

MATERIALS TRANSFER AGREEMENT

Provider: US Environmental Protection Agency

Provider Contact (not signator)

Name: Antony J. Williams

Address: 109 T.W. Alexander Drive, Room D131I, Mail Drop D143-02, Research Triangle Park, NC 27711

Phone: 919-541-1033

Email: williams.antony@epa.gov

Recipient: US Geological Survey

Recipient Contact (not signator)

Name: Gabrielle Black

Address: 6000 J St, Placer Hall 5001C

Phone: 916-278-9090

Email: gpblack@usgs.gov

1. Provider and Recipient are authorized to perform technical assistance with other Federal agencies, units of State or local government, industrial organizations, private corporations, public and private foundations, and nonprofit organizations (including universities) under the Stevenson-Wydler Act (15 U.S.C. §3710a, as amended).

2. Provider agrees to transfer to Recipient the following Research Material:

ChemSpace Tool, a proof-of-concept chemical space mapping tool to help address challenges in defining the detectable and identifiable chemical space of nontargeted analyses, which uses existing chemometric tools as well as filtering based on categories of chemicals identified in different types of samples. The software will be delivered in the form of an R-shiny application and containerized so that it can be run inside the recipient's environment.

3. This Research Material may not be used in human subjects. The Research Material will be used only for research purposes by Recipient's investigator in his/her Center, for the research project described below, under suitable containment conditions. This Research Material will not be used for screening, production or sale, for which a commercialization license may be required. Recipient agrees to comply with all Federal rules and regulations applicable to the Research Project and the handling of the Research Material.

EPA ONLY: If the data or material that are being transferred constitute human subjects research, please visit the following intranet site to determine if your project needs review and approval by the HSRRO: <https://intranet.ord.epa.gov/human-subject-research/hsr-projects-review>

There is no Human Subjects material being used in this research.

Research Plan will be reviewed and approved by HSRRO. No work will commence until HSRRO approval has been obtained.

4. If the data or material that are being transferred involve life sciences research, or more specifically any of the select agents or toxins listed and/or the definitions provided in EPA Order 1000.19 *Policy and Procedures for Managing Dual Use Research of Concern*, then Principal Investigators should consult EPA's Institutional Contact for Dual Use Research of Concern (ICDUR) at DURC@epa.gov before completing the following section. If not, then check the first box below.

This research does not meet any of the definitions of Dual Use Research of Concern (DURC) and no additional review or oversight are required. The PI must report to the ICDUR any results or changes in the research that meet any of the definitions of DURC.

This research meets one or more definitions of DURC and requires additional oversight under the *USG Policy for Institutional Oversight of DURC*. The parties to this Agreement are required to comply with EPA Order 1000.19, *Policy and Procedures for Managing Dual Use Research of Concern*.

For information about DURC and EPA Order 1000.19, please visit:

<http://intranet.ord.epa.gov/homeland-security/dual-use-research-concern-durc-policies>

5. This Research Material will be used by Recipient's investigator solely in connection with the following research project ("Research Project") described with specificity as follows (*insert description here or use an attachment page if necessary*):

Recipient will use the Research Material in the evaluation of contaminants in environmental media, namely, in identifying compounds likely to be identified in nontargeted chemical analysis. The recipient will select options in the Research Material that mimic laboratory method steps potentially including, but not limited to, instrument platform, ionization technique, extraction solvents, solid phase extraction method, elution buffers, analytical columns, mobile phase and mobile phase additives to predict which compounds included in the Research Material database are extractable, detectable, and identifiable using the Recipients methodologies. This predicted subset of chemical compounds will be used to better understand what compounds are and are not within the scope of the Recipients nontargeted chemical analysis research projects. Additionally, the information gleaned from use of the Research Material will help the Recipient calculate accuracy and quality control metrics on their datasets. The Recipient will use the Research Material on a multitude of projects and methods development projects including nontargeted chemical analyses of water, soil, sludges, biota, tissue, and bed sediment.

6. In all oral presentations or written publications concerning the Research Project, RECIPIENT will

acknowledge PROVIDER's contribution of this Research Material unless requested otherwise. RECIPIENT agrees to protect the information claimed as confidential business information from unauthorized disclosure to the extent permitted by law and consistent with EPA's regulations under 40 C.F.R. Part 2, Subpart B. The foregoing shall not apply to information that is or becomes publicly available or which is disclosed to RECIPIENT without a confidentiality obligation. In asserting a claim for protection, the PROVIDER must stamp its Research Material as "CLAIMED AS CONFIDENTIAL BUSINESS INFORMATION." Documents that are stamped with "CLAIMED AS CONFIDENTIAL BUSINESS INFORMATION" represent that the PROVIDER is asserting a confidentiality claim for a period of three (3) years. Any oral disclosures from PROVIDER to RECIPIENT, which PROVIDER wishes to assert as confidential business information, shall be identified as being confidential business information at the time of the disclosure and by written notice, stamped in the manner stated above, and delivered to RECIPIENT within thirty (30) days after the date of the oral disclosure. RECIPIENT may publish or otherwise publicly disclose the results of the Research Project, but if PROVIDER has given claimed confidential business information to RECIPIENT, such public disclosure may be made only after PROVIDER has had thirty (30) days to review the proposed disclosure to determine if it includes any claimed confidential business information, to the extent such review period is permitted by law.

7. This Research Material represents a significant investment on the part of Provider and is considered proprietary to Provider. Recipient's investigator therefore agrees to retain control over this Research Material and further agrees not to transfer the Research Material to other people not under his/her direct supervision without advance written approval of Provider. Provider reserves the right to distribute the Research Material to others and to use it for its own purposes. When the Research Project is completed, the Research Material will be returned to the Provider or disposed, if directed by Provider, to the extent such destruction is permitted by law.

8. This Research Material is provided as a service to the research community. It is being supplied to Recipient with no warranties, express or implied, including any warranty of merchantability or fitness for a particular purpose. Provider makes no representations that the use of the Research Material will not infringe any patent or proprietary rights of third parties.

9. Recipient shall retain title to any patent or other intellectual property rights in inventions made by its employees in the course of the Research Project. However, if said inventions contain any portion of the Research Material, are derived from the Research Material, or could not have been produced but for the use of the Research Material, Recipient agrees to contact the Provider to determine what ownership interests, if any, the Provider may have, and, where applicable, to negotiate in good faith the terms of a commercial license. Inventorship for a patent application or a commercialized product based on said inventions shall be determined according to United States patent law.

10. When Provider is the EPA: Recipient agrees not to claim, infer, or imply endorsement by the Government of the United States of America (hereinafter referred to as "Government") of the Research Project, the institution, or personnel conducting the Research Project or any resulting product(s).

11. When Recipient is the EPA: Provider will not be liable to EPA for any claims or damages arising from EPA's use of the Research Material.

12. The Provider shall have the right to terminate this Agreement at any time if Recipient breaches any of the terms of this Agreement. Upon termination, Recipient shall return to the Provider all unused portions of the Research Materials.

13. Will EPA develop any products or services from information or materials provided by the Recipient?

Yes – go to item A

No – skip to #14 (next clause)

Item A: The EPA Center must coordinate on matters related to Quality Assurance with their QA Specialist.

If necessary, the Center will develop/has developed a Quality Assurance Plan in coordination with the Quality Assurance Specialist.

No QA requirements are needed.

14. All notices pertaining to or required by this Agreement shall be in writing and shall be signed by an authorized representative and shall be delivered by hand (including private courier mail service) or sent by certified mail, return receipt requested, with postage prepaid, addressed as follows:

Provider's Contact Information:

Russell Thomas, Ph.D.

Director of EPA's Center for Computational Toxicology and Exposure

thomas.russell@epa.gov

919-541-4219

Recipient's Contact Information:

Gabrielle Black, Ph.D. Chemist

US Geological Survey

gpblack@usgs.gov

916-278-9090

With a copy to:

Kathleen Graham
FTTA Program Coordinator
Graham.kathleen@epa.gov
(303) 312-6137
FTTA@epa.gov

15. Paragraphs 3, 6, 8, 9, 10, and 11 shall survive termination.

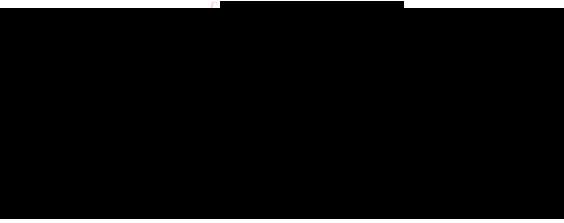
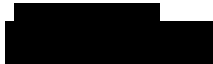
16. This Agreement shall be construed in accordance with law as applied by the Federal courts in the District of Columbia.

17. The undersigned Provider and Recipient expressly certify and affirm that the contents of any statements made herein are truthful and accurate.

18. This agreement shall enter into force as of the date of the last signature of the parties and shall remain in effect for 36 months from said date.

Authorized Representative of Institution SIGNATURES

FOR THE RECIPIENT

By:  _____ Date 


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