textiles; Plastics and Rubber;	Dermal; Inhalation;	Bladder;	Manufacturing; Research	Smoking and Tobacco; Food/ Food Additives; Personal Care	✓
8-Methoxypsoralen	Cosmetics;	Dermal; Ingestion;	Skin;	Manufacturing	Personal Care; Medical/Dental; Food/ Food Additives	
Aflatoxin	Biological Agents; Food and Food Production;	Inhalation; Ingestion; Dermal;	Liver;	Farming/ Agriculture	Food/ Food Additives	
Aristolochic Acid	Pharmaceuticals;	Ingestion;	Renal Pelvis and Ureter; Bladder;	NA	Personal Care; Food/ Food Additives	
Arsenic	Mining and Smelting; Herbicides and Pesticides; Metals and Electronics;	Ingestion;	Bladder; Skin; Lung; Stomach / Digestive; Liver; Kidney; Lymphatic; Hemopoietic;	Manufacturing; Processing; Vehicle; Farming/ Agriculture; Medical/Dental	Electronics/ Small Appliances; Cleaning Products; Art/ Office Supplies; Home Maintenance; Vehicle; Food/ Food Additives; Smoking and Tobacco; Drinking Water; Environmental Releases	✓
Azathioprine	Pharmaceuticals;	Ingestion; Injection; Inhalation;	Skin; Non-Hodgkin Lymphoma; Leukemia; Liver; Gallbladder;	Manufacturing; Medical/Dental	NA	

Continued, next page...

Table B1: Exposure pathways and cancer types for known carcinogens

Chemical	Major Prod. Categories	Exp. route	Associated cancers	Occupational exposure	General exposure	NHANES data
Benzene	Solvents; Petroleum, Oil, Biofuels; Pharmaceuticals; Textiles; Plastics and Rubber; Inks and Dyes;	Inhalation; Ingestion; Dermal;	Leukemia; Hodgkin's Lymphoma;	Manufacturing; Vehicle; Processing	Vehicle; Smoking and Tobacco; Art/ Office Supplies; Cleaning Products; Home Maintenance; Landscape/ Yard; Food/ Food Additives; Environmental Releases; Drinking Water	✓
Beryllium	Metals and Electronics; Mining and Smelting;	Inhalation; Ingestion;	Lung;	Processing; Minings; Manufacturing	Environmental Releases; Smoking and Tobacco; Drinking Water	✓
Bis-(2-Chloroethyl)Sulfide	Chemical Weapons;	Inhalation; Dermal;	Lung;	Military	Environmental Releases; Recreational Fishing	
Busulfan	Pharmaceuticals;	Ingestion;	Leukemia;	Manufacturing; Medical/ Dental	Medical/ Dental	
C.i. Azoic Diazo Component 112	Leather Tanning and Processing; Paper and Pulp Processing; Textiles; Laboratory Supplies;	Inhalation; Ingestion; Dermal;	Bladder;	Processing; Research	Home Maintenance; Environmental Releases; Drinking Water; Food/ Food Additives	
Cadmium	Mining and Smelting; Metals and Electronics; Paints, Finishes, Adhesives; Textiles; Plastics and Rubber;	Ingestion; Inhalation;	Lung; Prostate; Bladder; Kidney; Pancreatic; Breast;	Processing; Minings; Manufacturing	Food/ Food Additives; Drinking Water; Smoking and Tobacco; Art/ Office Supplies; Recreational Fishing; Environmental Releases; Medical/ Dental	✓
Chlorambucil	Pharmaceuticals;	Ingestion; Inhalation; Dermal;	Leukemia;	Medical/ Dental	Medical/ Dental; Manufacturing; Processing; Environmental Releases	
Chromium (VI) Ion	Metals and Electronics; Paints, Finishes, Adhesives; Textiles; Leather Tanning and Processing;	Ingestion; Inhalation; Dermal;	Lung; Sinus/Nasal;	Construction/ Building; Processing; Vehicle	Home Maintenance; Art/ Office Supplies; Environmental Releases; Drinking Water	
Coal Tar	Pharmaceuticals; Metals and Electronics; Herbicides and Pesticides; Paints, Finishes, Adhesives; Plastics and Rubber;	Ingestion; Inhalation; Dermal;	Skin; Lung; Leukemia; Digestive; Bladder; Kidney; Oral; Stomach; Prostate;	Processing; Construction/ Building; Manufacturing	Personal Care; Cleaning Products; Medical/ Dental; Environmental Releases; Pet Care	

Continued, next page...

Table B1: Exposure pathways and cancer types for known carcinogens

Chemical	Major Prod. Categories	Exp. route	Associated cancers	Occupational exposure	General exposure	NHANES data
Diethylstilbestrol	Pharmaceuticals;	Ingestion; Injection;	Cervix; Prostate; Breast; Pancreatic;	NA	Medical/ Dental	
Erionite	Mining and Smelting; Food and Food Production; Waste Incineration and Treatment;	Inhalation; Ingestion;	Lung;	Farming/ Agriculture; Construction/ Building; Mining; Processing; Food/ Food Additives	Home Maintenance; Environmental Releases; Pet Care; Food/ Food Additives	
Ethylene Oxide	Laboratory Supplies; Herbicides and Pesticides; Food and Food Production; Disinfectants; Cosmetics;	Inhalation; Ingestion;	Leukemia; Stomach; Lymphatic;	Medical/ Dental; Manufacturing; Farming/ Agriculture	Environmental Releases; Smoking and Tobacco; Personal Care; Home Maintenance; Food/ Food Additives; Vehicle	✓
Formaldehyde	Herbicides and Pesticides; Disinfectants; Laboratory Supplies; Food and Food Production; Plastics and Rubber; Paints, Finishes, Adhesives; Metals and Electronics; Leather Tanning and Processing; Paper and Pulp Processing; Textiles;	Inhalation; Dermal;	Leukemia; Nasal/Sinus;	Manufacturing; Research; Medical/ Dental; Farming/ Agriculture	Smoking and Tobacco; Food/ Food Additives; Home Maintenance; Environmental Releases; Vehicle; Cleaning Products; Art/ Office Supplies	✓
Melphalan	Pharmaceuticals; Chemical Weapons; Herbicides and Pesticides;	Injection; Ingestion; Inhalation; Dermal;	Leukemia;	Medical/ Dental; Manufacturing	Medical/ Dental; Environmental Releases	
Phenacetin	Pharmaceuticals; Cosmetics; Laboratory Supplies;	Ingestion; Inhalation; Dermal;	Kidney;	Pet Care; Medical/ Dental; Manufacturing	Pet Care; Medical/ Dental; Personal Care	
Pitch, Coal Tar, High-Temp.	Metals and Electronics; Paints, Finishes, Adhesives; Textiles; Leather Tanning and Processing; Pharmaceuticals;	Inhalation; Dermal;	Lung; Kidney; Skin;	Manufacturing; Construction/ Building; Processing; Vehicle; Cleaning Products	Art/ Office Supplies; Medical/ Dental; Home Maintenance; Vehicle; Cleaning Products	

Continued, next page...

Table B1: Exposure pathways and cancer types for known carcinogens

Chemical	Major Prod. Categories	Exp. route	Associated cancers	Occupational exposure	General exposure	NHANES data
Quartz (SiO ₂)	Food and Food Production; Cosmetics; Metals and Electronics; Petroleum, Oil, Biofuels; Mining and Smelting; Glass and Ceramics; Plastics and Rubber; Paper and Pulp Processing; Paints, Finishes, Adhesives; Herbicides and Pesticides;	Inhalation; Dermal;	Lung;	Manufacturing; Construction/ Building; Processing; Cleaning Products; Vehicle; Farming/ Agriculture	Art/ Office Supplies; Medical/ Dental; Home Maintenance; Cleaning Products; Vehicle; Cleaning Products; Personal Care	
Radon	Pharmaceuticals; Petroleum, Oil, Biofuels; Mining and Smelting; Laboratory Supplies;	Radiation; Inhalation; Ingestion;	Lung; Salivary Glands; Stomach; Colon; Bladder; Ovary; Brain; Skin;	Medical/ Dental; Military; Mining; Research; Construction/ Building	Environmental Releases; Medical/ Dental; Environmental Releases; Drinking Water; Construction/ Building	
Semustine	Pharmaceuticals;	Dermal; Injection;	Lung; Leukemia;	Manufacturing; Medical/ Dental	Medical/ Dental	
Tamoxifen	Pharmaceuticals;	Injection; Inhalation;	Uterus;	Manufacturing; Processing	Medical/ Dental	
Thiotepa	Herbicides and Pesticides; Pharmaceuticals;	Injection; Ingestion; Inhalation; Dermal;	Leukemia;	Medical/ Dental; Manufacturing	Medical/ Dental; Environmental Releases	
Trichloroethylene	Metals and Electronics; Textiles; Paints, Finishes, Adhesives; Inks and Dyes; Plastics and Rubber; Wood and Lumber; Leather Tanning and Processing;	Inhalation; Dermal;	Liver; Kidney; Non-Hodgkin's Lymphoma;	Art/ Office Supplies; Manufacturing; Processing; Military	Art/ Office Supplies; Home Maintenance; Cleaning Products; Drinking Water; Environmental Releases; Food/ Food Additives	✓
Vinyl Chloride	Plastics and Rubber; Food and Food Production; Metals and Electronics; Wood and Lumber; Leather Tanning and Processing;	Ingestion; Inhalation; Dermal;	Liver; Brain; Lung; Lymphoma; Leukemia;	Manufacturing; Processing; Vehicle; Construction/ Building	Smoking and Tobacco; Environmental Releases; Drinking Water; Food/ Food Additives; Vehicle; Home Maintenance	✓
(⁻³² P)Phosphane	Pharmaceuticals;	Injection;	Leukemia;	NA	Medical/ Dental	

Continued, next page...

Table B1: Exposure pathways and cancer types for known carcinogens

Chemical	Major Prod. Categories	Exp. route	Associated cancers	Occupational exposure	General exposure	NHANES data
1,2-Dichloropropane	Paints, Finishes, Adhesives; Food and Food Production; Paper and Pulp Processing; Petroleum, Oil, Biofuels; Plastics and Rubber;	Ingestion; Inhalation; Dermal;	Biliary Tract;	Manufacturing; Processing; Farming/ Agriculture	Home Maintenance; Environmental Releases; Drinking Water	✓
2,3,4,7,8-Pentachlorodibenzofuran	Petroleum, Oil, Biofuels; Herbicides and Pesticides; Plastics and Rubber; Inks and Dyes; Waste Incineration and Treatment; Laboratory Supplies; Wood and Lumber;	Inhalation; Dermal; Ingestion;	Non-Hodgkin Lymphoma; Breast;	Manufacturing; Processing; Farming/ Agriculture; Construction/ Building; Vehicle	Environmental Releases; Food/ Food Additives; Drinking Water; Vehicle	✓
3,3',4,4',5-Pentachlorobiphenyl	Inks and Dyes; Paper and Pulp Processing; Paints, Finishes, Adhesives; Metals and Electronics; Herbicides and Pesticides; Laboratory Supplies; Waste Incineration and Treatment;	Ingestion;;	Melanoma; Liver; Bile-Duct; Gastro-Intestinal;	Research; Processing	Food/ Food Additives; Environmental Releases; Drinking Water	✓
4,4'-Methylenedis(2-Chloroaniline)	Paints, Finishes, Adhesives; Metals and Electronics; Plastics and Rubber; Food and Food Production;	Dermal; Inhalation; Ingestion;	Bladder;	Manufacturing; Vehicle	Food/ Food Additives; Environmental Releases	
Acetaldehyde	Food and Food Production; Cosmetics; Mining and Smelting; Paints, Finishes, Adhesives; Wood and Lumber; Herbicides and Pesticides; Inks and Dyes; Paper and Pulp Processing; Pharmaceuticals; Leather Tanning and Processing;	Inhalation; Ingestion; Dermal;	Lung; Oral-Cavity;	Mining; Construction/ Building; Processing; Manufacturing	Smoking and Tobacco; Food/ Food Additives; Cleaning Products; Art/ Office Supplies; Home Maintenance; Personal Care; Vehicle; Construction/ Building; Environmental Releases	

Continued, next page...

Table B1: Exposure pathways and cancer types for known carcinogens

Chemical	Major Prod. Categories	Exp. route	Associated cancers	Occupational exposure	General exposure	NHANES data
Benzo[a]Pyrene	Food and Food Production; Wood and Lumber; Metals and Electronics; Petroleum, Oil, Biofuels; Mining and Smelting; Glass and Ceramics; Laboratory Supplies; Inks and Dyes; Pharmaceuticals;	Inhalation; Ingestion; Dermal;	Lung; Skin;	Processing; Manufacturing; Mining; Research	Food/ Food Additives; Vehicle; Art/ Office Supplies; Smoking and Tobacco; Personal Care; Drinking Water; Environmental Releases; Home Maintenance	
Chlornaphazine	Pharmaceuticals;	Injection;	Bladder;	NA	Medical/ Dental	
Ethanol	Food and Food Production; Pharmaceuticals; Disinfectants; Solvents; Cosmetics; Petroleum, Oil, Biofuels; Paints, Finishes, Adhesives; Herbicides and Pesticides; Laboratory Supplies; Leather Tanning and Processing;	Ingestion; Inhalation; Dermal;	Mouth; Larynx; Pharynx; Esophagus; Cholorectal; Liver; Breast;	Manufacturing; Processing; Cleaning Products; Farming/ Agriculture; Medical/ Dental; Construction/ Building; Personal Care	Cleaning Products; Medical/ Dental; Environmental Releases; Food/ Food Additives; Drinking Water; Personal Care; Art/ Office Supplies	
Etoposide	Pharmaceuticals;	Injection; Ingestion; Dermal;	Leukemia;	Medical/ Dental	Medical/ Dental; Environmental Releases	
Lindane	Herbicides and Pesticides; Cosmetics; Pharmaceuticals; Laboratory Supplies;	Ingestion; Inhalation; Dermal;	Non-Hodgkin Lymphoma; Leukemia; Prostate; Testicular;	Farming/ Agriculture; Research	Personal Care; Medical/ Dental; Food/ Food Additives; Drinking Water; Environmental Releases	✓
Pentachlorophenol	Wood and Lumber; Disinfectants; Herbicides and Pesticides; Paints, Finishes, Adhesives; Leather Tanning and Processing; Paper and Pulp Processing;	Dermal; Inhalation; Ingestion;	Non-Hodgkin Lymphoma;	Processing; Manufacturing	Art/ Office Supplies; Food/ Food Additives; Drinking Water; Home Maintenance; Landscape/ Yard	✓
Plutonium	Metals and Electronics; Military; Nuclear;	Inhalation; Dermal;	Lung; Liver; Bone;	Research; Military; Processing; Research	Environmental Releases; Food/ Food Additives; Drinking Water	

Continued, next page...

Table B1: Exposure pathways and cancer types for known carcinogens

Chemical	Major Prod. Categories	Exp. route	Associated cancers	Occupational exposure	General exposure	NHANES data
Polychlorinated Biphenyls	Metals and Electronics; Paints, Finishes, Adhesives; Herbicides and Pesticides; Inks and Dyes; Food and Food Production; Laboratory Supplies; Plastics and Rubber; Waste Incineration and Treatment;	Inhalation;	Melanoma; Liver; Bile-Duct; Gastro-Intestinal;	Farming/ Agriculture; Manufacturing; Research	Environmental Releases; Drinking Water; Food/ Food Additives	✓
Radium-226	Mining and Smelting; Paints, Finishes, Adhesives; Pharmaceuticals; Metals and Electronics;	Ingestion; Inhalation;	Bone; Nasal;	Research; Medical/ Dental; Manufacturing; Processing	Medical/ Dental; Drinking Water; Food/ Food Additives; Environmental Releases	
Radium-228	Mining and Smelting; Paints, Finishes, Adhesives; Pharmaceuticals; Metals and Electronics;	Ingestion; Inhalation;	Bone; Nasal;	Research; Medical/ Dental; Manufacturing; Processing	Medical/ Dental; Drinking Water; Food/ Food Additives; Environmental Releases	
Shale Oils	Mining and Smelting;	Ingestion; Inhalation; Dermal;	Skin;	Manufacturing; Processing	NA	
Thorium	Glass and Ceramics; Metals and Electronics; Mining and Smelting; Food and Food Production; Nuclear;	Inhalation; Injection; Ingestion; Dermal;	Liver; Lung; Pancreatic; Bone;	Processing; Manufacturing	Drinking Water; Art/ Office Supplies; Electronics/ Small Appliances; Food/ Food Additives	
Treosulfan	Pharmaceuticals;	Ingestion;	Leukemia;	NA	Medical/ Dental	
17alpha-Ethinylestradiol	Pharmaceuticals; Cosmetics; Food and Food Production;	Ingestion; Dermal; Inhalation;	Breast; Ovarian;	Medical/ Dental; Manufacturing	Medical/ Dental; Food/ Food Additives; Drinking Water	
17beta-Estradiol	Pharmaceuticals; Cosmetics; Food and Food Production;	Ingestion; Dermal; Injection; Inhalation;	Breast; Ovarian;	Medical/ Dental; Manufacturing; Processing	Medical/ Dental; Personal Care; Food/ Food Additives; Drinking Water; Environmental Releases	

Continued, next page...

Table B1: Exposure pathways and cancer types for known carcinogens

Chemical	Major Prod. Categories	Exp. route	Associated cancers	Occupational exposure	General exposure	NHANES data
Actinolite Asbestos	Paints, Finishes, Adhesives; Paper and Pulp Processing; Concrete and Construction; Shipping; Mining and Smelting; Metals and Electronics; Plastics and Rubber; Waste Incineration and Treatment; Glass and Ceramics; Petroleum, Oil, Biofuels;	Inhalation; Ingestion;	Larynx; Mesothelioma; Lung; Ovarian; Stomach; Pharynx; Colorectum;	Home Maintenance; Manufacturing/ Building; Vehicle; Mining	Landscape/ Yard; Art/ Office Supplies; Personal Care; Environmental Releases	
Ammonium Chromate	Metals and Electronics; Textiles; Inks and Dyes; Laboratory Supplies;	Inhalation;	Lung; Sinus/Nasal;	Art/ Office Supplies; Manufacturing; Research	Art/ Office Supplies	
Ammonium Dichromate	Metals and Electronics; Inks and Dyes; Leather Tanning and Processing; Petroleum, Oil, Biofuels; Glass and Ceramics; Wood and Lumber; Paints, Finishes, Adhesives; Textiles; Food and Food Production; Cosmetics; Chemical Weapons; Herbicides and Pesticides;	Inhalation; Ingestion; Dermal;	Lung;	Art/ Office Supplies; Processing; Manufacturing	Art/ Office Supplies; Personal Care; Environmental Releases	
Amosite	Paints, Finishes, Adhesives; Paper and Pulp Processing; Concrete and Construction; Shipping; Mining and Smelting; Metals and Electronics; Plastics and Rubber; Waste Incineration and Treatment; Glass and Ceramics; Petroleum, Oil, Biofuels;	Inhalation; Ingestion;	Larynx; Mesothelioma; Lung; Ovarian; Stomach; Pharynx; Colorectum;	Home Maintenance; Manufacturing/ Building; Vehicle; Mining	Landscape/ Yard; Art/ Office Supplies; Personal Care; Environmental Releases	
Aristolochic Acids	Pharmaceuticals; Food and Food Production;	Ingestion; Dermal;	Urothelial;	Farming/ Agriculture	Medical/ Dental; Landscape/ Yard; Food/ Food Additives	

Continued, next page...

Table B1: Exposure pathways and cancer types for known carcinogens

Chemical	Major Prod. Categories	Exp. route	Associated cancers	Occupational exposure	General exposure	NHANES data
Arsenic Oxide (As ₂ O ₃)	Glass and Ceramics; Leather Tanning and Processing; Wood and Lumber; Mining and Smelting; Metals and Electronics; Textiles; Inks and Dyes; Mining and Smelting; Paints, Finishes, Adhesives; Food and Food Production; Herbicides and Pesticides; Pharmaceuticals;	Ingestion; Inhalation; Dermal;	Skin; Lung; Stomach; Liver; Bladder; Kidney; Lymphatic; Hematopoietic;	Processing; Manufacturing; Farming/ Agriculture; Mining	Food/ Food Additives; Art/ Office Supplies; Landscape/ Yard; Drinking Water; Smoking and Tobacco; Environmental Releases	
Asbestos	Metals and Electronics; Cosmetics; Plastics and Rubber; Concrete and Construction; Paints, Finishes, Adhesives; Laboratory Supplies; Textiles;	Ingestion; Inhalation;	Lung; Mesothelioma; Larynx;	Vehicle; Construction/ Building; Home Maintenance; Manufacturing	Vehicle; Personal Care; Electronics/ Small Appliances; Home Maintenance; Art/ Office Supplies	
Beryllium Carbonate Hydroxide (3/2/2)	Metals and Electronics; Mining and Smelting;	Inhalation; Dermal;	Lung;	Processing; Manufacturing; Mining	NA	
Beryllium Chloride	Mining and Smelting; Metals and Electronics;	Inhalation; Dermal;	Lung;	Processing	NA	
Beryllium Hydroxide	Metals and Electronics; Mining and Smelting;	Inhalation; Dermal;	Lung;	Processing	NA	
Beryllium Oxide (Beo)	Metals and Electronics; Military; Glass and Ceramics; Plastics and Rubber; Laboratory Supplies; Mining and Smelting; Paints, Finishes, Adhesives;	Inhalation; Dermal;	Lung;	Manufacturing; Processing; Mining; Electronics/ Small Appliances; Construction/ Building	NA	
Beryllium Sulfate	Glass and Ceramics; Metals and Electronics; Mining and Smelting;	Inhalation; Dermal;	Lung;	Manufacturing; Processing; Mining; Electronics/ Small Appliances; Construction/ Building	Art/ Office Supplies; Environmental Releases; Drinking Water	

Continued, next page...

Table B1: Exposure pathways and cancer types for known carcinogens

Chemical	Major Prod. Categories	Exp. route	Associated cancers	Occupational exposure	General exposure	NHANES data
C.i. Direct Black 38	Inks and Dyes; Textiles; Leather Tanning and Processing; Paper and Pulp Processing; Wood and Lumber; Laboratory Supplies; Plastics and Rubber;	Inhalation; Dermal; Ingestion;	Bladder;	Personal Care; Processing; Art/ Office Supplies; Manufacturing	Personal Care; Art/ Office Supplies; Environmental Releases	
C.i. Direct Blue 6	Inks and Dyes; Textiles; Leather Tanning and Processing; Paper and Pulp Processing; Wood and Lumber; Laboratory Supplies; Plastics and Rubber;	Inhalation; Dermal; Ingestion;	Bladder;	Personal Care; Processing; Art/ Office Supplies; Manufacturing	Personal Care; Art/ Office Supplies; Environmental Releases	
C.i. Direct Brown 95	Inks and Dyes; Textiles; Leather Tanning and Processing; Paper and Pulp Processing; Wood and Lumber; Laboratory Supplies; Plastics and Rubber;	Inhalation; Dermal; Ingestion;	Bladder;	Personal Care; Processing; Art/ Office Supplies; Manufacturing	Personal Care; Art/ Office Supplies; Environmental Releases	
Cadmium Chloride	Herbicides and Pesticides; Textiles; Glass and Ceramics; Metals and Electronics; Plastics and Rubber;	Ingestion; Inhalation;;	Lungs; Prostate; Kidney;	Farming/ Agriculture; Processing	Art/ Office Supplies; Drinking Water; Food/ Food Additives	
Calcium Arsenate	Herbicides and Pesticides; Disinfectants;	Inhalation; Dermal; Ingestion;	Lung; Skin; Lymphatic;	Farming/ Agriculture; Manufacturing; Processing	Drinking Water; Environmental Releases	
Calcium Monochromate	Metals and Electronics; Paints, Finishes, Adhesives;	Inhalation; Ingestion; Dermal;	Lung; Sinus/Nasal;	Manufacturing; Processing; Home Maintenance	Electronics/ Small Appliances; Drinking Water; Home Maintenance; Environmental Releases	
Chloromethyl Methyl Ether	Food and Food Production; Laboratory Supplies; Paints, Finishes, Adhesives; Metals and Electronics;	Inhalation; Dermal;	Lung;	Manufacturing; Processing	Environmental Releases	

Continued, next page...

Table B1: Exposure pathways and cancer types for known carcinogens

Chemical	Major Prod. Categories	Exp. route	Associated cancers	Occupational exposure	General exposure	NHANES data
Chromium Lead Oxide	Metals and Electronics; Paints, Finishes, Adhesives; Inks and Dyes; Plastics and Rubber; Paper and Pulp Processing; Glass and Ceramics; Textiles;	Inhalation; Ingestion;	Lung; Sinus/Nasal;	Home Maintenance; Manufacturing; Processing; Construction/ Building	Home Maintenance; Food/ Food Additives	
Chromium Trioxide	Wood and Lumber; Metals and Electronics; Paints, Finishes, Adhesives; Pharmaceuticals;	Inhalation; Dermal; Ingestion;	Lung; Sinus/Nasal;	Processing; Construction/ Building; Home Maintenance; Vehicle; Manufacturing	Home Maintenance	
Chrysotile	Concrete and Construction; Food and Food Production; Mining and Smelting; Metals and Electronics; Paints, Finishes, Adhesives; Paper and Pulp Processing; Plastics and Rubber; Laboratory Supplies;	Inhalation;	Lung;	Vehicle; Construction/ Building; Mining; Processing; Home Maintenance; Manufacturing; Processing	Vehicle; Personal Care; Electronics/ Small Appliances; Home Maintenance; Art/ Office Supplies; Drinking Water; Medical/ Dental	
Cristobalite	Pharmaceuticals; Food and Food Production; Metals and Electronics; Mining and Smelting; Glass and Ceramics; Plastics and Rubber; Paints, Finishes, Adhesives; Cosmetics; Wood and Lumber; Paper and Pulp Processing; Disinfectants; Herbicides and Pesticides; Concrete and Construction; Petroleum, Oil, Biofuels;	Inhalation;	Lung;	Mining; Construction/ Building; Farming/ Agriculture; Processing; Manufacturing; Medical/ Dental	Home Maintenance; Environmental Releases; Personal Care; Food/ Food Additives; Art/ Office Supplies; Smoking and Tobacco; Medical/ Dental	
Cyclophosphamide	Pharmaceuticals;	Ingestion; Injection; Dermal; Inhalation;	Bladder; Leukemia;	Manufacturing; Processing	Medical/ Dental	
Cyclosporin A	Pharmaceuticals;	Ingestion; Injection; Dermal; Inhalation;	Lymphoma; Skin;	Manufacturing; Processing	Medical/ Dental; Pet Care	

Continued, next page...

Table B1: Exposure pathways and cancer types for known carcinogens

Chemical	Major Prod. Categories	Exp. route	Associated cancers	Occupational exposure	General exposure	NHANES data
Estrone	Pharmaceuticals; Cosmetics; Food and Food Production;	Ingestion; Dermal;	Uterus; Breast;	NA	Personal Care; Medical/ Dental; Food/ Food Additives	
Lead Arsenate	Metals and Electronics; Herbicides and Pesticides;	Inhalation; Ingestion; Dermal;	Lung; Skin; Liver; Kidney; Bladder;	Farming/ Agriculture; Manufacturing; Pet Care	Drinking Water; Environmental Releases; Processing; Smoking and Tobacco	
Mestranol	Pharmaceuticals;	;	;	NA	Medical/ Dental	
Mineral Oil - Includes Paraffin Oil	Petroleum, Oil, Biofuels; Herbicides and Pesticides; Plastics and Rubber; Metals and Electronics; Inks and Dyes; Cosmetics; Disinfectants; Pharmaceuticals; Textiles; Paints, Finishes, Adhesives; Solvents; Food and Food Production;	Dermal; Ingestion;	Skin; Testicular/Scrotal; Gastrointestinal; Rectal; Bladder;	Farming/ Agriculture; Art/ Office Supplies; Manufacturing; Processing	Cleaning Products; Art/ Office Supplies; Personal Care; Medical/ Dental; Food/ Food Additives	
Nickel (II) Oxide	Metals and Electronics; Glass and Ceramics; Paints, Finishes, Adhesives; Mining and Smelting; Laboratory Supplies;	Inhalation;	Lung; Sinus/ Nasal;	Vehicle; Art/ Office Supplies; Manufacturing; Processing; Mining	Art/ Office Supplies	
Nickel Hydroxide (Ni(OH) ₂)	Metals and Electronics; Mining and Smelting; Plastics and Rubber;	Inhalation; Ingestion; Dermal;	Lung; Sinus/ Nasal;	Electronics/ Small Appliances; Processing	NA	
Nickel Subsulfide	Metals and Electronics; Mining and Smelting; Petroleum, Oil, Biofuels;	Inhalation; Ingestion; Dermal;	Lung; Sinus/ Nasal;	Electronics/ Small Appliances; Processing	NA	
Nickel Sulfate	Metals and Electronics; Mining and Smelting; Inks and Dyes; Glass and Ceramics; Textiles;	Inhalation; Ingestion; Dermal;	Lung; Sinus/ Nasal;	Electronics/ Small Appliances; Manufacturing; Processing	Cleaning Products; Drinking Water	
Nickel Sulfide	Petroleum, Oil, Biofuels; Metals and Electronics;	Inhalation;	Lung; Sinus/ Nasal;	Processing	NA	
Nickel(II) Acetate	Textiles;	Inhalation;	Lung; Sinus/ Nasal;	Manufacturing	NA	

Continued, next page...

Table B1: Exposure pathways and cancer types for known carcinogens

Chemical	Major Prod. Categories	Exp. route	Associated cancers	Occupational exposure	General exposure	NHANES data
Potassium Arsenite Anhydrous	Glass and Ceramics; Metals and Electronics; Herbicides and Pesticides; Wood and Lumber; Laboratory Supplies; Pharmaceuticals;	Inhalation; Ingestion; Dermal;	Skin; Lung; Stomach; Liver; Bladder; Kidney; Lymphatic; Hematopoietic;	Pet Care; Manufacturing; Processing; Farming/ Agriculture	Medical/ Dental	
Potassium Chromate(Vi)	Leather Tanning and Processing; Paints, Finishes, Adhesives; Textiles; Metals and Electronics; Laboratory Supplies;	Inhalation; Ingestion; Dermal;	Lung; Sinus/Nasal;	Manufacturing; Processing	Art/ Office Supplies	
Potassium Dichromate	Pharmaceuticals; Textiles; Leather Tanning and Processing; Paints, Finishes, Adhesives; Wood and Lumber; Metals and Electronics; Disinfectants; Food and Food Production;	Ingestion; Inhalation; Dermal;	Lung; Sinus/Nasal;	Processing; Manufacturing	Personal Care; Art/ Office Supplies; Cleaning Products; Home Maintenance; Food/ Food Additives	
Premarin	Pharmaceuticals;	Ingestion;	Uterus; Breast;	NA	Medical/ Dental	
Riebeckite	Concrete and Construction; Plastics and Rubber; Mining; Paints, Finishes, Adhesives;	Inhalation; Dermal; Ingestion;	Lung;	Construction/ Building; Home Maintenance; Manufacturing; Processing; Mining	Food/ Food Additives; Smoking and Tobacco; Home Maintenance	
Sodium Arsenate	Herbicides and Pesticides; Glass and Ceramics; Textiles; Cosmetics; Pharmaceuticals;	Inhalation; Dermal; Ingestion;	Bladder; Lung; Liver; Kidney; Skin;	Processing; Manufacturing; Pet Care	Personal Care	
Sodium Arsenite	Herbicides and Pesticides; Leather Tanning and Processing; Disinfectants; Pharmaceuticals; Inks and Dyes; Laboratory Supplies; Wood and Lumber; Textiles;	Ingestion; Inhalation; Dermal;	Bladder; Lung; Liver; Kidney; Skin;	Pet Care; Processing; Manufacturing; Farming/ Agriculture; Art/ Office Supplies	Art/ Office Supplies	

Continued, next page...

Table B1: Exposure pathways and cancer types for known carcinogens

Chemical	Major Prod. Categories	Exp. route	Associated cancers	Occupational exposure	General exposure	NHANES data
Sodium Chromate	Leather Tanning and Processing; Textiles; Inks and Dyes; Wood and Lumber; Herbicides and Pesticides; Laboratory Supplies; Petroleum, Oil, Biofuels; Paints, Finishes, Adhesives;	Dermal; Inhalation; Ingestion;	Lung; Sinus/Nasal;	Manufacturing; Processing; Medical/ Dental	Electronics/ Small Appliances; Home Maintenance; Environmental Releases; Food/ Food Additives	
Sodium Dichromate	Metals and Electronics; Wood and Lumber; Inks and Dyes; Waste Incineration and Treatment; Disinfectants; Herbicides and Pesticides; Glass and Ceramics; Food and Food Production; Textiles; Petroleum, Oil, Biofuels;	Dermal; Inhalation; Ingestion;	Lung; Sinus/Nasal;	Processing; Manufacturing; Home Maintenance	Cleaning Products; Home Maintenance; Food/ Food Additives; Drinking Water	
Strontium Chromate	Metals and Electronics; Plastics and Rubber; Leather Tanning and Processing; Paints, Finishes, Adhesives;	Inhalation; Dermal;	Lung; Sinus/Nasal;	Manufacturing; Processing; Home Maintenance	Home Maintenance	
Sulfuric Acid	Food and Food Production; Metals and Electronics; Paper and Pulp Processing; Leather Tanning and Processing; Textiles; Herbicides and Pesticides; Mining and Smelting; Glass and Ceramics; Inks and Dyes; Cosmetics; Pharmaceuticals; Plastics and Rubber; Waste Incineration and Treatment; Wood and Lumber; Paints, Finishes, Adhesives; Petroleum, Oil, Biofuels;	Inhalation; Ingestion; Dermal;	Larynx; Lung; Sinus/Nasal;	Art/ Office Supplies; Manufacturing; Processing; Mining; Farming/ Agriculture; Cleaning Products	Art/ Office Supplies; Electronics/ Small Appliances; Home Maintenance; Landscape/ Yard; Personal Care; Pet Care; Environmental Releases; Cleaning Products; Sports Equipment; Vehicle; Food/ Food Additives; Drinking Water	

Continued, next page...

Table B1: Exposure pathways and cancer types for known carcinogens

Chemical	Major Prod. Categories	Exp. route	Associated cancers	Occupational exposure	General exposure	NHANES data
Thorium(Iv) Oxide	Pharmaceuticals; Glass and Ceramics; Nuclear; Mining; Metals and Electronics; Laboratory Supplies;	Inhalation; Injection; Ingestion; Dermal;	Liver; Leukemia; Bone; Bile Duct; Pancreatic;	Manufacturing; Processing; Medical/ Dental; Research; Pet Care; Mining	Medical/ Dental; Environmental Releases	
Tremolite Asbestos	Glass and Ceramics; Paints, Finishes, Adhesives; Cosmetics; Mining and Smelting; Metals and Electronics; Waste Incineration and Treatment;	Inhalation;	Lung;	Construction/ Building; Manufacturing; Processing; Mining	Construction/ Building; Personal Care; Environmental Releases; Drinking Water	
Tridymite	Pharmaceuticals; Food and Food Production; Metals and Electronics; Mining and Smelting; Glass and Ceramics; Plastics and Rubber; Paints, Finishes, Adhesives; Cosmetics; Wood and Lumber; Paper and Pulp Processing; Disinfectants; Herbicides and Pesticides; Concrete and Construction; Petroleum, Oil, Biofuels;	Inhalation;	Lung;	Mining; Construction/ Building; Farming/ Agriculture; Processing; Manufacturing; Medical/ Dental	Home Maintenance; Environmental Releases; Personal Care; Food/ Food Additives; Art/ Office Supplies; Smoking and Tobacco; Medical/ Dental	
Zinc Chromate	Paints, Finishes, Adhesives; Inks and Dyes; Leather Tanning and Processing; Food and Food Production; Metals and Electronics;	Inhalation; Ingestion; Dermal;	Lung; Sinus/Nasal;	Manufacturing; Processing	Food/ Food Additives	
Bis(Chloromethyl) Ether	Textiles; Plastics and Rubber; Laboratory Supplies;	;	Lung;	Manufacturing; Processing	NA	

Notes: Compiled exposure source and type, product category, and associate cancer type information for the complete list of 112 known carcinogens. See Section 2.4 and Section 3.4 for description of exposure pathway information collection and product categorization methods.

C Information on data construction and exposure literature review

Additional details of NHANES data processing and analysis

This analysis primarily relied on a merged and harmonized version of the continuous NHANES data; we analyzed examination-level biomonitoring samples drawn cross-sections of individuals between 2003-2004 and 2017-2018 (Nguyen et al., 2023). We examined blood, serum, and urine biomarkers only for chemicals that were on the ubiquitous carcinogen list. We followed best practices in the use of NHANES: examination or subsample survey weights were used to account for NHANES’ sampling design (Nguyen et al., 2023; CDC, 2020), efforts were taken to ensure metabolites of carcinogens were correctly assigned to the parent chemical (Stanfield et al., 2024), and the analysis followed the CDC’s convention of not reporting geometric mean estimates for stratification subgroups with 40% of test results below the reported limit of detection (CDC, 2024). For subgroups with sufficient detected observations, we imputed a value of $\frac{LOD}{\sqrt{2}}$ for samples below the detection limit when reporting geometric mean values. This is a common approximation method of censored values below the detection threshold for screening work. For carcinogens studied in multiple NHANES cycles, statistics were calculated based on only the most recent cycle of data. The most recent cycle varied by chemical: the earliest cycle considered in this analysis was from 2003-2004 and the latest was from 2017-2018.

There were some instances in which multiple biomarker media or measurement variations were available for a given chemical or metabolite. We made the following set of selected choices. We relied on lipid-adjusted statistics for the following chemicals: heptachlor epoxide, Dieldrin, and p,p’-DDE. We relied on statistics from blood rather than urine concentrations for the following metals that were tested in both media: lead, cadmium, cobalt. We calculated urine concentration statistics for total and speciated arsenic chemicals, which tend to be less detectable. Biomonitoring results listed as “Arsenic” refer to total arsenic throughout the analysis. We calculated statistics based on total PFOS and PFOA concentrations rather than separately analyzing the branched and linear isomers. Total chromium levels in blood and urine are available in the NHANES microdata; because only hexavalent chromium is on our carcinogen list and cannot be differentiated from trivalent chromium via existing biomonitoring, we elected not to present results for total chromium.

Demographic variables used in the analysis of individual-level biomonitoring data include race/ethnicity, age, sex, and household income. We relied on the “RIDRETH1” race and ethnicity recode variable throughout our analysis. This decision resulted in uniform categories across the different NHANES cycle, but limited our racial/ethnic groups to Hispanic, non-Hispanic Black, non-Hispanic White, and Other.¹ A decomposition of the Other category that differentiates non-Hispanic Asian respondents was available from the 2011-2012 cycle onwards using the “RIDRETH3” variable, but to maintain within-analysis consistency across the dozens of chemicals assessed, we solely relied on the more limited racial/ethnic sub-groupings. As our measurement for household income, we relied on the “INDFMPIR” variable, which is an index for the ratio of family income to poverty. Using this variable, we categorized surveyed individuals as below 2x the poverty line or above 2x the poverty line. The small set of individuals for which this variable was missing were dropped from our analysis. We coded individual’s age and sex based on the variables “RIDAGEYR” and “RIAGENDR”. Remaining statistical details of GM and P95 concentration ratios are in the main text.

¹Prior to 2007-2008, NHANES oversampled for Mexican Americans only (versus all Hispanic people), so the small number of estimates based on data from pre-2007 are representative of Mexican Americans rather than Hispanics.

For a subset of biomonitored chemicals – namely, organochloride pesticides, dioxins/furans, PCBs, and brominated flame retardants – the most recent NHANES samples available are pooled. In those cases, we directly acquired and relied on the original, unharmonized NHANES microdata from the 2011-2012 and 2015-2016 cycles (CDC, 2020). Due to the pooled nature of these biomonitoring data and their sampling design, we could only assess arithmetic means and create differentiated statistics by race/ethnicity, sex, and age (CDC, 2021a). As before, we followed the CDC’s convention of not reporting mean estimates for stratification subgroups with 40% of test results below the reported limit of detection. Because all pooled samples were from cycles in 2011-2012 or later, we relied on the “RIDRETH3” race/ethnicity indicator. Thus, pooled biomonitoring results in the paper were presented with separate non-Hispanic Asian and Other groupings. We used these estimates from the pooled data to calculate arithmetic mean ratios (*AMRs*) analogous to the geometric mean ratio described in the main text, again using the entire national population as the reference group.

Additional details of DW sample processing, national representativeness

Despite rigorous quality assurance and quality control assessments that were conducted prior to publishing the SYR and UCMR drinking water sampling data, their analytical use comes with complexities (US Environmental Protection Agency, 2016a, 2022a). These are largely related to missing detection limits, inconsistencies in state-reported concentration units for sampling and detection limits, and heterogeneity in the types of samples drawn by community water systems. Our processing of SYR data followed much of Ravalli et al. (2022), with details described as follows:

1. We first removed all unfinished and raw drinking water samples.
2. For samples flagged as non-detects that are typically reported in $\mu\text{g/L}$, all reported limits of detection (LOD) were standardized to $\mu\text{g/L}$. For samples flagged as non-detects that are typically reported in mg/L , all reported limits of detection (LOD) were standardized to mg/L . The exception is asbestos which is measured in fibers (mf/L).
3. The modal LOD was calculated for each chemical sampled. Any non-detect sample with a reported LOD four orders of magnitude or smaller (10^{-4}) than the modal LOD were imputed with the modal LOD due to likely reporting errors from the state or CWS. Modal LODs for each SYR chemical are reported below in Table C1.
4. Samples without a reported LOD were imputed with the modal LOD for the chemical.
5. Using the reported or imputed LOD as described above, all non-detect samples were assigned sample concentration values equal to $\frac{\text{LOD}}{\sqrt{2}}$.

Our processing of UCMR data was similar in nature, though less complex. By design, UCMR2 through UCMR4 samples were drawn from finished water only. An EPA-established, chemical-specific Minimum Reporting Level (MRL) was provided for all non-detected samples; using this MRL, we imputed sample concentration values equal to $\frac{\text{MRL}}{\sqrt{2}}$ for all non-detects.²

²UCMR data can be considered a census of large systems, but they do not fully represent water quality at small systems. While the vast majority of the US population is served by large systems, we acknowledge that our statistics may somewhat under-represent smaller systems.

Next, we combined samples across monitoring regimes. Only the most recent round of monitoring samples for a given chemical was included in our final data: if a chemical was monitored in SYR4, we dropped any samples of it that were taken during SYR3. Similarly, if a chemical was monitored in UCMR4, we dropped any samples of it that were taken during UCMR2 or UCMR3.

We then followed Ravalli et al. (2022) and restricted our main analyses to only chemicals with more than 10% of samples above the LOD/MRL. In the absence of this restriction, any statistics generated could lend a sense of false precision. By requiring a 10% sample detection threshold, we generated statistics built from observed signals in the data rather than a reliance on distributional assumptions below censoring thresholds.³ Fifteen of the 70 original chemicals met the detection requirement, and we proceeded using this subsample.

For each CWS and retained chemical, we calculated an average carcinogen concentration based on all of the system’s collected samples from 2006-2020. We used customer demographics information for each CWS as calculated by Austin et al. (2024) using the EPA-ORD water system boundaries described in the paper. These included a count of the population served by the CWS, the fraction of constituents of each race/ethnicity, the fraction with household income below the poverty line and the fraction with household income below twice the poverty line, all as of 2022. With this information, we calculated population-weighted average carcinogen concentrations in drinking water (DW_{cs}^*) for each demographic subgroup:

$$DW_{cs}^* = \sum_J \frac{DW_{jc} Pop_{js}}{Pop_s}$$

where J indexes the complete set of CWSs for which a given carcinogen has sampling data, DW_{jc} is the average concentration level of carcinogen c in the drinking water of CWS j , Pop_{js} is the population count of demographic subgroup s in CWS j , and Pop_s is the national population count of subgroup s across CWSs for which a carcinogen has data.⁴

With these average drinking water concentrations for each subgroup and carcinogen, we calculated carcinogen-specific relative concentration ratios for each racial/ethnic or household income subgroup as in Equation 1, again with the national average concentration for the carcinogen acting as a reference value. These relative concentration ratios served as our measure of differences in exposure via drinking water.

Table C1 below provides summary measures to help illustrate the processing description above. Column (2) describes the percentage of a chemical’s samples with a detectable concentration. Column (3) shows the number of CWSs that reported any sample of the chemical. Column (4) shows the 2022 population count for the set of CWSs that reported sampling the chemical. Column (5) describes the percentage of CWSs for which at least 10% of samples had detectable concentrations. Column (6) shares the chemical’s modal reported limit of detection. Column (7) shares the chemical’s unit of measurement. Column (8) shows the data source from which samples were drawn for each chemical.

³Our selected 10% threshold is not fully immune from this concern; as such we caveated our results in the limitations section of the main paper. Future research could examine correlations between sociodemographic characteristics of a community and the LODs/RLs used by the CWS serving them, with an eye to understanding how such correlations could affect subgroup-specific drinking water exposure metrics

⁴A reviewer pointed out that much of the meaningful historic variation in disinfection byproduct (DBP) exposure occurred within a given distribution system, and thus certain subpopulations may be differentially affected even though the system-wide average of DBP concentrations does not deviate from national norms. These within-system deviations have dampened over time, and therefore our reliance on the most recent possible vintage of SYR data may somewhat mitigate this concern. But this fact nonetheless should be kept in mind while interpreting results for DBPs.

Table C1: Drinking water data: underlying data info and representativeness

(1)	(2)	(3)	(4)	(5)	(6)	(7)	(8)
Chemical	% Detect	PWS count	Pop. Count	% PWS Det.	Modal DL	DL Unit	Source
HAA5*	100.00	4,751	254,581,887	100.00		ug/L	UCMR4
HAA6BR*	100.00	4,751	254,581,887	100.00		ug/L	UCMR4
HAA9*	100.00	4,751	254,581,887	100.00		ug/L	UCMR4
DICHLOROACETIC ACID*	78.09	28,112	210,841,018	68.42	1	ug/L	SYR4
DICHLOROMONOBROMOMETHANE*	75.75	31,728	223,654,395	72.54	1	ug/L	SYR4
CHROMIUM-6*	75.61	4,659	253,306,908	88.35	0.03	ug/L	UCMR3
TRICHLOROMETHANE*	75.04	31,763	223,711,184	72.46	1	ug/L	SYR4
TRICHLOROACETIC ACID*	70.47	28,105	210,161,481	58.68	1	ug/L	SYR4
ARSENIC*	44.97	27,858	216,001,168	38.26	0.002	mg/L	SYR3
DIBROMOACETIC ACID*	41.21	28,106	210,775,662	53.49	1	ug/L	SYR4
TRIBROMOMETHANE*	33.13	31,703	223,399,206	46.09	1	ug/L	SYR4
BROMATE*	22.97	256	32,153,441	35.94	5	ug/L	SYR4
MICROCYSTIN-LR*	20.00	5	88,293	20.00	0.02	ug/L	UCMR4
1,4-DIOXANE*	11.71	4,655	253,224,006	20.21	0.07	ug/L	UCMR3
NDMA*	10.94	1,042	142,332,591	24.28	0.002	ug/L	UCMR2
TRICHLOROETHYLENE	6.41	28,837	213,565,611	1.15	0.5	ug/L	SYR3
TETRACHLOROETHYLENE	6.09	28,838	213,585,357	1.56	0.5	ug/L	SYR3
BIS(2-ETHYLHEXYL) PHTHALATE	4.19	19,146	167,952,482	7.72	0.6	ug/L	SYR3
1,2-DIBROMO-3-CHLOROPROPANE	3.26	20,328	173,545,539	0.48	0.02	ug/L	SYR3
ASBESTOS	2.95	3,132	69,737,620	2.91	0.2	mf/L	SYR3
1,1-DICHLOROETHANE	2.33	4,656	253,242,431	3.57	0.03	ug/L	UCMR3
1,1-DICHLOROETHYLENE	1.57	28,829	213,328,575	0.36	0.5	ug/L	SYR3
CADMIUM	1.51	25,781	201,881,945	3.29	0.001	mg/L	SYR3
COBALT	1.35	4,660	253,354,223	3.05	1	ug/L	UCMR3
ETHYLBENZENE	1.09	28,844	213,292,583	2.90	0.5	ug/L	SYR3
PFOA	1.07	4,660	253,354,223	2.02	0.02	ug/L	UCMR3
BERYLLIUM, TOTAL	0.99	25,563	201,078,820	1.89	0.001	mg/L	SYR3
4-ANDROSTENE-3,17-DIONE	0.83	1,082	154,022,064	5.36	0.0003	ug/L	UCMR3
PFOS	0.77	4,660	253,354,223	1.46	0.04	ug/L	UCMR3
CARBON TETRACHLORIDE	0.76	28,833	213,405,463	0.80	0.5	ug/L	SYR3
1,2,3-TRICHLOROPROPANE	0.70	4,656	253,242,431	0.88	0.03	ug/L	UCMR3
TESTOSTERONE	0.56	1,082	154,022,064	3.33	0.0001	ug/L	UCMR3
METHYLENE CHLORIDE	0.54	28,826	213,218,763	1.27	0.5	ug/L	SYR3
1,1,1-TRICHLOROETHANE	0.46	28,828	213,238,681	0.40	0.5	ug/L	SYR3
1,2-DICHLOROETHANE	0.43	28,828	213,275,283	0.36	0.5	ug/L	SYR3
HEPTACHLOR EPOXIDE	0.37	21,276	174,134,820	0.24	0.02	ug/L	SYR3
1,2-DIBROMOMETHANE	0.35	21,043	173,763,025	0.30	0.01	ug/L	SYR3
O-TOLUIDINE	0.32	4,823	255,422,296	0.95	0.007	ug/L	UCMR4
1,2-DICHLOROPROPANE	0.31	28,827	213,230,545	0.29	0.5	ug/L	SYR3
QUINOLINE	0.31	4,823	255,422,296	1.16	0.02	ug/L	UCMR4
BENZENE	0.30	28,829	213,305,081	0.46	0.5	ug/L	SYR3
NDEA	0.27	1,042	142,332,591	0.58	0.005	ug/L	UCMR2
NPYR	0.25	1,042	142,332,591	1.44	0.002	ug/L	UCMR2
2,4-D	0.23	20,227	176,364,002	0.37	0.1	ug/L	SYR3
P-DICHLOROBENZENE	0.20	28,829	212,919,949	0.54	0.5	ug/L	SYR3
VINYL CHLORIDE	0.20	28,484	212,298,096	0.32	0.5	ug/L	SYR3
PAHs	0.12	19,647	170,136,287	0.19	0.1	ug/L	SYR3
CHLORDANE	0.11	20,103	169,740,088	0.10	0.2	ug/L	SYR3
STYRENE	0.11	28,823	213,223,684	0.39	0.5	ug/L	SYR3
ALACHOR	0.08	23,244	187,744,943	0.12	0.2	ug/L	SYR3
ALPHA-HEXACHLOROCYCLOHEXANE	0.07	4,821	255,433,030	0.37	0.01	ug/L	UCMR4
BHC-GAMMA	0.07	21,517	176,036,138	0.07	0.02	ug/L	SYR3
HEPTACHLOR	0.06	21,309	174,230,503	0.08	0.04	ug/L	SYR3
NDBA	0.05	1,042	142,332,591	0.00	0.004	ug/L	UCMR2
TOXAPHENE	0.05	20,665	172,289,075	0.07	1	ug/L	SYR3
1,1,2-TRICHLOROETHANE	0.04	28,826	213,228,594	0.13	0.5	ug/L	SYR3

Continued, next page...

Table C1: Drinking water data: underlying data info and representativeness

(1)	(2)	(3)	(4)	(5)	(6)	(7)	(8)
Chemical	% Detect	PWS count	Pop. Count	% PWS Det.	Modal DL	DL Unit	Source
17-ALPHA-ETHYNYLESTRADIOL	0.04	1,082	154,022,064	0.18	0.0009	ug/L	UCMR3
2,3,7,8-TCDD	0.04	1,726	51,384,238	0.17	0.000005	ug/L	SYR3
DECACHLOROBIPHENYL	0.04	11,602	115,238,214	0.12	0.1	ug/L	SYR3
17-BETA-ESTRADIOL	0.03	1,082	154,022,064	0.00	0.0004	ug/L	UCMR3
GLYPHOSATE	0.03	9,483	109,125,633	0.12	6	ug/L	SYR3
HEXACHLOROBENZENE	0.03	21,251	173,474,370	0.08	0.1	ug/L	SYR3
NMEA	0.02	1,042	142,332,591	0.29	0.003	ug/L	UCMR2
ETHOPROP	0.01	4,821	255,433,030	0.06	0.03	ug/L	UCMR4
TRIBUFOS	0.01	4,820	255,408,264	0.04	0.07	ug/L	UCMR4
1,3-BUTADIENE	0.00	4,656	253,242,431	0.00	0.1	ug/L	UCMR3
ACETOCHLOR	0.00	1,042	142,332,591	0.00	2	ug/L	UCMR2
ESTRONE	0.00	1,082	154,022,064	0.00	0.002	ug/L	UCMR3
NDPA	0.00	1,042	142,332,591	0.00	0.007	ug/L	UCMR2
TNT	0.00	3,794	216,424,871	0.00	0.8	ug/L	UCMR2

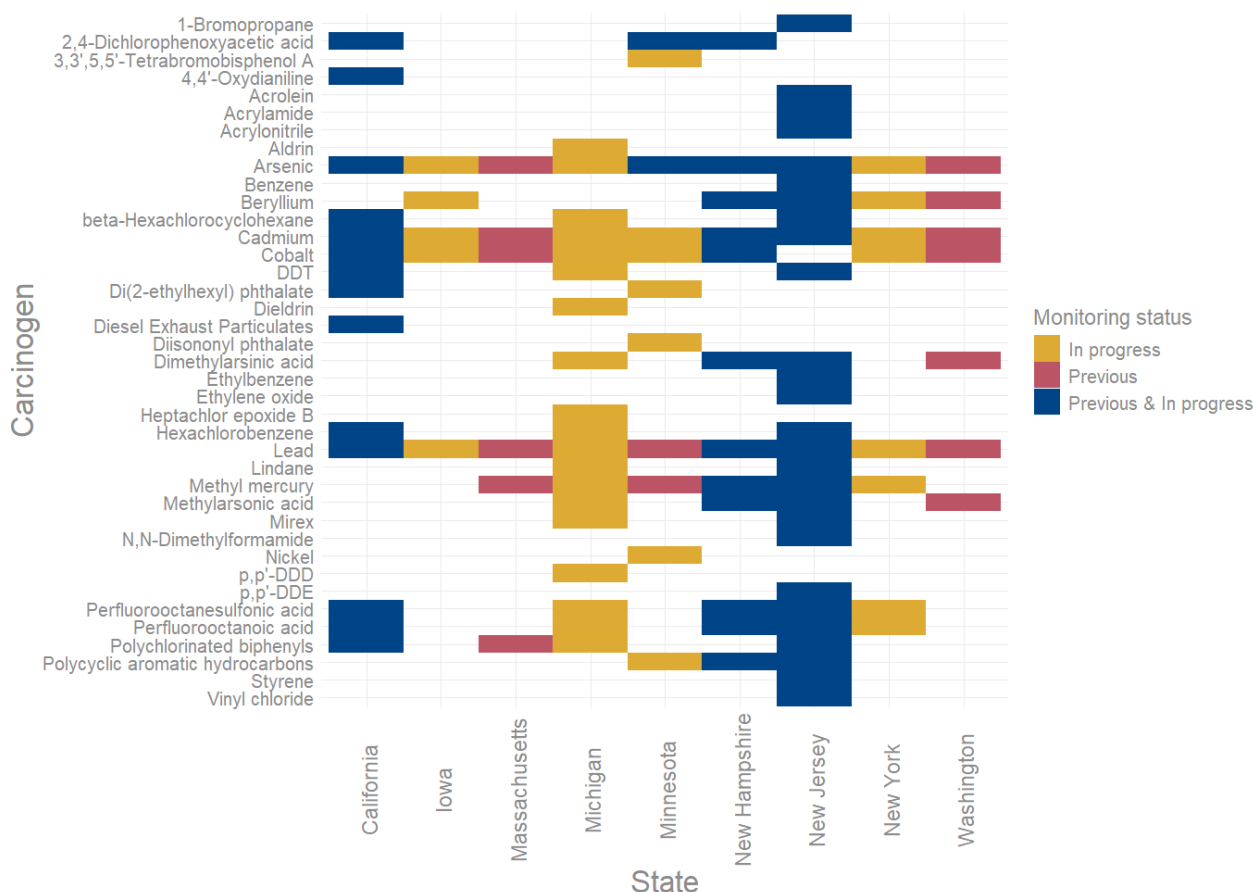
Return to: Section 2.2.3. Section 3.2.2.

D Summary of carcinogen biomonitoring by states⁵

Several US states possess their own biomonitoring programs. Many of these programs were started and have continued operations using money from competitive CDC state biomonitoring grants. Together, these states form the core of the National Biomonitoring Network, a collaborative effort consisting of federal, state, and local laboratories that aim to ensure quality laboratory practices, helping produce comparable biomonitoring data across levels of government (Latshaw et al. (2017), Nassif et al. (2021)).

In scoping available biomonitoring data for the paper's main analysis, we contacted nine states. The following appendix summarizes information collected in early 2024 about several states' individual biomonitoring efforts, including the extent to which carcinogens are or were being monitored, and any available summary exposure statistics stratified by demographic subgroups. Figure D1 below summarizes findings on surveillance data availability for carcinogens across the states.

Figure D1: Carcinogen surveillance data availability by state



Across the states that made information available, 39 carcinogens in total were biomonitoring. All of these carcinogens except 4,4'-Oxydianiline, 1-Bromopropane, Aldrin, p,p'-DDD, and TBBPA were monitored in

⁵Several individuals involved in state biomonitoring programs were incredibly generous with their time. We are very grateful to the following individuals for their helpful conversations and, when possible, the provision of data, as this project got off the ground: Nerissa Wu, Kathleen Attfield, Tina Fan, Chang Yu, Melissa Josefiak, Madhumita Chatterjee, Jessica Nelson, Sheila Amenumey, Meg Blanchet, Rachel Long, Rebecca Hunt, Gonza Namulanda, and Robin Dodson.

NHANES at some point from 1988-2018. The modest additional carcinogen coverage from state biomonitoring is logical. States rely on the same laboratory methods used nationally by NHANES and are unlikely to have time and resources dedicated to expanding such capabilities. Additionally, a key justification for state biomonitoring programs is the need to understand how local and regional exposures compare to the national distribution. This requires surveillance of chemicals for which the national exposure distribution is known.

Because of the limited value-add that state level biomonitoring results can contribute relative to NHANES, our analysis relied on the national data alone. Nevertheless, a summary of findings from the aborted scoping exercise follows:

California

California was one of the earliest state biomonitoring programs to receive a CDC grant (2009), and today is likely the most advanced in terms of capabilities. By law, California has a formal biomonitoring selection process in place to filter down the universe of chemicals into a single designation list of several hundred. From this designation list, a Scientific Guidance Panel recommends prioritization of a subset based on degree of potential exposure, likelihood of a chemical being a carcinogen/toxicant, the limits of laboratory detection, and more. From this priority list, California DPH selects a final group of chemicals for biomonitoring based on laboratory considerations, feasibility, public resources, and special relevance of a chemical to a particular sub-population being studied. In practice, the state typically has resources to monitor less than 100 chemicals.

California has performed a wide variety of biomonitoring projects over the years; some surveilled a statewide population sample while others focused on particular regions, certain demographic or occupational groups, or populations affected by health outcomes of concern. Ensuring biomonitoring for a diverse set of populations is a primary goal of CA DPH. In sum, California monitored 113 chemicals over this period. 12 of these chemicals are on the carcinogen list.

Of these studies, the California Regional Exposure (CARE) Study and Biomonitoring Exposures Study (BEST) provide the best coverage of chemicals and populations more representative of regions or the entire state. The expanded wave of BEST is now complete, but CA DPH does not have weighted and demographically-stratified summary statistics on hand at this time. CA DPH was able to share a new public report including demographically-stratified carcinogen summary statistics for CARE-LA (LA County) and CARE-2 (Riverside, San Bernardino, Imperial, Mono, and Inyo Counties) which were conducted from 2018-2020.

Washington

In 2010-2011, Washington state conducted a state-level biomonitoring program. Urine samples were obtained from a representative sample of households across the state and paired with a household questionnaire used to determine demographics. The Washington Tracking Network shares the results of this biomonitoring, stratified by demographic variables of interest, on a public dashboard.

The dashboard data were procured, cleaned and standardized. Of the 25 chemical chemicals surveilled in WA, only 7 are on the carcinogen list. Five are metals: arsenic, beryllium, cadmium, cobalt, and lead. The other two are carcinogenic metabolites of arsenic: DMA and MMA.

New Jersey

New Jersey was awarded CDC grants in 2014 and 2019 for development of its biomonitoring program, NJHANES. During the 2014-2019 grant cycle, statewide surveillance was performed on metals in blood and urine, on PFAS in serum, and on PCBs in serum. Demographic information was also collected during a corresponding household survey.

NJ DOH was unable to immediately provide machine-readable chemical-by-chemical summary statistics stratified by demographics or access to the underlying microdata. They did refer to published and working papers that include results stratified by sex, though these do not include measures varying by race/ethnicity or household income (Du et al. (2020); Yu et al. (2020)).

In addition to these results, NJ DOH kindly provided a complete list of NJHANES analytes for the 2019-2024 project cycle. This list included 122 analytes, including around 20 variations of heavy metals, several polycyclic aromatic hydrocarbons (PAH) metabolites, several VOC metabolites, 12 PFAS and dozens of polychlorinated biphenyls (PCBs) and polybrominated diphenyl ethers (PBDEs). Of these 122 analytes, 27 are carcinogens. Five are metals, two are metabolites of arsenic, and the remainder can be classified as PFAS (PFOA and PFOS), PAHs, PCBs, metabolites of VOCs (e.g. 1-Bromopropane, Benzene, Styrene, etc.), and pesticides or their byproducts (DDT, DDE, Mirex, and hexachlorocyclohexanes).

New Hampshire

New Hampshire was awarded CDC grants in 2014 and 2019 for development of their biomonitoring program, NH TrACE. During the 2014-2019 grant cycle, statewide surveillance was performed on roughly 20 heavy metals, roughly 15 PFAS, and roughly 10 pesticides. Demographic information was also collected during a corresponding household survey. Linked water testing was also conducted for private wells and publicly-sourced water, providing information on household exposure to metals, radiologicals, PFAS, and a large variety of VOCs and pesticides via the drinking water pathway. In sum, these data provide substantial potential for stratified exposure summary statistics by demographics of interest.

NH produced a summary report of findings from this initial cycle in 2019. Unfortunately, this report does not contain any statistics of interest and the underlying raw microdata are not publicly available at the present time. While there is very limited racial and ethnic diversity in NH's population sample, there may be interesting heterogeneity along the household income dimension.

NH has continued NH TrACE during the 2019-2024 award cycle, surveilling 35 of the chemicals studied previously as well as metabolites of polycyclic aromatic hydrocarbons. A summary report on findings is expected late in 2024 and may include demographically-stratified summary statistics. In addition, smaller-scale biomonitoring studies are targeting residents who rely on private well water, Medicaid recipients, and upstate residents located near an EPA Superfund site; these studies will test for a similar set of chemicals.

Of the chemicals monitored during both cycles, only the 11 shown below are on the carcinogen list. These carcinogens consist of five heavy metals, two metabolites of arsenic (DMA and MMA), 2 PFAS, metabolites of PAHs, and a single herbicide (2,4-D).

Minnesota

Minnesota established a biomonitoring program by state law in 2007. This program has conducted a variety of projects including a study of PFAS exposure in urban adults, a study of arsenic levels in child living in a neighborhood with known soil contamination, plastic and paraben exposure levels for pregnant women in

Minneapolis, and mercury and other heavy metal blood concentrations of newborns in the Twin Cities and Lake Superior region of the state.

Staff at MDoH kindly shared demographically-stratified results from these projects, as available. More specifically, results come from the state's MN FEET study and the 2018 Healthy Rural and Urban Kids pilot project. Of note there is meaningful heterogeneity in exposure to arsenic, methylmercury, lead, and 2,4-D across racial/ethnic and income subgroups.

In 2019, the MDoH was awarded a CDC grant to perform statewide biomonitoring on the child-aged population. Data collection is ongoing; MDoH intends to perform disparity analyses for all chemical analytes and will share in a public report when available. The state is surveilling a set of 62 chemicals, all by urine, including the standard metals panel, PFAS, pesticides, phthalates, flame retardants, phenols, and metabolites of PAHs. Of these chemicals, nine are carcinogens.

Iowa

Iowa was awarded a CDC grant for development of a biomonitoring program in 2019. The program is currently in progress, with a summary report describing (sub-)population-level findings expected in December 2024. Surveillance focuses two particular sub-populations: IA residents who rely on private wells for drinking water and pregnant women. The program will study exposure to 32 individual chemicals. These consist of the usual set of metals, a group of neonicotinoid insecticides that may be found in private wells, and several phenols for which exposure during pregnancy is of potential concern.

Of these 32 monitored chemicals, only the set of 5 metals shown below are on the carcinogen list. Because the program is ongoing, no stratified statistics are current available.

New York

New York state was awarded a CDC grant for development of a biomonitoring program in 2019. Sample collection is complete in some regions of the state, while participant recruitment is ongoing in others. Eligibility was restricted to adults aged 20 years or older who reside at randomly selected addresses across the state.

The program is taking blood samples to screen for eleven unique PFAS and seven heavy metals. Urine samples are tested for a group of 17 heavy metals and pesticides including organophosphates, phenoxy acids, and pyrethroids. Unfortunately, the exact pesticide chemicals being screened for were not public information at time of writing and NY state did not respond to outreach.

Of the known chemicals being monitored by NY state, only 5 metals and 2 PFAS, shown below, are on the carcinogen list. Again, because the program is ongoing, no stratified statistics are currently available.

Massachusetts

Massachusetts was awarded a CDC grant for development of their state biomonitoring during the 2014 award cycle. From 2014-2019, MA DPH conducted both a statewide representative study and a smaller-scale study targeting low income residents. Unfortunately, MA DPH was unable to provide stratified biomonitoring statistics from these studies at this time; the program's funding expired before a final report with such information could be drafted. MA DPH staff plans to finish this report in the near future for public consumption, but in the meantime was able to provide a list of analytes that were monitored during the study period.

Michigan

Michigan has varying biomonitoring projects complete and underway. Michigan DHHS received a biomonitoring grant in 2019 to perform a large-scale, state-representative exposure study. The state is measuring 197 analytes, including 45 PFAS, 100 PCBs, 17 organochlorine pesticides, 24 heavy metals, 10 PDBEs, and PBB. Most of the sample collection has already been completed. Of these 197 analytes, 19 are on the carcinogen list.

Other notable governmental biomonitoring efforts

Several states not listed above were recipients of biomonitoring grants from the CDC in 2014 but did not publicly provide useful data from their programs nor effective contact information. These states include Virginia and the Four Corners State Consortium (Utah, Arizona, Colorado, and New Mexico). The National Biomonitoring Network includes in its ranks several states that do not yet have biomonitoring programs but aspire to in the future; these states include TX, LA, AR, TN, SC, NC, WI, VT, RI, and DC.

In multiple conversations with scientists at state health departments, it was noted that New York City has previously done two rounds of biomonitoring (NYC HANES). In the most recent 2013-2014 survey, however, blood and urine samples for relevant carcinogens were limited; only information on Pb and Hg were available. The earlier 2004 survey round produced academic publications where demographically-stratified summary statistics were presented for Pb, Hg, and Cd, as well as several pesticides (McKelvey et al. (2007); McKelvey et al. (2013)).

ATSDR and its state health department partners conducted the Biomonitoring of Great Lakes Populations program from 2010-2020. This program targeted vulnerable populations (including subsistence anglers, immigrants, and tribal communities) in the Great Lakes region (MN, WI, MI, and NY). Carcinogenic legacy contaminants and chemicals of emerging concern were measured in blood and urine, including Pb, Hg, Mirex, Hexachlorobenzene, DDT, DDE, PCBs, PAHs, PFAS, and Toxaphene. Data collected from the multiple cross-sectional studies were not available in a consistent and accessible public fashion, so were not included in this appendix.

E Full list of candidate carcinogens

Table E1 below contains information on the full set of carcinogens considered in this analysis. The following describes the information in each column as well as the data sources.

Column (1) describes the chemical name. Column (2) is the CompTox DTXSID used to uniquely identify each carcinogen throughout the analysis. Column (3) is the chemical’s CAS number. Column (4) denotes whether the chemical was a member of the IARC’s carcinogen groupings: group 1 is known carcinogenic, group 2A is probably carcinogenic, and group 2B is possibly carcinogenic. Column (5) denotes whether the chemical was a member of the NTP’s ROC groupings: “Known” is a known carcinogen, and “RAHC” is a reasonably anticipated human carcinogen. Column (6) denotes whether a chemical is listed as a carcinogen on California’s Proposition 65 list. Column (7) denotes whether a chemical is a substance of very high concern due to carcinogenicity in the EU, UK, or both. Column (8) denotes whether a chemical is listed as active on the TSCA Inventory (T stands for true). Column (9) denotes it was reported as actively manufactured or imported into the U.S. per EPA’s 2020 Chemical Data Reporting. Column (10) denotes whether the chemical was reported in version 1 of the harmonized Multimedia Monitoring Database. Column (11) denotes the numeric count of patents in which a chemical was mentioned, per PubChem. Column (12) denotes the numeric count of scientific articles in which a chemical was mentioned, per PubChem. And Column (13) denotes whether based on the aforementioned criteria, we ultimately considered the carcinogen to be ubiquitous.

Table E1: Complete list of candidate carcinogens

Chemical Name	DTXSID	CAS	IARC	ROC	Prop65	SVHC	TSCA Active	CDR 2020	MMDBV1	Pat. Count	Lit. Count	Ubiq. Flag
(+)-Diclofop-methyl	DTXSID0032605	51338-27-3			T				T	18020	422	T
(2E,4E)-2,4-Hexadienal	DTXSID2025391	142-83-6	2B		T		T			2041	280	T
(-32'-P)Phosphane	DTXSID80932734	14596-37-3	1							7438	58	T
1,1'-Biphenyl, 2,2',3,3',	DTXSID9065572	13654-09-6		RAHC						3374	285	T
1,1,1,2-Tetrachloroethane	DTXSID2021317	630-20-6	2B		T		T		T	37408	1534	T
1,1,1-Trichloroethane	DTXSID0021381	71-55-6			T		T	T	T	71988	3850	T
1,1,2,2-Tetrachloroethane	DTXSID7021318	79-34-5	2B		T		T	T	T	117420	1210	T
1,1,2-Trichloroethane	DTXSID5021380	79-00-5			T		T	T	T	28732	757	T
1,1-Dichloroethane	DTXSID1020437	75-34-3			T		T	T	T	52627	6186	T
1,1-Dichloroethylene	DTXSID8021438	75-35-4	2B		T		T	T	T	24947	2693	T
1,1-Dimethylhydrazine	DTXSID1020516	57-14-7	2B	RAHC	T		T			37431	2195	T
1,2,3,4,5,6-Hexachlorocyc	DTXSID7020687	608-73-1	2B	RAHC	T				T	104118	10426	T
1,2,3-Trichloropropane	DTXSID9021390	96-18-4	2A	RAHC	T	Both	T	T	T	6848	331	T
1,2-(Methylenedioxy)-4-pr	DTXSID7020475	94-58-6	2B		T		T	T	T	612	33	T
1,2-Benzenediol	DTXSID3020257	120-80-9	2B		T		T	T	T	109916	22836	T
1,2-Dibromo-3-chloropropa	DTXSID3020413	96-12-8	2B	RAHC	T		T		T	6152	1119	T
1,2-Dibromoethane	DTXSID3020415	106-93-4	2A	RAHC	T		T	T	T	75644	3132	T
1,2-Dichloroethane	DTXSID6020438	107-06-2	2B	RAHC	T	Both	T	T	T	46484	8353	T
1,2-Dichloropropane	DTXSID0020448	78-87-5	1		T		T	T	T	25792	741	T
1,2-Diethylhydrazine	DTXSID1043754	1615-80-1	2B		T					510	63	
1,2-Dimethylhydrazine	DTXSID0041228	540-73-8	2A		T					7574	2763	T
1,2-Diphenylhydrazine	DTXSID7020710	122-66-7		RAHC	T		T		T	4109	631	T
1,2-Epoxybutane	DTXSID6020569	106-88-7	2B				T	T		45747	505	T
1,2-Phenylenediamine	DTXSID3025881	95-54-5	2B		T		T	T	T	100546	10564	T
1,2-Phenylenediamine dihy	DTXSID3021140	615-28-1	2B				T	T		9832	1443	T
1,2-Propylene oxide	DTXSID5021207	75-56-9	2B	RAHC	T	Both	T	T	T	71341	13480	T
1,3-Butadiene	DTXSID3020203	106-99-0	1	Kwn	T		T	T	T	131292	18381	T
1,3-Dichloro-2-propanol	DTXSID6025010	96-23-1	2B		T		T	T	T	16827	434	T
1,3-Dichloropropene	DTXSID1022057	542-75-6	2B	RAHC	T		T	T	T	12543	319	T
1,3-Dinitropropene	DTXSID5074975	75321-20-9	2B		T					16	124	
1,3-Propane sultone	DTXSID8021195	1120-71-4	2A	RAHC	T	Both	T			35746	576	T
1,4-Dichloro-2-butene	DTXSID9027310	764-41-0			T		T		T	1794	39	T
1,4-Dichloro-2-nitrobenze	DTXSID7052602	89-61-2	2B		T		T		T	1788	82	T
1,4-Dichlorobenzene	DTXSID1020431	106-46-7	2B	RAHC	T		T	T	T	88600	3229	T
1,4-Dioxane	DTXSID4020533	123-91-1	2B	RAHC	T	EU	T	T	T	98992	33889	T
1,6-Dinitropropene	DTXSID90872819	42397-64-8	2B	RAHC	T					78	219	T
1,8-Dinitropropene	DTXSID2073514	42397-65-9	2B	RAHC	T					225	244	T

Return to: Section 2.1.1. Continued, next page...

Table E1: Complete list of candidate carcinogens

Chemical Name	DTXSID	CAS	IARC	ROC	Prop65	SVHC	TSCA Active	CDR 2020	MMDBV1	Pat. Count	Lit. Count	Ubic. Flag
1-Amino-2,4-dibromanthra	DTXSID4039235	81-49-2	2B	RAHC	T		T			417	37	T
1-Amino-2-methylantraqui	DTXSID7020057	82-28-0		RAHC	T		T			3706	150	T
1-Bromo-3-chloropropane	DTXSID1051565	109-70-6	2B		T		T	T		12322	512	T
1-Bromopropane	DTXSID6021874	106-94-5	2B	RAHC	T		T	T		48362	1069	T
1-Chloro-2-nitrobenzene	DTXSID0020280	88-73-3	2B		T		T			11434	964	T
1-Chloro-4-(trifluorometh	DTXSID7024821	98-56-6	2B		T		T	T		5199	203	T
1-Chloro-4-nitrobenzene	DTXSID5020281	100-00-5	2B		T		T			10406	1312	T
1-Ethyl-1-nitrosourea	DTXSID8020593	759-73-9	2A	RAHC	T		T			4396	4967	T
1-Hydroxyanthraquinone	DTXSID1020722	129-43-1	2B		T					3821	676	T
1-Methoxy-2-nitrobenzene	DTXSID3020962	91-23-6	2A	RAHC	T		T			2587	301	T
1-Naphthalenol, 1-(N-meth	DTXSID9020247	63-25-2			T		T		T	56447	5942	T
1-Naphthylamine	DTXSID7020920	134-32-7			T		T		T	78091	4475	T
1-Nitropyrene	DTXSID6020983	5522-43-0	2A	RAHC	T		T		T	1401	1067	T
1-tert-Butoxy-2-propanol	DTXSID8025967	57018-52-7	2B		T		T			13984	50	T
1-[(3-(2-Chlorophenyl)-2-	DTXSID901034223	135319-73-2			T							
17alpha-Ethinylestradiol	DTXSID5020576	57-63-6		Kwn	T		T		T	54975	16986	T
17beta-Estradiol	DTXSID0020573	50-28-2		Kwn	T				T	92825	143623	T
2',3'-Dideoxycytidine	DTXSID0023747	7481-89-2	2B		T					49690	1815	T
2,2'-Bioxirane	DTXSID0041307	1464-53-5		RAHC	T		T			9677	1138	T
2,2-Dimethylvinyl chlorid	DTXSID5020520	513-37-1	2B	RAHC	T					704	73	
2,3,4,7,8-Pentachlorodibe	DTXSID7030066	57117-31-4	1						T	428	278	T
2,3,7,8-Tetrachlorodibenz	DTXSID2021315	1746-01-6	1	Kwn	T				T	75481	16728	T
2,3-Benzofuran	DTXSID6020141	271-89-6	2B		T		T			107257	3657	T
2,3-Dibromopropanol	DTXSID7021817	96-13-9	2B	RAHC	T	EU	T		T	3890	94	T
2,4,5-Trimethylaniline	DTXSID9021398	137-17-7			T					707	75	
2,4,6-Trichlorophenol	DTXSID5021386	88-06-2	2B	RAHC	T		T		T	6269	1983	T
2,4,6-Trinitrotoluene	DTXSID7024372	118-96-7			T		T	T		32001	4985	T
2,4-Diaminoanisole	DTXSID5041358	615-05-4	2B		T		T			4840	168	T
2,4-Diaminotoluene	DTXSID4020402	95-80-7	2B	RAHC	T	Both	T	T		69530	980	T
2,4-Dichloronitrobenzene	DTXSID3024998	611-06-3	2B		T		T		T	1510	108	T
2,4-Dichlorophenoxyacetic	DTXSID0020442	94-75-7	2B				T	T	T	51498	23242	T
2,4-Dinitrotoluene	DTXSID0020529	121-14-2	2B		T	Both	T	T	T	9869	1503	T
2,6-Dimethylaniline	DTXSID8026307	87-62-7	2B		T		T	T		11788	643	T
2,6-Dinitrotoluene	DTXSID5020528	606-20-2	2B		T		T		T	1994	604	T
2-Acetylaminofluorene	DTXSID0039227	53-96-3		RAHC	T		T		T	2355	4181	T
2-Amino-1-methyl-6-phenyl	DTXSID3037628	105650-23-5	2B	RAHC	T				T	1571	1413	T
2-Amino-3-methyl-3H-imida	DTXSID4020745	76180-96-6	2A	RAHC	T				T	262	473	T

Return to: Section 2.1.1. Continued, next page...

Table E1: Complete list of candidate carcinogens

Chemical Name	DTXSID	CAS	IARC	ROC	Prop65	SVHC	TSCA Active	CDR 2020	MMDBV1	Pat. Count	Lit. Count	Ubiq. Flag
2-Amino-4-chlorophenol	DTXSID5024473	95-85-2	2B		T		T			5706	193	T
2-Amino-5-(5-nitro-2-fury	DTXSID7020059	712-68-5	2B		T					81	26	
2-Amino-alpha-carboline	DTXSID7020001	26148-68-5	2B		T				T	72	170	T
2-Aminoanthraquinone	DTXSID6020068	117-79-3		RAHC	T		T			12302	193	T
2-Aminofluorene	DTXSID1049223	153-78-6			T					1627	1066	T
2-Anisidine	DTXSID5023877	90-04-0	2A	RAHC	T	Both	T	T		19169	969	T
2-Bromopropane	DTXSID7030197	75-26-3			T		T	T		29140	529	T
2-Ethylhexyl acrylate	DTXSID9025297	103-11-7	2B		T		T	T		80423	486	T
2-Mercaptobenzothiazole	DTXSID1020807	149-30-4	2A		T		T	T		105584	2307	T
2-Methoxy-5-methylaniline	DTXSID1020350	120-71-8	2B	RAHC	T	Both	T	T		5313	131	T
2-Methoxyaniline hydrochl	DTXSID8020092	134-29-2	2A	RAHC	T					94	91	
2-Methyl-1-nitroanthraqui	DTXSID7020847	129-15-7	2B		T					138	33	
2-Methylaniline	DTXSID1026164	95-53-4	1	Kwn	T	Both	T	T	T	85377	5852	T
2-Methylaniline hydrochlo	DTXSID7021364	636-21-5			T		T			645	238	T
2-Methylimidazole	DTXSID4022107	693-98-1	2B		T		T	T		97732	2231	T
2-Naphthylamine	DTXSID2020921	91-59-8	1	Kwn	T				T	25477	3105	T
2-Nitrofluorene	DTXSID2020971	607-57-8	2B		T		T		T	3168	560	T
2-Nitropropane	DTXSID6020981	79-46-9	2B	RAHC	T		T	T	T	9814	762	T
2-Nitrotoluene	DTXSID4025791	88-72-2	2A	RAHC	T		T		T	24089	1572	T
2-Phenylphenol	DTXSID2021151	90-43-7			T		T		T	81633	2003	T
3'-Azido-3'-deoxythymidin	DTXSID8020127	30516-87-1	2B		T					66977	20676	T
3,3',4,4',5-Pentachlorobi	DTXSID3032179	57465-28-8	1						T	1103	912	T
3,3',4,4'-Tetrachlorozob	DTXSID6026086	14047-09-7	2A		T					83	93	
3,3',5,5'-Tetrabromobisph	DTXSID1026081	79-94-7	2A		T	EU	T	T	T	69436	1669	T
3,3'-Dichloro-4,4'-diamin	DTXSID5074842	28434-86-8	2B		T					1068	23	
3,3'-Dichlorobenzidine	DTXSID6020432	91-94-1	2B	RAHC	T		T		T	10366	266	T
3,3'-Dichlorobenzidine di	DTXSID1020433	612-83-9		RAHC	T		T	T		158	24	T
3,3'-Dimethylbenzidine	DTXSID5024059	119-93-7	2B	RAHC	T		T	T	T	28842	1116	T
3,3'-Dimethylbenzidine di	DTXSID6020511	612-82-8		RAHC	T		T			72	25	T
3,4-Dihydroxycinnamic aci	DTXSID5020231	331-39-5	2B		T					9282	2010	T
3,7-Dinitrofluoranthene	DTXSID20147305	105735-71-5	2B		T					1	31	
3,9-Dinitrofluoranthene	DTXSID40177028	22506-53-2	2B		T						41	
3-(3,5-Dichlorophenyl)-N-	DTXSID3024154	36734-19-7			T				T	44692	1283	T
3-Amino-9-ethylcarbazole	DTXSID2020054	6109-97-3			T						48	
3-Bromo-2,2-bis(bromometh	DTXSID9024641	1522-92-5				EU				2137	33	T
3-Chloro-1,2-propanediol	DTXSID4020664	96-24-2	2B		T		T	T		15314	1961	T
3-Chloro-2-methylpropene	DTXSID1020279	563-47-3	2B	RAHC	T		T	T		9777	359	T

Return to: Section 2.1.1. Continued, next page...

Table E1: Complete list of candidate carcinogens

Chemical Name	DTXSID	CAS	IARC	ROC	Prop65	SVHC	TSCA Active	CDR 2020	MMDBV1	Pat. Count	Lit. Count	Ubiq. Flag
3-Chloro-4-(dichloromethyl)-3-methylcholanthrene	DTXSID6020276	77439-76-0	2B		T					107	248	T
3-Methylcholanthrene	DTXSID0020862	56-49-5			T		T		T	3441	10553	T
3-Nitrobenzanthrone	DTXSID60881271	17117-34-9	2B							20	167	T
4'-(9-Acridinylamino)methyl-4'-bis(dimethylamino)-4,4'-diaminobiphenyl methyl-4'-methylenebis(2-chloro-4'-methylenebis(N,N-dimethyl-4'-methylenebis(o-toluidine))	DTXSID4022604	51264-14-3	2B		T					92967	2843	T
4,4'-Bis(dimethylamino)-4,4'-diaminobiphenyl methyl-4'-methylenebis(2-chloro-4'-methylenebis(N,N-dimethyl-4'-methylenebis(o-toluidine))	DTXSID00204642	561-41-1				Both				3	0	
4,4'-Diaminobiphenyl methyl-4'-methylenebis(2-chloro-4'-methylenebis(N,N-dimethyl-4'-methylenebis(o-toluidine))	DTXSID6022422	101-77-9	2B	RAHC	T	Both	T	T		44067	1102	T
4,4'-Methylenebis(2-chloro-4'-methylenebis(N,N-dimethyl-4'-methylenebis(o-toluidine))	DTXSID5020865	101-14-4	1	RAHC	T	Both		T	T	26656	291	T
4,4'-Methylenebis(N,N-dimethyl-4'-methylenebis(o-toluidine))	DTXSID5020869	101-61-1	2B	RAHC	T	Both	T			9674	185	T
4,4'-Methylenebis(o-toluidine)	DTXSID5020867	838-88-0	2B		T	Both	T			13684	76	T
4,4'-Methylenedianiline d	DTXSID9020871	13552-44-8		RAHC	T					6	26	
4,4'-Oxydianiline	DTXSID0021094	101-80-4	2B	RAHC	T	Both	T	T		81122	886	T
4,4'-Thiodianiline	DTXSID9021344	139-65-1	2B	RAHC	T		T			28691	128	T
4-(N-Methyl-N-nitrosamino)-4-amino-2-nitrophenol	DTXSID3020881	64091-91-4		RAHC	T				T	2005	1756	T
4-Amino-2-nitrophenol	DTXSID6020064	119-34-6			T		T			1604	89	T
4-Androstene-3,17-dione	DTXSID8024523	63-05-8			T							
4-Biphenylamine	DTXSID5020071	92-67-1	1	Kwn	T	Both	T		T	19459	16688	T
4-Chloro-1,2-diaminobenzene	DTXSID5020283	95-83-0	2B	RAHC	T		T		T	12729	1448	T
4-Chloro-2-methylaniline	DTXSID1041508	95-69-2	2A	RAHC	T		T			4362	166	T
4-Chloro-2-methylaniline	DTXSID0020288	3165-93-3		RAHC			T			4196	204	T
4-Chloroaniline	DTXSID9020295	106-47-8	2B		T		T	T		518	52	T
4-Chloroaniline hydrochloride	DTXSID4020296	20265-96-7			T		T		T	30952	2557	T
4-Chlorobenzotrithiolide	DTXSID2027593	5216-25-1			T		T			105	181	T
4-Methyl-2-pentanone	DTXSID5021889	108-10-1	2B		T		T	T		960	59	T
4-Methylimidazole	DTXSID9025617	822-36-6	2B		T		T	T	T	169480	7089	T
4-Nitrobiphenyl	DTXSID9041522	92-93-3			T		T			27997	525	T
4-Nitropyrene	DTXSID5074844	57835-92-4	2B	RAHC	T		T		T	1923	457	T
4-Nitrosodiphenylamine	DTXSID1021031	156-10-5			T		T		T	48	73	T
4-Vinyl-1-cyclohexene diol	DTXSID0020604	106-87-6	2B	RAHC	T		T			1176	101	T
4-Vinylcyclohexene	DTXSID3021437	100-40-3	2B		T		T	T		25715	296	T
5,5-Diphenylhydantoin	DTXSID8020541	57-41-0	2B	RAHC	T				T	15726	464	T
5-Azacytidine	DTXSID9020116	320-67-2	2A	RAHC	T				T	64206	41453	T
5-Chloro-2-methylaniline	DTXSID5020287	95-79-4			T		T			105360	10779	T
5-Chloro-3-methyl-4-nitro-5-methoxyphenol	DTXSID3058454	6814-58-0			T				T	2809	105	T
5-Methoxyphenol	DTXSID1025560	484-20-8	2A		T					159	19	
5-Methylchrysene	DTXSID6063143	3697-24-3	2B	RAHC	T				T	3739	1743	T
5-Nitroacenaphthene	DTXSID3020960	602-87-9	2B		T		T		T	363	176	T
6-Methyl-2-thiouracil	DTXSID2020890	56-04-2	2B		T		T		T	2805	92	T
										2388	1622	T

Return to: Section 2.1.1. Continued, next page...

Table E1: Complete list of candidate carcinogens

Chemical Name	DTXSID	CAS	IARC	ROC	Prop65	SVHC	TSCA Active	CDR 2020	MMDBV1	Pat. Count	Lit. Count	Ubiqu. Flag
6-Nitrochrysene	DTXSID9075454	7496-02-8	2A	RAHC	T				T	177	232	T
6-Propyl-2-thiouracil	DTXSID5021209	51-52-5	2B	RAHC	T		T			21250	9048	T
7,12-Dimethylbenz(a)anthr	DTXSID1020510	57-97-6			T		T		T	8066	8782	T
7H-Dibenzo[c,g]carbazole	DTXSID9059755	194-59-2	2B	RAHC	T					517	131	
8-Methoxypsoralen	DTXSID8020830	298-81-7	1	Kwn	T		T			23415	5583	T
AF-2	DTXSID9020033	3688-53-7	2B		T					1057	283	T
Acetaldehyde	DTXSID5039224	75-07-0	2B	RAHC	T		T	T	T	103204	37295	T
Acetaldehyde methylformyl	DTXSID0039225	16568-02-8			T					23	16	
Acetamide	DTXSID7020005	60-35-5	2B		T		T			39321	6893	T
Acetochlor	DTXSID8023848	34256-82-1			T				T	31891	976	T
Acid Red 26	DTXSID8021228	3761-53-3	2B		T		T			545	20	T
Acifluorfen sodium	DTXSID7023853	62476-59-9			T		T		T	9359	83	T
Acrolein	DTXSID5020023	107-02-8	2A									
Acrylamide	DTXSID5020027	79-06-1	2A	RAHC	T		T	T	T	99308	13624	T
Acrylonitrile	DTXSID5020029	107-13-1	2B	RAHC	T		T	T	T	99862	39858	T
Actinolite asbestos	DTXSID40108998	77536-66-4		Kwn			T	T	T	118855	15690	T
Actinomycin D	DTXSID9020031	50-76-0			T					127517	23440	T
Adriamycin hydrochloride	DTXSID3030636	25316-40-9		RAHC	T					98190	63557	T
Aflatoxin	DTXSID4020036	1402-68-2	1	Kwn	T				T			T
Aflatoxin M1	DTXSID40891797	6795-23-9	2B							370	1391	T
Alachlor	DTXSID1022265	15972-60-8			T				T	42062	2297	T
Aldrin	DTXSID8020040	309-00-2			T				T	16830	5301	T
Amides, coco, N,N-bis(hyd	DTXSID0024842	68603-42-9	2B		T		T	T	T			T
Amitrole	DTXSID0020076	61-82-5		RAHC	T		T			43395	3022	T
Ammonium chromate	DTXSID3064857	7788-98-9		Kwn			T			3929	20	T
Ammonium dichromate	DTXSID70872953	7789-09-5		Kwn			T	T		2436	451	T
Amosite	DTXSID5030741	12172-73-5		Kwn					T			T
Aniline	DTXSID8020090	62-53-3	2A									
Aniline hydrochloride	DTXSID3020091	142-04-1	2A		T		T	T	T	103761	44443	T
Anthracene	DTXSID0023878	120-12-7			T		T	T	T	10062	3325	T
Anthracene oil	DTXSID9029067	90640-80-5					T			32643	21569	T
Anthracene oil, anthracen	DTXSID701020329	90640-81-6					T					T
Anthracene oil, anthracen	DTXSID001020330	90640-82-7										
Anthracene oil, anthracen	DTXSID701020331	91995-15-2										
Anthracene oil, anthracen	DTXSID401020332	91995-17-4										
Anthraquinone	DTXSID3020095	84-65-1	2B		T		T	T	T	125902	18957	T
Antimony trioxide	DTXSID4023880	1309-64-4	2B	RAHC	T		T	T		15180	194	T

Return to: Section 2.1.1. Continued, next page...

Table E1: Complete list of candidate carcinogens

Chemical Name	DTXSID	CAS	IARC	ROC	Prop65	SVHC	TSCA Active	CDR 2020	MMDBV1	Pat. Count	Lit. Count	Ubic. Flag
Aramite	DTXSID3020097	140-57-8	2B		T			T	T	3338	136	T
Arecoline	DTXSID3022617	63-75-2	2B							7103	2568	T
Aristolochic acid	DTXSID0040969	313-67-7	1	Kwn						1989	1162	T
Aristolochic acid II	DTXSID00197166	475-80-9		Kwn						15	36	
Aristolochic acids	DTXSID70217421	67123-64-2		Kwn	T						0	
Arsenic	DTXSID4023886	7440-38-2	1	Kwn	T		T	T	T	64758	699	T
Arsenic acid	DTXSID1034341	7778-39-4				Both	T	T	T	1692	12034	T
Arsenic oxide (As ₂ O ₃)	DTXSID0020103	1327-53-3		Kwn		Both	T	T			3288	T
Arsenic(V) pentoxide	DTXSID1034343	1303-28-2				Both	T	T		2078	596	T
Asbestos	DTXSID4023888	1332-21-4		Kwn	T		T	T	T			T
Asbestos, anthophyllite	DTXSID00108999	77536-67-5		Kwn								
Asphalt, oxidized	DTXSID4028250	64742-93-4	2A				T	T				T
Attapulgit	DTXSID6050469	12174-11-7	2B		T							
Auramine	DTXSID7043821	492-80-8	2B		T		T	T		22133	1566	T
Azaserine	DTXSID9020118	115-02-6	2B		T					27327	681	T
Azathioprine	DTXSID4020119	446-86-6	1	Kwn	T					134088	48761	T
Azobenzene	DTXSID8020123	103-33-3			T		T		T	37127	6278	T
Basic Blue 26	DTXSID0038889	2580-56-5				Both	T	T		20957	599	T
Benthiavalcarb-isopropyl	DTXSID9058232	177406-68-7			T					6766	37	T
Benz(a)anthracene	DTXSID5023902	56-55-3	2B	RAHC	T	Both	T	T	T	13146	6002	T
Benz(j)aceanthrylene	DTXSID30174041	202-33-5	2B							164	56	
Benzene	DTXSID3039242	71-43-2	1	Kwn	T		T	T	T	242482	155764	T
Benzo(b)fluoranthene	DTXSID0023907	205-99-2	2B	RAHC	T					2150	3174	T
Benzo(j)fluoranthene	DTXSID8052691	205-82-3	2B	RAHC	T				T	1663	610	T
Benzo[a]pyrene	DTXSID2020139	50-32-8	1	RAHC	T		T	T	T	27386	22457	T
Benzo[c]phenanthrene	DTXSID4075459	195-19-7	2B						T	1457	389	T
Benzo[k]fluoranthene	DTXSID0023909	207-08-9	2B	RAHC	T				T	1993	3131	T
Benzophenone	DTXSID0021961	119-61-9	2B		T		T	T	T	160994	10012	T
Benzotrichloride	DTXSID1020148	98-07-7		RAHC	T		T	T	T	9211	390	T
Benzyl Violet 4B	DTXSID7021441	1694-09-3	2B		T		T	T		13155	47	T
Benzyl chloride	DTXSID0020153	100-44-7			T		T	T	T	61728	3710	T
Beryllium	DTXSID4023913	7440-41-7	1	Kwn	T		T	T	T	35119	2729	T
Beryllium carbonate hydro	DTXSID10984600	66104-24-3		Kwn						2	0	
Beryllium chloride	DTXSID10858756	7787-47-5		Kwn			T			3565	359	T
Beryllium hydrogen phosph	DTXSID70929198	13598-15-7		Kwn						20	20	
Beryllium hydroxide	DTXSID8083932	13327-32-7		Kwn			T	T	T		305	T
Beryllium oxide (BeO)	DTXSID30872818	1304-56-9		Kwn			T	T		61787	2187	T

Return to: Section 2.1.1. Continued, next page...

Table E1: Complete list of candidate carcinogens

Chemical Name	DTXSID	CAS	IARC	ROC	Prop65	SVHC	TSCA Active	CDR 2020	MMDBV1	Pat. Count	Lit. Count	Ubiq. Flag
Beryllium sulfate	DTXSID0020159	13510-49-1		Kwn			T	T		2045	318	T
Beryllium sulfate tetrahy	DTXSID3024603	7787-56-6		Kwn						167	44	
Beryllium zinc orthosilic	DTXSID0094898	25638-88-4		Kwn						15	0	
Bis(4-(dimethylamino)phen	DTXSID2020894	90-94-8	2B	RAHC	T	Both	T			74755	163	T
Bis(chloroethyl) ether	DTXSID9020168	111-44-4			T		T	T		18415	420	T
Bis-(2-chloroethyl)sulfid	DTXSID0037100	505-60-2	1	Kwn	T		T	T		12884	2755	T
Bitumens, extracts of ste	DTXSID801046853	128683-24-9			T							
Bleomycin	DTXSID1030862	11056-06-7	2B									
Bromate	DTXSID8023923	15541-45-4			T				T	28822	978	T
Bromkal 80	DTXSID701016695	61288-13-9		RAHC								
Bromochloroacetic acid	DTXSID4024642	5589-96-8	2B	RAHC	T				T	177	167	T
Bromodichloroacetic acid	DTXSID4024644	71133-14-7		RAHC	T				T	93	103	T
Bromodichloromethane	DTXSID1020198	75-27-4	2B	RAHC	T		T		T	3883	950	T
Bromoethane	DTXSID6020199	74-96-4			T		T	T		74963	2288	T
Bromoform	DTXSID1021374	75-25-2			T		T	T		26927	3334	T
Busulfan	DTXSID3020910	55-98-1	1	Kwn	T					114398	14081	T
Butyl glycidyl ether	DTXSID9024691	2426-08-6	2B		T		T	T		40210	253	T
Butylated hydroxylanisole	DTXSID7020215	25013-16-5	2B	RAHC	T		T	T				T
C.I. Acid Red 114	DTXSID8021224	6459-94-5	2B	RAHC	T		T			585	46	T
C.I. Azoic Diazo Component	DTXSID2020137	92-87-5	1	Kwn	T		T	T		94774	7477	T
C.I. Basic Red 9 monohydr	DTXSID1021247	569-61-9	2B	RAHC	T		T			1	414	T
C.I. Basic Violet 14	DTXSID6021246	632-99-5	2B		T		T			16815	8153	T
C.I. Direct Black 38	DTXSID7020184	1937-37-7		Kwn	T	Both	T			1	74	T
C.I. Direct Blue 14	DTXSID4026268	72-57-1	2B		T		T			87327	3061	T
C.I. Direct Blue 15	DTXSID7020186	2429-74-5	2B	RAHC	T		T			6	45	T
C.I. Direct Blue 218	DTXSID2020187	28407-37-6	2B		T		T	T			20	T
C.I. Direct Blue 6	DTXSID2020185	2602-46-2		Kwn	T		T				30	T
C.I. Direct Brown 95	DTXSID3020201	16071-86-6		Kwn	T		T				27	T
C.I. Direct Red 28	DTXSID4022816	573-58-0				Both	T			32994	3652	T
C.I. Disperse Black 6	DTXSID3025091	119-90-4	2B	RAHC	T		T			30535	3244	T
C.I. Disperse Black 6 dih	DTXSID1020485	20325-40-0			T			T		168	703	T
C.I. Disperse Blue 1	DTXSID7020188	2475-45-8	2B	RAHC	T		T			4526	106	T
C.I. Disperse Yellow 3	DTXSID6021450	2832-40-8			T		T			3167	63	T
C.I. Oxidation Base 12A	DTXSID7020398	39156-41-7		RAHC	T		T			142	81	T
C.I. Pigment Orange 5	DTXSID6029258	3468-63-1			T		T	T		20920	22	T
C.I. Pigment Red 104	DTXSID9051321	12656-85-8				Both	T	T				T
C.I. Pigment Red 53	DTXSID3021229	5160-02-1			T		T	T		26377	36	T

Return to: Section 2.1.1. Continued, next page...

Table E1: Complete list of candidate carcinogens

Chemical Name	DTXSID	CAS	IARC	ROC	Prop65	SVHC	TSCA Active	CDR 2020	MMDBV1	Pat. Count	Lit. Count	Ubic. Flag
C.I. Pigment Yellow 34	DTXSID8051336	1344-37-2				Both	T	T			0	T
C.I. Solvent Orange 2	DTXSID6025808	2646-17-5	2B		T		T			5088	30	T
C.I. Solvent Red 80	DTXSID6024838	6358-53-8	2B		T		T			1447	25	T
C.I. Solvent Yellow 1	DTXSID6024460	60-09-3	2B		T	Both	T			26083	995	T
C.I. Solvent Yellow 14	DTXSID4021135	842-07-9			T		T			4292	246	T
C.I. Solvent Yellow 2	DTXSID5020491	60-11-7	2B	RAHC	T		T		T	7038	3289	T
C10-12 chloroalkanes	DTXSID10872316	108171-26-2		RAHC	T							
Cadmium	DTXSID1023940	7440-43-9	1	Kwn	T	Both	T	T				T
Cadmium chloride	DTXSID6020226	10108-64-2		Kwn		Both	T			22	20	T
Cadmium dinitrate	DTXSID7044504	10325-94-7				Both	T	T		4	0	T
Cadmium fluoride (CdF2)	DTXSID6064878	7790-79-6				Both	T			1797	393	T
Cadmium hydroxide (Cd(OH)	DTXSID9066671	21041-95-2				Both	T	T		4	450	T
Cadmium oxide	DTXSID0024715	1306-19-0				Both	T	T		37551	2272	T
Cadmium sulfate (1:1)	DTXSID1020229	10124-36-4				Both	T			6495	2680	T
Cadmium sulfate (1:1) hyd	DTXSID0020230	7790-84-3				EU				110	0	
Cadmium sulfate-water (1	DTXSID90934502	15244-35-6				EU				3	27	
Cadmium sulfide	DTXSID30893280	1306-23-6				Both	T			27878	6772	T
Cadmium(II) chloride hydr	DTXSID4040183	7790-78-5				EU					0	
Calcium arsenate	DTXSID6058182	7778-44-1		Kwn		Both	T			5466	550	T
Calcium monochromate	DTXSID0024717	13765-19-0		Kwn			T			4140	267	T
Captafol	DTXSID4020242	2425-06-1	2A	RAHC	T				T	23954	564	T
Captan	DTXSID9020243	133-06-2			T		T		T	41381	2804	T
Carbazole	DTXSID4020248	86-74-8	2B		T		T		T	128344	6874	T
Carbon black	DTXSID7051216	1333-86-4	2B		T		T	T				T
Carbon nanotubes	DTXSID301020377	308068-56-6	2B									
Carbon tetrachloride	DTXSID8020250	56-23-5	2B	RAHC	T		T	T		36125	44870	T
Carbonic acid, cadmium sa	DTXSID60883417	513-78-0				Both	T			2935	439	T
Carboxymethylnitrosourea	DTXSID3020251	60391-92-6			T					9	3	
Carrageenan, acid-degrade	DTXSID8020254	53973-98-1	2B		T							
Chinomethionate	DTXSID2032342	2439-01-2			T		T		T	34021	232	T
Chloral	DTXSID7024744	75-87-6	2A		T		T	T	T	36366	3395	T
Chloral hydrate	DTXSID7020261	302-17-0	2A		T		T					
Chlorambucil	DTXSID7020263	305-03-3	1	Kwn	T					49686	18512	T
Chloramphenicol	DTXSID7020265	56-75-7	2A	RAHC			T		T	126750	10668	T
Chloramphenicol sodium su	DTXSID2024747	982-57-0			T					134621	59298	T
Chlordane	DTXSID7020267	57-74-9	2B		T				T	694	178	T
					T				T	12895	4074	T
Chlordimeform	DTXSID2037508	6164-98-3			T					16957	549	T

Return to: Section 2.1.1. Continued, next page...

Table E1: Complete list of candidate carcinogens

Chemical Name	DTXSID	CAS	IARC	ROC	Prop65	SVHC	TSCA Active	CDR 2020	MMDBV1	Pat. Count	Lit. Count	Ubiq. Flag
Chlorendic acid	DTXSID2020268	115-28-6	2B	RAHC	T		T		T	10426	39	T
Chlornaphazine	DTXSID3060087	494-03-1	1		T					17394	135	T
Chlorobenzilate	DTXSID9020299	510-15-6			T		T		T	15921	358	T
Chloroethane	DTXSID1020302	75-00-3			T		T	T	T	108095	3798	T
Chloroform	DTXSID1020306	67-66-3	2B	RAHC	T		T	T	T	141399	159729	T
Chloromethyl methyl ether	DTXSID6020307	107-30-2		Kwn	T		T	T		26826	643	T
Chloroprene	DTXSID5020316	126-99-8	2B	RAHC	T		T		T	35844	1226	T
Chlorothalonil	DTXSID0020319	1897-45-6	2B		T		T		T	130110	2390	T
Chlorotrianisene	DTXSID1021299	569-57-3			T					33476	294	T
Chlorozotocin	DTXSID9020320	54749-90-5	2A	RAHC	T					16724	0	T
Chromic(VI) acid	DTXSID8034455	7738-94-5				Both	T	T		13236	6880	T
Chromium (VI) ion	DTXSID7023982	18540-29-9	1	Kwn	T				T	44120	4086	T
Chromium chromate (Cr2(Cr	DTXSID10894154	24613-89-6				Both	T	T		2353	20	T
Chromium lead oxide	DTXSID201015373	11119-70-3		Kwn								
Chromium trioxide	DTXSID0040125	1333-82-0		Kwn		Both	T	T		40206	5280	T
Chrysene	DTXSID0022432	218-01-9	2B		T	Both	T		T	38289	6233	T
Chrysotile	DTXSID0030742	12001-29-5		Kwn							0	
Cidofovir	DTXSID3043734	113852-37-2			T					30333	2179	T
Cinnamyl anthranilate	DTXSID3020330	87-29-6			T		T			130	3	T
Cisplatin	DTXSID4024983	15663-27-1	2A	RAHC	T		T					T
Clofibrate	DTXSID3020336	637-07-0			T					52079	8173	T
Clomiphene citrate (1:1)	DTXSID8020337	50-41-9			T						7963	T
Coal tar	DTXSID1027683	8007-45-2	1	Kwn			T					T
Cobalt	DTXSID1031040	7440-48-4	2B	RAHC	T		T	T	T	275685	22413	T
Cobalt chloride	DTXSID9040180	7646-79-9		RAHC		Both	T	T		61740	2243	T
Cobalt oxide	DTXSID801014561	11104-61-3		RAHC			T	T				T
Cobalt sulfate	DTXSID1031042	10124-43-3		RAHC	T	Both	T	T		40389	1059	T
Cobalt sulfate heptahydra	DTXSID7020340	10026-24-1	2B	RAHC	T		T	T		2870	152	T
Cobalt sulfide	DTXSID301015176	12653-56-4		RAHC			T	T				T
Cobalt tungsten carbide	DTXSID20209477	60674-89-7		RAHC							0	
Cobalt(II) acetate	DTXSID6026373	71-48-7		RAHC		Both	T	T		40289	28	T
Cobalt(II) carbonate	DTXSID7052151	513-79-1				Both	T	T		15977	382	T
Cobalt(II) nitrate	DTXSID9064970	10141-05-6		RAHC		Both	T	T		42592	4861	T
Cobalt(II) oxide	DTXSID6051649	1307-96-6			T		T	T		24528	6127	T
Cresote	DTXSID2023987	8001-58-9	2A		T		T	T	T			T
Cristobalite	DTXSID5041801	14464-46-1		Kwn			T	T				T
Crotonaldehyde	DTXSID8024864	4170-30-3	2B				T	T	T	12859	288	T

Return to: Section 2.1.1. Continued, next page...

Table E1: Complete list of candidate carcinogens

Chemical Name	DTXSID	CAS	IARC	ROC	Prop65	SVHC	TSCA Active	CDR 2020	MMDBV1	Pat. Count	Lit. Count	Ubiq. Flag
Gumene	DTXSID1021827	98-82-8	2B RAHC	RAHC	T		T	T	T	150762	6690	T
Cupferron	DTXSID1020352	135-20-6	2B RAHC	RAHC	T		T			1678	56	T
Cycasin	DTXSID2057574	14901-08-7	2B		T						99	
Cyclopenta[cd]pyrene	DTXSID7074822	27208-37-3	2A		T				T	78	186	T
Cyclophosphamide	DTXSID5020364	50-18-0		Kwn	T					199306	119647	T
Cyclophosphamide monohydr	DTXSID6024888	6055-19-2			T					2152	54077	T
Cyclosporin A	DTXSID0020365	59865-13-3		Kwn	T					152041	63260	T
Cytembena	DTXSID5020368	21739-91-3			T						32	
D&C Red No. 8	DTXSID6062173	2092-56-0			T		T			2330	15	T
DDT	DTXSID4020375	50-29-3	2A RAHC	RAHC	T		T		T	57337	16796	T
DE 71	DTXSID30873481	117148-05-7			T							
Dacarbazine	DTXSID0020369	4342-03-4	2B RAHC	RAHC	T					133831	8394	T
Daminozide	DTXSID9020370	1596-84-5			T					25668	593	T
Danthron	DTXSID9020328	117-10-2	2B RAHC	RAHC	T		T			5852	896	T
Daunorubicin	DTXSID7022883	20830-81-3	2B		T					153251	19185	T
Di(2-ethylhexyl) phthalat	DTXSID5020607	117-81-7	2B RAHC	RAHC	T		T	T	T	46944	6680	T
Diazinon	DTXSID9020407	333-41-5	2A				T		T	66874	5763	T
Diazoaminobenzene	DTXSID9024934	136-35-6		RAHC	T		T			6406	235	T
Dibenz(c,h)acridine	DTXSID30176929	224-53-3	2B							49	47	
Dibenz[a,c]anthracene	DTXSID9049245	215-58-7			T				T	1278	630	T
Dibenz[a,h]acridine	DTXSID3059761	226-36-8	2B RAHC	RAHC	T				T	201	117	T
Dibenz[a,h]anthracene	DTXSID9020409	53-70-3	2A RAHC	RAHC	T		T		T	2576	2872	T
Dibenz[a,i]acridine	DTXSID4059758	224-42-0	2A RAHC	RAHC	T				T	310	115	T
Dibenz[a,i]anthracene	DTXSID8074811	224-41-9			T					399	117	
Dibenzo(a,e)pyrene	DTXSID3052690	192-65-4		RAHC	T				T	594	167	T
Dibenzo[a,h]pyrene	DTXSID4059752	189-64-0	2B RAHC	RAHC	T				T	459	175	T
Dibenzo[a,i]pyrene	DTXSID9059751	189-55-9	2B RAHC	RAHC	T				T	370	190	T
Dibenzo[a,l]pyrene	DTXSID9059753	191-30-0	2A RAHC	RAHC	T				T	349	582	T
Diberyllium orthosilicate	DTXSID10929197	13598-00-0		Kwn						1215	0	
Dibromoacetic acid	DTXSID1023815	631-64-1	2B RAHC	RAHC	T		T		T	1853	327	T
Dibromoacetonitrile	DTXSID3024940	3252-43-5	2B		T		T		T	270	154	T
Dibromochloroacetic acid	DTXSID3031151	5278-95-5		RAHC						62	92	
Dichloroacetic acid	DTXSID2020428	79-43-6	2B RAHC	RAHC	T		T		T	55489	2982	T
Dichloromethane	DTXSID0020868	75-09-2	2A RAHC	RAHC	T		T	T	T	149383	77642	T
Dichlorvos	DTXSID5020449	62-73-7	2B		T		T		T	42009	3965	T
Dichromic acid	DTXSID30872954	13530-68-2					T			6772	259	T
Dieldrin	DTXSID9020453	60-57-1			T	Both			T	16067	8718	T

Return to: Section 2.1.1. Continued, next page...

Table E1: Complete list of candidate carcinogens

Chemical Name	DTXSID	CAS	IARC	ROC	Prop65	SVHC	TSCA Active	CDR 2020	MMDBV1	Pat. Count	Lit. Count	Ubic. Flag
Diesel fuel, marine	DTXSID4025030	77650-28-3	2B									
Diethanolamine	DTXSID3021932	111-42-2	2B	RAHC	T	Both	T	T	T	47679	5065	T
Diethyl sulfate	DTXSID1024045	64-67-5	2A	RAHC	T		T	T		49727	1182	T
Diethylstilbestrol	DTXSID3020465	56-53-1	1	Kwn	T		T		T	55160	14993	T
Diglycidyl resorcinol eth	DTXSID2020470	101-90-6	2B	RAHC	T		T	T		15492	85	T
Digoxin	DTXSID5022934	20830-75-5	2B				T		T	77175	31032	T
Dihydroxymethylfuratrizin	DTXSID401045239	794-93-4	2B		T							
Diisononyl phthalate	DTXSID4022521	28553-12-0			T		T	T	T			T
Diisopropyl sulfate	DTXSID3073131	2973-10-6	2B		T					913	33	
Dimethyl phosphonate	DTXSID5020493	868-85-9			T		T	T		13380	518	T
Dimethyl sulfate	DTXSID5024055	77-78-1	2A	RAHC	T	Both	T	T		42200	5606	T
Dimethylarsinic acid	DTXSID7020508	75-60-5	2B		T		T		T	14388	2445	T
Dimethylcarbamoyl chlorid	DTXSID1020512	79-44-7	2A	RAHC	T		T			17225	189	T
Diphenylhydantoin sodium	DTXSID2023775	630-93-3		RAHC	T		T			4185	13876	T
Dipropyl 2,5-pyridinedica	DTXSID8032544	136-45-8			T					1369	32	
Diuron	DTXSID0020446	330-54-1			T		T	T	T	71096	9483	T
Doxorubicin	DTXSID8021480	23214-92-8	2A	RAHC						236403	122238	T
Elmiron	DTXSID1025229	37319-17-8	2B									
Epichlorohydrin	DTXSID1020566	106-89-8	2A	RAHC	T		T	T	T	36160	6363	T
Eritonite	DTXSID40872384	12510-42-8	1	Kwn	T							
Estragole	DTXSID0020575	140-67-0			T		T	T	T	11012	2070	T
Estrone	DTXSID4022367	53-16-7		Kwn	T		T		T	47750	20816	T
Estropipate	DTXSID3023005	7280-37-7			T				T	7021	117	T
Ethanol	DTXSID9020584	64-17-5	1				T	T	T	1198010	541282	T
Ethoprop	DTXSID4032611	13194-48-4			T				T	23499	775	T
Ethyl acrylate	DTXSID4020583	140-88-5	2B		T		T	T	T	43681	2353	T
Ethyl methanesulfonate	DTXSID6025309	62-50-0	2B	RAHC	T		T		T	25950	6466	T
Ethylbenzene	DTXSID3020596	100-41-4	2B		T		T	T	T	71444	13414	T
Ethylene oxide	DTXSID0020600	75-21-8	1	Kwn	T		T	T	T	146849	18049	T
Ethylene thiourea	DTXSID5020601	96-45-7		RAHC	T		T	T	T	18924	1091	T
Ethyleneimine	DTXSID8020599	151-56-4	2B		T		T			37074	5558	T
Etoposide	DTXSID5023035	33419-42-0	1		T					164082	50135	T
Etridiazole	DTXSID3032547	2593-15-9			T				T	29386	189	T
Fenoxycarb	DTXSID7032393	72490-01-8			T				T	28211	537	T
Fluoro-edenite fibrous am	DTXSID401335773	67164-60-7			T							T
Folpet	DTXSID0021385	133-07-3			T		T	T	T	38543	677	T
Formaldehyde	DTXSID7020637	50-00-0	1	Kwn	T		T	T	T	219873	141838	T

Return to: Section 2.1.1. Continued, next page...

Table E1: Complete list of candidate carcinogens

Chemical Name	DTXSID	CAS	IARC	ROC	Prop65	SVHC	TSCA Active	CDR 2020	MMDBV1	Pat. Count	Lit. Count	Ubiq. Flag
Formaldehyde, polymer wit	DTXSID801011036	25214-70-4				Both		T				T
Formic acid 2-[4-(5-nitro	DTXSID6020640	3570-75-0	2B		T					1031	41	
Fumonisin B1	DTXSID6020644	116355-83-0	2B		T				T	4	1602	T
Furaltadone	DTXSID4048016	139-91-3			T					1552	125	
Furan	DTXSID6020646	110-00-9	2B	RAHC	T	Both	T	T	T	65498	17619	T
Furazolidone	DTXSID4041997	67-45-8			T		T			7273	1361	T
Furfuryl alcohol	DTXSID2025347	98-00-0	2B		T		T	T	T	55638	4841	T
Furilazole	DTXSID4041999	121776-33-8			T					13302	8	T
Furmecyclox	DTXSID2024119	60568-05-0			T				T	11900	31	T
Fusarin C	DTXSID501020204	79748-81-5			T					79	70	
Gallium arsenide	DTXSID2023779	1303-00-0			T		T			29274	8722	T
Ganciclovir	DTXSID8041032	82410-32-0			T					86110	10479	T
Gasoline	DTXSID50105296	86290-81-5	2B					T	T			T
Gemfibrozil	DTXSID0020652	25812-30-0			T				T	44518	5564	T
Gentian Violet	DTXSID5020653	548-62-9	2B		T	Both	T			68384	28454	T
Ginkgo biloba extract	DTXSID8031366	90045-36-6	2B									
Glass wool	DTXSID30891701	308066-92-4		RAHC	T							
Glu-P-1	DTXSID5020657	67730-11-4	2B		T				T	187	168	T
Glu-P-2	DTXSID0020658	67730-10-3	2B		T				T	190	87	T
Glycidaldehyde	DTXSID9020665	765-34-4	2B		T					2755	253	T
Glycidol	DTXSID4020666	556-52-5	2A	RAHC	T		T			69799	1444	T
Glycidyl methacrylate	DTXSID0025361	106-91-2	2A		T		T	T		89538	3057	T
Glyphosate	DTXSID1024122	1071-83-6	2A		T				T	79067	10668	T
Griseofulvin	DTXSID8020674	126-07-8	2B		T				T	44062	7553	T
HC Blue 1	DTXSID6020191	2784-94-3	2B		T		T			730	32	T
Heptachlor	DTXSID3020679	76-44-8	2B		T				T	18672	3856	T
Heptachlor epoxide B	DTXSID1024126	1024-57-3			T				T		149	T
Hexabromobiphenyl	DTXSID3025382	36355-01-8		RAHC								
Hexachloro-1,3-butadiene	DTXSID7020683	87-68-3			T		T		T	8799	1555	T
Hexachlorobenzene	DTXSID2020682	118-74-1	2B	RAHC	T		T	T	T	36868	7342	T
Hexachlorodibenzo-p-dioxi	DTXSID00872820	34465-46- 8			T							T
Hexachloroethane	DTXSID7020689	67-72-1	2B	RAHC	T		T	T	T	20890	1223	T
Hexamethylphosphoramide	DTXSID6020694	680-31-9	2B	RAHC	T		T		T	59098	3707	T
Hydrazine	DTXSID3020702	302-01-2	2A	RAHC	T	Both	T	T	T	109597	24430	T
Hydrazine sulfate	DTXSID8020703	10034-93-2		RAHC	T		T			10655	3530	T
Hydrochlorothiazide	DTXSID2020713	58-93-5	2B				T		T	63505	18228	T
Imazali	DTXSID8024151	35554-44-0			T				T	42807	1029	T

Return to: Section 2.1.1. Continued, next page...

Table E1: Complete list of candidate carcinogens

Chemical Name	DTXSID	CAS	IARC	ROC	Prop65	SVHC	TSCA Active	CDR 2020	MMDBV1	Pat. Count	Lit. Count	Ubiq. Flag
Indeno[1,2,3-cd]pyrene	DTXSID8024153	193-39-5	2B	RAHC	T		T		T	942	2224	T
Indium phosphide	DTXSID3031444	22398-80-7	2A		T		T					T
Indium tin oxide	DTXSID10904045	71243-84-0			T		T			625	977	T
Iprovalicarb	DTXSID0047662	140923-17-7			T					19905	34	T
Iron dextran	DTXSID1031464	9004-66-4	2B	RAHC	T							
Isobutyl nitrite	DTXSID3020750	542-56-3	2B		T		T			2120	213	T
Isoprene	DTXSID2020761	78-79-5	2B	RAHC	T		T	T		51421	9824	T
Isoptirazam	DTXSID7058208	881685-58-1			T					13597	78	T
Isoxafutole	DTXSID5034723	141112-29-0			T				T	23480	204	T
Kava kava resin	DTXSID5031505	9000-38-8	2B									
Kepone	DTXSID1020770	143-50-0	2B	RAHC	T				T	6446	1598	T
Kresoxim-methyl	DTXSID2032558	143390-89-0			T				T	17414	451	T
Lactofen	DTXSID7024160	77501-63-4			T				T	25006	159	T
Lasiocarpine	DTXSID1020772	303-34-4	2B		T					312	57	
Lead	DTXSID2024161	7439-92-1	2B	RAHC	T		T	T	T	8205	39266	T
Lead acetate	DTXSID10274073	15347-57-6		RAHC								
Lead arsenate	DTXSID4042092	7784-40-9		Kwn		Both	T			5668	61	T
Lead phosphate (3:2)	DTXSID5064706	7446-27-7		RAHC	T		T			261	52	T
Lead(II) acetate	DTXSID6020773	301-04-2			T		T	T		35	1379	T
Lead(II) arsenate (3:2)	DTXSID9074274	3687-31-8				Both				4	0	
Lead(II) chromate	DTXSID1064792	7758-97-6				Both	T	T		42242	93	T
Leucomalachite green	DTXSID7031531	129-73-7	2B		T		T			2848	376	T
Levofuraltadone	DTXSID6043834	3795-88-8	2B								20	
Lindane	DTXSID2020686	58-89-9	1	RAHC	T		T		T	104118	10426	T
Lomustine	DTXSID2023222	13010-47-4	2A	RAHC	T					92047	497	T
Lynestrenol	DTXSID4021478	52-76-6			T					5706	1407	T
MON-4660	DTXSID0044575	71526-07-3			T					5721	22	T
Malathion	DTXSID4020791	121-75-5	2A		T				T	52709	9549	T
Malonaldehyde, sodium sal	DTXSID4020793	24382-04-5			T					2	0	
Mancozeb	DTXSID0034695	8018-01-7			T							
Maneb	DTXSID9020794	12427-38-2			T					35471	617	T
MeA-alpha-C	DTXSID5043766	68006-83-7	2B		T				T	112	103	T
MelQ	DTXSID6020800	77094-11-2	2B	RAHC	T					112	320	T
MelQx	DTXSID1020801	77500-04-0	2B	RAHC	T					179	631	T
Medroxyprogesterone aceta	DTXSID0025527	71-58-9	2B		T		T			54459	11356	T
Megestrol acetate	DTXSID9040683	595-33-5			T					92294	3245	T
Melamine	DTXSID6020802	108-78-1	2B				T	T		61063	5959	T

Return to: Section 2.1.1. Continued, next page...

Table E1: Complete list of candidate carcinogens

Chemical Name	DTXSID	CAS	IARC	ROC	Prop65	SVHC	TSCA Active	CDR 2020	MMDBV1	Pat. Count	Lit. Count	Ubiqu. Flag
Melphalan	DTXSID6020804	148-82-3	1	Kwn	T					128050	8336	T
Mepanipyrim	DTXSID4042121	110235-47-7			T				T	32301	147	T
Merphalan	DTXSID9031569	531-76-0	2B		T					1429	8236	T
Mestranol	DTXSID0020814	72-33-3		Kwn	T				T	13022	3539	T
Metam-sodium	DTXSID2029167	137-42-8			T				T	18989	506	T
Methyl acrylate	DTXSID0024183	96-33-3	2B		T		T	T	T	43077	5211	T
Methyl carbamate	DTXSID8020834	598-55-0			T		T	T	T	22180	825	T
Methyl iodide	DTXSID0024187	74-88-4			T		T	T	T	22793	12485	T
Methyl mercury	DTXSID2031615	16056-34-1			T				T	3187	14	T
Methyl methanesulfonate	DTXSID7020845	66-27-3	2A	RAHC	T		T		T	23115	3150	T
Methylarsonic acid	DTXSID4020903	124-58-3	2B						T	4544	1517	T
Methylazoxymethanol	DTXSID1035076	590-96-5			T					123	527	T
Methylazoxymethanol aceta	DTXSID1025568	592-62-1	2B		T						663	T
Methyl Eugenol	DTXSID5025607	93-15-2	2B	RAHC	T		T		T	14589	1767	T
Methylhydrazine	DTXSID4020874	60-34-4			T		T		T	45729	1341	T
Methylnitrosoguanidi	DTXSID2020846	70-25-7	2A	RAHC	T		T		T	17230	3906	T
Metiram	DTXSID9034737	9006-42-2			T				T			T
Metronidazole	DTXSID2020892	443-48-1	2B	RAHC	T					117361	39734	T
Microcystin LR	DTXSID3031654	101043-37-2	2B		T				T	1670	0	T
Mineral oil - includes pa	DTXSID3034743	8012-95-1		Kwn			T	T	T			T
Mirex	DTXSID7020895	2385-85-5	2B	RAHC	T				T	10508	2169	T
Mitomycin C	DTXSID2020898	50-07-7	2B		T		T			172352	33326	T
Mitoxantrone	DTXSID4046947	65271-80-9	2B							122629	14317	T
Mitoxantrone dihydrochlor	DTXSID0045173	70476-82-3			T					79132	4572	T
Molybdenum trioxide	DTXSID7020899	1313-27-5	2B		T		T	T		50745	8876	T
Monobasic lead acetate	DTXSID1020774	1335-32-6		RAHC	T		T					T
Monocrotaline	DTXSID9020902	315-22-0	2B		T				T	3035	3163	T
Myrcene	DTXSID6025692	123-35-3	2B		T		T	T		27601	4553	T
N'-Nitrosornicotine	DTXSID4021476	16543-55-8		RAHC	T				T	1602	505	T
N,N'-Bis(2-chloroethyl)-N	DTXSID8022743	154-93-8	2A	RAHC	T					108198	11831	T
N,N'-Diacylbenzidine	DTXSID9036854	613-35-4	2B		T		T			105	73	T
N,N,4-Trimethylaniline	DTXSID0021832	99-97-8	2B		T		T	T		28377	687	T
N,N-Dimethyl-N'-[5-[2-(5-	DTXSID801126079	25962-77-0	2B									
N,N-Dimethylacetamide	DTXSID5020499	127-19-5	2B		T		T	T		53671	9164	T
N,N-Dimethylformamide	DTXSID6020515	68-12-2	2A		T		T	T		140231	60584	T
N-(2-Cyanoethyl)-N-methyl	DTXSID70975579	60153-49-3	2B		T					33	31	
N-Amyl-N-methylnitrosamin	DTXSID30157617	13256-07-0			T					17	91	

Return to: Section 2.1.1. Continued, next page...

Table E1: Complete list of candidate carcinogens

Chemical Name	DTXSID	CAS	IARC	ROC	Prop65	SVHC	TSCA Active	CDR 2020	MMDBV1	Pat. Count	Lit. Count	Ubiq. Flag
N-Heptyl-N-methylnitrous	DTXSID70936822	16338-99-1			T						4	
N-Hexyl-N-methylnitrous a	DTXSID20951227	28538-70-7			T					1	1	
N-Methyl-N-nonylnitrous a	DTXSID80997267	75881-19-5			T						1	
N-Methyl-N-octylnitrous a	DTXSID50955970	34423-54-6			T						1	
N-Methylolacrylamide	DTXSID3020885	924-42-5			T	EU		T		97163	321	T
N-Nitroso-N-methyl-N-dode	DTXSID4021000	55090-44-3			T					298	25	
N-Nitroso-N-methyl-N-tetr	DTXSID4021004	75881-20-8			T						6	
N-Nitroso-N-methyldecylam	DTXSID9021005	75881-22-0			T					1	2	
N-Nitroso-N-methylethylam	DTXSID6021036	10595-95-6	2B		T				T	204	67	T
N-Nitroso-N-methylurea	DTXSID4021006	684-93-5	2A	RAHC			T			7905	6496	T
N-Nitroso-N-methylurethan	DTXSID4021472	615-53-2	2B		T		T			679	394	T
N-Nitrosodi-n-propylamine	DTXSID6021032	621-64-7	2B	RAHC	T		T		T	1276	333	T
N-Nitrosodibutylamine	DTXSID2021026	924-16-3	2B	RAHC	T		T		T	644	446	T
N-Nitrosodiethanolamine	DTXSID7021027	1116-54-7	2B	RAHC	T		T			542	351	T
N-Nitrosodiethylamine	DTXSID2021028	55-18-5	2A	RAHC	T		T		T	2615	7220	T
N-Nitrosodimethylamine	DTXSID7021029	62-75-9	2A	RAHC	T		T		T	7004	6528	T
N-Nitrosodiphenylamine	DTXSID6021030	86-30-6			T		T	T		8944	345	T
N-Nitrosohexamethyleneimi	DTXSID0021040	932-83-2			T					75	40	
N-Nitrosomethyl-N-propyla	DTXSID90238991	924-46-9			T					13	6	
N-Nitrosomethylvinylamine	DTXSID3042211	4549-40-0	2B	RAHC	T						0	
N-Nitrosomorpholine	DTXSID4021056	59-89-2	2B	RAHC	T				T	979	1013	T
N-Nitrosopiperidine	DTXSID8021060	100-75-4	2B	RAHC	T		T		T	961	511	T
N-Nitrosopyrrolidine	DTXSID8021062	930-55-2	2B	RAHC	T		T		T	896	641	T
N-Nitrososarcosine	DTXSID9074309	13256-22-9	2B	RAHC	T					350	120	
N-[4-(5-Nitro-2-furyl)-2-	DTXSID9020952	531-82-8	2B		T					349	44	
Nafenopin	DTXSID8020911	3771-19-5	2B		T					577	583	T
Nalidixic acid	DTXSID3020912	389-08-2			T					37656	12281	T
Naphthalene	DTXSID8020913	91-20-3	2B	RAHC	T		T	T		78694	34015	T
Nickel	DTXSID2020925	7440-02-0	2B	RAHC	T		T	T	T	591386	26453	T
Nickel (II) oxide	DTXSID7025710	1313-99-1		Kwn	T		T				649	T
Nickel carbonate	DTXSID3025708	3333-67-3			T		T					T
Nickel carbonyl	DTXSID0024212	13463-39-3			T		T		T	6	1446	T
Nickel hydroxide (Ni(OH)2	DTXSID90274011	12054-48-7		Kwn	T		T	T		37378	106	T
Nickel subsulfide	DTXSID2025711	12035-72-2		Kwn	T		T	T				T
Nickel sulfate	DTXSID6023787	7786-81-4		Kwn			T	T		72521	6057	T
Nickel sulfide	DTXSID801014466	11113-75-0		Kwn			T	T				T
Nickel(II) acetate	DTXSID7020926	373-02-4		Kwn	T		T	T		32350	49	T

Return to: Section 2.1.1. Continued, next page...

Table E1: Complete list of candidate carcinogens

Chemical Name	DTXSID	CAS	IARC	ROC	Prop65	SVHC	TSCA Active	CDR 2020	MMDBV1	Pat. Count	Lit. Count	Ubiqu. Flag
Nickelocene	DTXSID8025709	1271-28-9			T		T				287	T
Nifuradene	DTXSID2020973	555-84-0	2B		T					1299	25	
Niridazole	DTXSID6045244	61-57-4	2B		T					3178	788	T
Nitrapyrin	DTXSID0024216	1929-82-4			T		T	T		14054	768	T
Nitrioltriactic acid	DTXSID6020939	139-13-9	2B	RAHC	T		T	T	T	121944	4905	T
Nitrioltriactic acid tri	DTXSID5020940	18662-53-8			T					977	36	
Nitrobenzene	DTXSID3020964	98-95-3	2B	RAHC	T		T	T	T	117874	16032	T
Nitrofen	DTXSID7020970	1836-75-5	2B	RAHC	T		T		T	12704	967	T
Nitrofurazone	DTXSID5020944	59-87-0			T		T	T		3908	252	T
Nitrogen mustard	DTXSID2020975	51-75-2	2A		T		T			102630	7997	T
Nitrogen mustard N-oxide	DTXSID7020976	126-85-2	2B		T					4268	5672	T
Nitrogen mustard hydrochl	DTXSID8025757	55-86-7		RAHC	T					22951	5809	T
Nitrogen mustard oxide	DTXSID0023375	302-70-5			T					24324	5662	T
Nitromethane	DTXSID2020977	75-52-5	2B	RAHC	T		T	T	T	50287	7517	T
Nitroso-2,6-dimethylmorph	DTXSID4021470	1456-28-6			T					18	88	
Nitrosomethyl-N-butylamin	DTXSID20220960	7068-83-9			T					452	25	
Nitrosomethylundecylamine	DTXSID9021055	68107-26-6			T						3	
Norethindrone	DTXSID9023380	68-22-4		RAHC	T				T	43524	8369	T
Norethynodrel	DTXSID3021069	68-23-5			T					9141	1067	T
Ochratoxin A	DTXSID7021073	303-47-9	2B	RAHC	T					2582	6496	T
Oryzalin	DTXSID8024238	19044-88-3			T				T	23015	1280	T
Oxadiazon	DTXSID3024239	19666-30-9			T				T	27954	356	T
Oxazepam	DTXSID1021087	604-75-1	2B		T				T	31498	5470	T
Oxymetholone	DTXSID5023794	434-07-1		RAHC	T					5877	943	T
Parathion	DTXSID7021100	56-38-2	2B		T				T	44073	7950	T
Pentachlorophenol	DTXSID7021106	87-86-5	1	RAHC	T		T	T	T	74940	8693	T
Pentaerythritol dibromide	DTXSID9020164	3296-90-0	2B	RAHC	T		T	T		1062	74	T
Pentazinc chromate octahy	DTXSID30197957	49663-84-5					EU			2	0	
Pentosan polysulfate sodi	DTXSID70102623	140207-93-8			T							T
Perfluorooctanesulfonic a	DTXSID3031864	1763-23-1			T		T		T	26669	3356	T
Perfluorooctanoic acid	DTXSID8031865	335-67-1	2B		T		T		T	23464	4336	T
Phenacetin	DTXSID1021116	62-44-2	1	RAHC	T		T	T		29377	7087	T
Phenazopyridine	DTXSID1023445	94-78-0			T				T	20885	902	T
Phenazopyridine hydrochlo	DTXSID1021118	136-40-3	2B	RAHC	T					7530	765	T
Phenesterin	DTXSID6021119	3546-10-9			T					23597	35	T
Phenobarbital	DTXSID5021122	50-06-6	2B		T				T	147350	41672	T
Phenolphthalein	DTXSID0021125	77-09-8	2B	RAHC	T		Both	T		86632	16225	T

Return to: Section 2.1.1. Continued, next page...

Table E1: Complete list of candidate carcinogens

Chemical Name	DTXSID	CAS	IARC	ROC	Prop65	SVHC	TSCA Active	CDR 2020	MMDBV1	Pat. Count	Lit. Count	Ubiqu. Flag
Phenoxybenzamine	DTXSID0023458	59-96-1								13428	9404	T
Phenoxybenzamine hydrochl	DTXSID0021127	63-92-3	2B	RAHC	T					1788	5402	T
Phenyl glycidyl ether	DTXSID8021145	122-60-1	2B		T		T	T		51345	723	T
Phenylhydrazine	DTXSID8021147	100-63-0			T		T			52396	7854	T
Pioglitazone	DTXSID3037129	111025-46-8	2A		T				T	74776	4456	T
Pyrimicarb	DTXSID1032569	23103-98-2			T			T		32192	957	T
Pitch, coal tar, high-tem	DTXSID1028285	65996-93-2	1	Kwn	T	Both						T
Plutonium	DTXSID4064682	7440-07-5	1		T					31171	3680	T
Polybrominated biphenyls	DTXSID5021174	59536-65-1	2A	RAHC	T				T			T
Polychlorinated biphenyls	DTXSID5024267	1336-36-3	1	RAHC	T		T					T
Polycyclic aromatic hydro	DTXSID3044043	130498-29-2		RAHC				T				T
Ponceau 3R	DTXSID7021231	3564-09-8	2B		T					258	32	
Potassium N-methyldithioc	DTXSID1034424	137-41-7			T				T	3958	149	T
Potassium arsenite anhydr	DTXSID10274159	13464-35-2		Kwn						4	82	
Potassium bromate	DTXSID6020195	7758-01-2	2B		T		T			11701	2523	T
Potassium chromate(VI)	DTXSID8064858	7789-00-6		Kwn		Both	T			10310	2536	T
Potassium dichromate	DTXSID5025948	7778-50-9		Kwn		Both	T			40647	15954	T
Premarin	DTXSID9021186	12126-59-9		Kwn								
Primidone	DTXSID7023510	125-33-7	2B		T				T	21937	5165	T
Procabazine	DTXSID4021189	671-16-9		RAHC	T					133053	8706	T
Procabazine hydrochlorid	DTXSID3021190	366-70-1	2A	RAHC	T					24535	3444	T
Procainidone	DTXSID9033923	32809-16-8			T				T	19068	755	T
Progesterone	DTXSID3022370	57-83-0		RAHC	T		T		T	115933	123256	T
Propachlor	DTXSID4024274	1918-16-7			T				T	29406	450	T
Propargite	DTXSID4024276	2312-35-8			T				T	29783	466	T
Propoxur	DTXSID7021948	114-26-1			T				T	39359	2022	T
Propyleneimine	DTXSID8024286	75-55-8	2B	RAHC	T		T	T		10917	169	T
Propyzamide	DTXSID2020420	23950-58-5			T				T	21198	577	T
Pymetrozine	DTXSID2032637	123312-89-0			T				T	12739	353	T
Pyridine	DTXSID9021924	110-86-1	2B		T		T	T	T	178813	80334	T
Quartz (SiO2)	DTXSID3032040	14808-60-7	1	Kwn	T			T				T
Quinoline	DTXSID1021798	91-22-5	2B		T		T	T		235320	15414	T
R-(+)-Pulegone	DTXSID2025975	89-82-7	2B		T		T			8069	505	T
Radium-224	DTXSID50912341	13233-32-4	1						T	503	159	T
Radium-226	DTXSID3051202	13982-63-3	1						T	1904	165	T
Radium-228	DTXSID6050415	15262-20-1	1						T	249	43	T
Radon	DTXSID701015778	10043-92-2	1	Kwn					T	42893	6555	T

Return to: Section 2.1.1. Continued, next page...

Table E1: Complete list of candidate carcinogens

Chemical Name	DTXSID	CAS	IARC	ROC	Prop65	SVHC	TSCA Active	CDR 2020	MMDBV1	Pat. Count	Lit. Count	Ubic. Flag
Refractories, fibers, alu	DTXSID90891702	142844-00-6	2B		T	Both		T				T
Reserpine	DTXSID7021237	50-55-5		RAHC	T		T			201285	28228	T
Resmethrin	DTXSID7022253	10453-86-8			T				T	14870	427	T
Rhodamine B	DTXSID6042369	81-88-9			T		T			181293	13925	T
Riddelline	DTXSID4026006	23246-96-0	2B	RAHC	T					4	72	T
Riebeckite	DTXSID5030743	12001-28-4		Kwn					T			T
Safrrole	DTXSID0001254	94-59-7	2B	RAHC	T		T			18387	1956	T
Sedaxane	DTXSID2058207	874967-67-6			T				T	14302	60	T
Selenium sulfide (SeS)	DTXSID9021265	7446-34-6		RAHC	T					10751	742	T
Sennustine	DTXSID8031603	13909-09-6	1	Kwn	T					41318	1004	T
Shale oils	DTXSID7028435	68308-34-9	1		T		T					T
Silicon carbide	DTXSID5052751	409-21-2	2A		T		T	T		170678	7975	T
Silicon carbide, fibrous	DTXSID701020379	308076-74-6	2B									
Sodium 2-phenylphenate	DTXSID2021153	132-27-4	2B		T		T			8929	357	T
Sodium arsenate	DTXSID5020102	7631-89-2		Kwn			T					T
Sodium arsenite	DTXSID5020104	7784-46-5		Kwn			T			18323	3161	T
Sodium chromate	DTXSID7032056	7775-11-3		Kwn		Both	T			10080	1441	T
Sodium dichromate	DTXSID8021274	10588-01-9		Kwn		Both	T	T		20057	1021	T
Sodium pentachlorophenate	DTXSID4040264	131-52-2		RAHC			T			11099	2350	T
Spirocliflofen	DTXSID6034928	148477-71-8			T				T	24470	195	T
Spironolactone	DTXSID6034186	52-01-7			T					60269	21554	T
Stanozolol	DTXSID3044128	10418-03-8			T					8003	1550	T
Sterigmatocystin	DTXSID2021280	10048-13-2	2B		T					506	1234	T
Streptozotocin	DTXSID2021282	18883-66-4	2B	RAHC	T					96389	19175	T
Strontium chromate	DTXSID70872832	7789-06-2		Kwn		Both	T	T		10850	239	T
Styrene	DTXSID2021284	100-42-5	2A	RAHC	T		T	T	T	309037	38045	T
Styrene oxide	DTXSID2021286	96-09-3	2A	RAHC	T		T			73762	2422	T
Sulfallate	DTXSID7021289	95-06-7	2B	RAHC	T				T	9381	96	T
Sulfasalazine	DTXSID0001256	599-79-1	2B		T					98963	8438	T
Sulfuric acid	DTXSID5029683	7664-93-9		Kwn			T	T	T	334467	122647	T
Talc	DTXSID5032109	14807-96-6	2B		T		T	T	T			T
Tamoxifen	DTXSID1034187	10540-29-1	1	Kwn	T				T	127664	47356	T
Teniposide	DTXSID8023638	29767-20-2	2A							93396	3043	T
Teriparatide	DTXSID001027900	52232-67-4			T							
Testosterone	DTXSID8022371	58-22-0			T		T		T	94121	135724	T
Tetrachloroethylene	DTXSID2021319	127-18-4	2A	RAHC	T		T	T	T	56538	7992	T
Tetraethyl lead	DTXSID7023801	78-00-2		RAHC			T	T	T	11519	1832	T

Return to: Section 2.1.1. Continued, next page...

Table E1: Complete list of candidate carcinogens

Chemical Name	DTXSID	CAS	IARC	ROC	Prop65	SVHC	TSCA Active	CDR 2020	MMDBV1	Pat. Count	Lit. Count	Ubiq. Flag
Tetrafluoroethylene	DTXSID6021325	116-14-3	2A	RAHC	T		T	T		48648	2986	T
Tetrahydrofuran	DTXSID1021328	109-99-9	2B		T		T	T	T	198206	46843	T
Tetranitromethane	DTXSID5021334	509-14-8	2B	RAHC	T		T			10594	1480	T
Thioacetamide	DTXSID9021340	62-55-5	2B	RAHC	T		T			18385	5414	T
Thiodi carb	DTXSID0032578	59669-26-0			T				T	142	113	T
Thiotepea	DTXSID0021339	52-24-4	1	Kwn	T					132998	7436	T
Thiouracil	DTXSID4021347	141-90-2	2B		T		T			47267	4128	T
Thiourea	DTXSID9021348	62-56-6		RAHC	T		T	T		36477	29607	T
Thorium	DTXSID6049800	7440-29-1	1				T		T	63012	147	T
Thorium(IV) oxide	DTXSID0051657	1314-20-1		Kwn	T		T			32451	3427	T
Titanium dioxide	DTXSID3021352	13463-67-7	2B		T		T	T				T
Toluene 2,4-diisocyanate	DTXSID7026156	584-84-9		RAHC			T	T	T	100258	3527	T
Toluene diisocyanate	DTXSID0024341	26471-62-5	2B	RAHC	T		T	T				T
Toluene-2,6-diisocyanate	DTXSID2026157	91-08-7		RAHC			T	T		57372	286	T
Toxaphene	DTXSID7021368	8001-35-2	2B	RAHC	T				T			T
Tremolite asbestos	DTXSID701026177	77536-68-6		Kwn						11539	148	T
Treosulfan	DTXSID0026173	299-75-2	1		T				T	33615	4021	T
Triantereene	DTXSID6021373	396-01-0	2B		T					1914	163	T
Tribromoacetic acid	DTXSID6021668	75-96-7		RAHC			T					T
Tribromoneopentyl alcohol	DTXSID40893054	36483-57-5				EU	T	T				T
Tribufos	DTXSID1024174	78-48-8			T		T	T	T	7277	446	T
Trichloromethine	DTXSID6074835	817-09-4	2B		T					215	78	
Trichloroacetic acid	DTXSID1021378	76-03-9	2B		T		T		T	8644	42054	T
Trichloroethylene	DTXSID0021383	79-01-6	1	Kwn	T	Both	T	T	T	56309	17227	T
Tridymite	DTXSID6049563	15468-32-3		Kwn		Both				149	3	
Triethyl arsenate	DTXSID20166036	15606-95-8										
Trim VX	DTXSID701052685	1204332-00-2			T							T
Trimethyl orthophosphate	DTXSID1021403	512-56-1			T		T	T		67999	913	T
Trimethylolpropane triacr	DTXSID0027773	15625-89-5	2B		T		T	T		45319	605	T
Triphenyltin hydroxide	DTXSID1021409	76-87-9			T		T	T		21418	335	T
Tris(1,3-dichloro-2-propyl	DTXSID9026261	13674-87-8			T		T	T	T	4732	325	T
Tris(2,3-dibromopropyl) p	DTXSID5021413	126-72-7	2A	RAHC	T		T		T	11314	369	T
Tris(2-chloroethyl) phosph	DTXSID5021411	115-96-8			T		T		T	19483	510	T
Trip-P-1	DTXSID0043767	62450-06-0	2B		T				T	483	300	T
Trip-P-2	DTXSID5043768	62450-07-1	2B		T				T	703	331	T
Uracil mustard	DTXSID8026270	66-75-1	2B		T					65211	251	T
Urethane	DTXSID9021427	51-79-6	2A	RAHC	T		T			173756	25913	T

Return to: Section 2.1.1. Continued, next page...

Table E1: Complete list of candidate carcinogens

Chemical Name	DTXSID	CAS	IARC	ROC	Prop65	SVHC	TSCA Active	CDR 2020	MMDBV1	Pat. Count	Lit. Count	Ubiq. Flag
Vanadium oxide (V2O5)	DTXSID2023806	1314-62-1	2B		T		T	T				T
Vinclozolin	DTXSID4022361	50471-44-8			T				T	39037	1394	T
Vincristine-prednisone-ni	DTXSID701045240	113803-47-7			T							
Vinyl acetate	DTXSID3021431	108-05-4	2B				T	T	T	107263	7107	T
Vinyl bromide	DTXSID8021432	593-60-2	2A	RAHC	T		T	T	T	38132	852	T
Vinyl chloride	DTXSID8021434	75-01-4	1	Kwn	T		T	T	T	96428	10265	T
Vinyl fluoride	DTXSID3021435	75-02-5	2A	RAHC	T		T	T		128091	1033	T
Z-Tetrachlorvinphos	DTXSID1032648	22248-79-9	2B		T				T	30515	144	T
Zileuton	DTXSID9023752	111406-87-2			T					17395	1671	T
Zinc chromate	DTXSID7047485	13530-65-9		Kwn			T	T		15302	544	T
Zinc potassium chromate	DTXSID2026313	11103-86-9				Both	T	T			20	T
alpha,alpha-bis[4-(Dimeth	DTXSID4064474	6786-83-0				Both	T			18	0	T
alpha-1,2,3,4,5,6-Hexachl	DTXSID2020684	319-84-6		RAHC			T		T	104118	10426	T
alpha-Methylstyrene	DTXSID9025661	98-83-9	2B		T		T	T		38793	1821	T
beta-Butyrolactone	DTXSID1020223	3068-88-0	2B		T		T			15557	201	T
beta-Hexachlorocyclohexan	DTXSID7020685	319-85-7		RAHC			T		T	104118	10426	T
beta-Propiolactone	DTXSID8021197	57-57-8	2B	RAHC	T		T			49583	1530	T
bis(2-Chloro-1-methylethy	DTXSID4020167	108-60-1			T		T		T	3221	84	T
bis(Chloromethyl) ether	DTXSID8020173	542-88-1		Kwn	T		T		T	3468	369	T
o-Aminoazotoluene	DTXSID1020069	97-56-3	2B	RAHC	T	Both	T			1341	286	T
p,p'-DDD	DTXSID4020373	72-54-8			T				T	1351	1885	T
p,p'-DDE	DTXSID9020374	72-55-9			T				T	955	5481	T
p-Nitroanisole	DTXSID9059208	100-17-4	2B		T		T			3860	793	T
trans-2-[(Dimethylamino)m	DTXSID7020502	55738-54-0			T						0	

Return to: Section 2.1.1.