Broad-Area Emission Surveys with Airborne Remote Sensing

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Introduction

- Airborne hyperspectral longwave-infrared (LWIR) imaging is a powerful technique for detecting, identifying, and tracking/sourcing gaseous emissions from compact sources
- Hyperspectral resolution enables full characterization of the thermal radiance distribution and detection/identification of gaseous and solid materials within the scene
 - Atmospheric Compensation (ISAC)
 - Spectral Matched Filter (SMF) or Adaptive Coherence Estimation (ACE)
 - Stepwise (forward) Generalized Least Squares (whitened space)
- Urban-industrial environments are replete with gaseous emissions deriving from a wide variety of commercial and residential operations
- Broad-area surveys are important in regulatory monitoring, mitigation of public health concerns, post-disaster hazard evaluation, regional climate assessment
 - Recovery from immediate effects of catastrophic events is often hampered by an array of latent dangers
 - Fugitive emissions of hazardous gases are an especially pernicious risk to first responders and nearby communities
 - Broad-area airborne LWIR surveys rapidly mounted provide flexible and unique capabilities for identification and monitoring of environmental hazards

Airborne Wide-Swath LWIR Spectral Imager ("Mako")



Whiskbroom scanning enables broad-area coverage and high areal rate

Parameter	Mako
Spectral coverage	7.6 – 13.2 μm
Spectral resolution	44 nm (128 channels)
IFOV	0.55 mrad
Along-track FOV	4.0°
Base frame rate	3255 Hz
Operational frame rate	814 Hz (4 co-adds)
Cross-track pixels	400 – 3600
Cross-track field-of-regard (relative to nadir)	±56.4° (max.)
FPA temperature	9.5 K
NESR (10 µm, 4 co-adds)	<0.4 µW cm ⁻² sr ⁻¹ µm ⁻¹
NEDT (10 µm, 300 K)	0.02 K
Scan type	Whiskbroom
Areal acquisition rate (2-m GSD)	33 km²/min (max.)





Spectral Detection and Identification

- Spectral Matched Filter (SMF): $SMF = \frac{\mathbf{A}^{T} \mathbf{C}^{-1} \mathbf{Y}}{(\mathbf{A}^{T} \mathbf{C}^{-1} \mathbf{A})^{1/2}}$
- Adaptive Coherence Estimator (ACE): $A^{T}C^{-1}Y$

ACE = $\frac{\mathbf{A}^{\mathrm{T}} \mathbf{C}^{-1} \mathbf{Y}}{\left(\mathbf{A}^{\mathrm{T}} \mathbf{C}^{-1} \mathbf{A}\right)^{1/2} \left(\mathbf{Y}^{\mathrm{T}} \mathbf{C}^{-1} \mathbf{Y}\right)^{1/2}}$

• *t*-statistic (used primarily for ID):

 $t = \frac{\mathbf{A}^{\mathrm{T}} \mathbf{C}^{-1} \mathbf{Y}}{\left(\mathbf{A}^{\mathrm{T}} \mathbf{C}^{-1} \mathbf{A}\right)^{1/2} \left(\left(\mathbf{Y} - \mathbf{A} \mathbf{X}\right)^{\mathrm{T}} \mathbf{C}^{-1} \left(\mathbf{Y} - \mathbf{A} \mathbf{X}\right)\right)^{1/2}}$



	Source	Definition
Y	Data	Pixel Spectrum (mean removed) of compensated data; measurement of ground-leaving radiance
с	Data	Covariance Matrix Describes variance within each spectral channel and correlation between spectral channels
A	Spectral Library	Target Signature (resampled to sensor spectral response); gas absorption coefficient or solid material reflectance
x	Calculated	Target Coefficient Matrix Obtained from GLS regression

- Identification (ID) by stepwise generalized least-squares (GLS)
- Spectral fitting in "whitened space"
 - Based on scene covariance matrix
- Algorithm proven in multiple field tests
 - Tested against ground-truth
 - Algorithm competitions and "blind" exercises
 - Up to 6 separate gases have been successfully unmixed and ID'd under test conditions
- Confidence levels







Tactical Processing: Urban-Industrial Environments

90 compounds of interest selected from a ~700-member spectral library

1,1,1,2-Tetrafluoroethane	Acrolein	Ethane	Methyl isobutyl ketone	Sulfur trioxide
1,1,1-Trichloroethane	Acrylonitrile	Ethanol	Methyl methacrylate	Sulfuryl fluoride
1,1,2-Trichloro-1,2,2-trifluoroethane	Ammonia	Ethene	Methylamine	tert-Butyl methyl ether
1,1-Dichloro-1-fluoroethane	Benzene	Ethyl acetate	Naphthalene	Tetrachloroethylene
1,1-Dichloroethane	Butane	Ethylbenzene	Nitric acid	Tetrahydrofuran
1,1-Difluoroethane	Carbon dioxide	Formaldehyde	Nitrogen dioxide	Toluene
1,2,4-Trimethylbenzene	Carbon tetrachloride	Heptane	Nitrogen dioxide and dimer	Trichloroethylene
1,2-Dichlorotetrafluoroethane	Carbonyl sulfide	Hexane	Nitrous oxide	Trichlorofluoromethane
1,2-Dibromoethane	Chlorodifluoromethane	Hydrogen peroxide	Octane	Vinyl chloride
1,2-Dichloroethane	Chloroform	Hydrogen sulfide	Ozone	a-Pinene
1,3-Butadiene	Chloropicrin	Isobutane	Pentane	d-Limonene
1,3-Dichloropropene	Cycloheptane	Isobutene	Peroxy acetyl nitrate (main bands)	m-Xylene
1,4-Dichlorobenzene	Cycloheptene	Isoprene	Phosgene	o-Xylene
1-Propanol	Cyclohexane	Isopropanol	Phosphine	p-Xylene
2,2-Dichlor-1,1,1-trifluorethane	Cyclohexene	Methane	Propane	
2-Butoxyethanol	Dichlorodifluoromethane	Methanol	Propene	
Acetaldehyde	Dichlorofluoromethane	Methyl bromide	Styrene	
Acetic acid	Dichloromethane	Methyl ethyl ketone	Sulfur dioxide	
Acetone	Diethyl ether	Methyl formate	Sulfur hexafluoride	

Tactical Analysis Output Sample

All emissions are time-stamped, geolocated, and chemically identified as illustrated in this excerpt from a typical automated tactical analysis report:

Plume Sources (LA Basin - 7/22/2014)				
Time (UTC)	(LON, LAT)	ID	t-stat	Source Information
17:40:24	-118.345305, 33.986106	1,1,1,2-Tetrafluoroethane	-14.8	Residence
17:41:24	-118.314280, 33.982817	Acetone	-26.9	Furniture and upholstery
17:41:33	-118.309144, 33.978966	Methanol	-14.6	Auto paint shop
17:43:02	-118.265361, 33.975626	Acetone	-13.8	Furniture factory
17:43:46	-118.242887, 33.992775	Acetone	-17.9	Iron worker
17:43:46	-118.243212, 33.972248	Acetone	-18.4	Screen printer
17:43:46	-118.242184, 34.010894	Chlorodifluoromethane	-10.2	Metal salvager
17:43:55	-118.237554, 34.002014	Ammonia	-23.6	Cold storage
17:43:59	-118.236606, 34.002475	Ammonia	-9.2	Cold storage
17:43:59	-118.235838, 33.990606	Ammonia	-27.4	Clothing store (probable furnace flue)
17:44:04	-118.234248, 33.986740	Ammonia	-11	Fabric finishing and lamination (probable furnace flue)
17:44:17	-118.226988, 33.995651	Ammonia	-24.6	Galvanizing
17:44:25	-118.223949, 34.000360	Ammonia	-15	Galvanizing
17:44:25	-118.223321, 34.006222	Ethanol	-22	Foods warehouse
17:44:30	-118.220774, 33.999024	Carbon dioxide	11.7	Waste handling/incinerator
17:44:30	-118.220774, 33.998867	Carbon dioxide	13.1	Waste handling/incinerator
17:44:34	-118.217737, 34.006523	Ammonia	-19.4	Produce warehouse
17:44:34	-118.217199, 33.996992	Carbon dioxide	11.3	Glass foundry
17:45:00	-118.204839, 33.982193	Ammonia	-24.6	Thermal processing of metals
17:45:00	-118.204715, 33.981642	Ammonia	-22.1	Thermal processing of metals
17:45:25	-118.191401, 33.997515	Ethene	-16.1	Produce Supplier
		Methane	-6	

Greenhouse Gas Focus: The Kyōto Basket

- The "Kyōto Basket" comprises five specific gases and two classes of halocarbons
- LWIR hyperspectral sensors address most of the Kyōto Basket gases:

Kyōto Basket Gas	Mako Sensitivity (kg/h)*
Carbon dioxide (CO ₂)	Very low (4800)
Methane (CH ₄)	Moderate (10)
Nitrous oxide (N ₂ O)	Moderate (10)
Hydrofluorocarbons (HFCs)	Very high/high (0.2–2.0)
Perfluorocarbons (PFCs)	Very high/high (0.2–1.0)
Sulfur hexafluoride (SF ₆)	Very high (0.1)
Nitrogen trifluoride (NF ₃)	Very high (0.5)

* Minimum detectable point-source emission flux assuming 5-m/s wind, 5-K thermal contrast, 2-m GSD, and urban/industrial spectral clutter statistics.

Kyōto Basket Species Observed in Urban Settings

Selected ACE detection filter images



- 1: Carbon dioxide (landfill flare)
- 3: Methane (international airport)
- 5: Nitrous oxide (industrial gas supplier)
- 7: Sulfur hexafluoride (aerospace plant)
- 9: 1,1-Difluoroethane, HFC-152a (residential building)
- 11: Pentafluoroethane + Difluoromethane, R-410 (office bldg.) (HFC-125) (HFC-32)

- 2: Carbon dioxide (oil refinery)
- 4: Methane (landfill)
- 6: Nitrous oxide (industrial gas supplier)
- 8: Sulfur hexafluoride (research facility)
- 10: 1,1,1,2-Tetrafluoroethane, HFC-134a (water dept. depot)
- 12: Pentafluoroethane + 1,1,1,2-Tetrafluoroethane + Difluoromethane, R-407 (gas station) (HFC-125) (HFC-134a) (HFC-32)

Broad-Area Survey of the Los Angeles Basin, 1 April 2019

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I,1,1-TRIFLUOROETHANE	
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ACETIC ACID DIMER	
	• >2500 km² acquired in 3 hours
	• 2-m GSD: all detected emission sources
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	commercial and residential areas
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I D NITROUS OXIDE	an ideal "laboratory" for testing and
	evaluating gas sensing techniques.
PROPANE	
> C STYRENE	• Multiple years of collections provide
	insight into emission trends
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Broad-Area Survey of the Los Angeles Basin: HFC-134a



Broad-Area Survey of the Los Angeles Basin: Ammonia



Broad-Area Survey of the Los Angeles Basin: Methane



Long-Range Plume Tracking



Area Covered: ~52 km² Collection Time: ~2.3 min

Ammonia slip from NO_x emission control system (SCR reactor) at gas-fired power plant; plume length ~8 km

Plume sources are two power plant stacks (red arrows). Note the well-defined bifurcate structure.

Additional plume sources downwind from primary sources (yellow arrows) are from oil refinery

Grayscale ACE overlain on satellite imagery

Methane release from wastewater treatment plant; plume length ~4 km

Methane generated at the plant supplies a nearby power generating station. Gas excess to capacity is usually flared.

False color ACE overlain on thermal radiance mosaic



Kern Oil Fields, Bakersfield, CA

Venerable production field with enhanced recovery (steam injection)





Source: DOE

Kern Oil Fields, Bakersfield, CA

Broad-area context with high spatial resolution



Thermal radiance scene image encoded in false color to clarify structural details on the ground

Multiple methane plumes from several sources coalesce into a diffuse haze detectable more than 1 km downwind

(GSD: 1 m)

Topical Question of the Day

LETTER

Resurgence in atmospheric abundance of CFC-11 (trichlorofluoromethane)

Nature, Vol. 569, pp. 546-550 (2019)

https://doi.org/10.1038/s41586-019-1193-4

Increase in CFC-11 emissions from eastern China based on atmospheric observations

M. Rigby^{1,15}, S. Park^{2,15}*, T. Saito^{3,15}, L. M. Western^{1,15}, A. L. Redington^{4,15}, X. Fang^{5,15}, S. Henne⁶, A. J. Manning⁴, R. G. Prinn⁵, G. S. Dutton^{7,8}, P. J. Fraser⁹, A. L. Ganesan¹⁰, B. D. Hall⁷, C. M. Harth¹¹, J. Kim¹¹, K.-R. Kim², P. B. Krummel⁹, T. Lee², S. Li¹², Q. Liang¹³, M. F. Lunt¹⁴, S. A. Montzka⁷, J. Mühle¹¹, S. O'Doherty¹, M.-K. Park¹², S. Reimann⁶, P. K. Salameh¹¹, P. Simmonds¹, R. L. Tunnicliffe¹, R. F. Weiss¹¹, Y. Yokouchi³ & D. Young¹

- LWIR HSI is highly sensitive to halocarbons
- Broad-area airborne surveys enable identification and geolocation of emission hotspots on regional scales
- Facilitates improved regulatory compliance

Foam blowing at construction site



HVAC unit "in distress"



Vehicle dismantler





Persistent Observations of Built Environments

Background

- Collaborative venture with New York University's Center for Urban Science and Progress
- Near-continuous operation from a static vantage point over a period of 8 days in April 2015
- 5402 data cubes collected (accompanied by context camera imagery)

• Synopsis

- Large clouds of HCFC-22 (chlorodifluoromethane), sometimes occupying more than 6 city blocks, persisting for several minutes, and disappearing/reappearing as the gas migrates through the urban canyons
 - HCFC-22 is being phased out due to its ozone depletion potential and its listing as a potent greenhouse agent with a global warming potential ~1800 times that of CO₂
- Large clouds of a mixture of difluoromethane (HFC-32) and pentafluoroethane (HFC-125), which is sold as R-410 and was designed to replace HCFC-22
 - These plumes were also large, but not as prominent as the HCFC-22 plumes
- Smaller plumes of 1,1,1,2-tetrafluoroethane (HFC-134a) were occasionally observed
- Also observed plumes of methane, ethene, CO_2 , and NH_3 emanating from many buildings



Sample Gas Detection Images

- In these examples the upper panel shows the ACE-filter gas detection image, while the lower panel shows the same scene in the LWIR with the pixel ensemble of strongest SNR overlain in red
- The HCFC-22 case at left shows how this prominent plume switches between observation in emission (when viewed against the cold sky) and observation in absorption (when viewed against warmer buildings) as it diffuses throughout the scene
- The SO₂ case at right depicts a cruise ship progressing up the Hudson River while burning sulfur-rich fuel



Time Series Analyses Expose "Rhythms of the City"





HCFC-22 is being phased out *per* the Montreal Protocol. All production and import is scheduled to be eliminated by 2020, although recycled HCFC-22 remains in circulation to service existing air conditioners.

Sensor operational periods

Observations suggest that venting of overpressured HVAC systems is synchronized within a few hours of local noon

Summary

- High spatio-spectral resolution airborne LWIR imaging spectrometry is a versatile tool for detecting and tracking GHG and other emissions within urban and non-urban environments
- High spatial resolution (1-2 m) permits trace-back of emission plumes to their source
 - Plumes can range in size from a few pixels to several km in extent
- High spectral resolution enables precise identification and discrimination of mixed gas plume components through the application of SMF and/or ACE algorithms
 - Quantification of emission rate is possible with knowledge of prevailing meteorology
- Application to multiple sectors
 - Regulatory monitoring
 - Mitigation of public health concerns
 - Post-disaster hazard evaluation
 - Regional climate assessment
- Operation in the emissive LWIR spectral region avoids reliance on solar illumination, allowing
 operations to be conducted day and night
- Further information: psl-isd@aero.org

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Further Reading

- "Tracking and quantification of gaseous chemical plumes from anthropogenic emission sources within the Los Angeles Basin," K. Buckland *et al.*, *Remote Sensing of Environment*, **201**, 275-296 (2017), doi:<u>10.1016/j.rse.2017.09.012</u>
- "Mapping refrigerant gases in the New York City skyline," M. Ghandehari *et al.*, Scientific Reports, 7, 2735 (2017), doi:<u>10.1038/s41598-017-02390-z</u>
- "MAHI: An airborne mid-infrared imaging spectrometer for industrial emissions monitoring," D. Tratt et al., IEEE Transactions on Geoscience and Remote Sensing, 55, 4558-4566 (2017), doi: 10.1109/TGRS.2017.2693979
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- "High areal rate longwave-infrared hyperspectral imaging for environmental remote sensing," D. Tratt et al., Proceedings of SPIE, 10639, 1063915 (2018), doi: 10.1117/12.2303834