

Preliminary Air Monitoring and Sampling Summary

Los Palos Warehouse Fire

Los Angeles, California

Prepared by: Onterris Response and Recovery, LLC

Air Monitoring Period: August 26, 2026

1.0 Introduction

Onterris Response and Recovery, LLC. (Onterris) personnel were contacted to provide air monitoring and environmental support regarding a cold-storage warehouse fire located at 1400 S. Los Palos Street in Los Angeles, California (herein referred to as the “Site”). Onterris personnel began real-time air monitoring on behalf of Lineage Logistics on June 23, 2026, at approximately 0006 Pacific Daylight Time (PDT). This report summarizes preliminary results of real-time air monitoring conducted by Onterris personnel from 0000 through 23:59 PDT on August 26, 2026, as well as results of VOC analysis in samples collected on August 1, August 3-4, and August 25, 2026, results of sulfur analysis in samples collected August 1 and August 3-4, 2026, and results of metal particulate analysis in samples collected on August 6 and August 24, 2026. All data provided in this summary are considered preliminary; these data have not undergone full quality assurance and quality control (QA/QC) review and are subject to change.

2.0 Air Monitoring and Sampling Methods

Prior to arrival onsite, Onterris personnel developed a Preliminary Air Sampling and Analysis Plan (SAP) to document and quantify analytes of concern potentially associated with the Site. Onterris personnel have updated the SAP based on site conditions and operational phases in coordination with the United States Environmental Protection Agency (US EPA) and South Coast Air Quality Management District (AQMD). The SAP (v1.2) is provided in **Appendix A** and includes air monitoring plans for Community/Fenceline Monitoring and Site Assessment. This preliminary summary includes Community and Fenceline Monitoring data with all testing locations listed in **Appendix B**.

Beginning July 11, 2026, Onterris field personnel discontinued air monitoring strategies employed during the Recovery Operational period and immediately transitioned to the Remediation Operational Period. The Onterris air monitoring and sampling approach during the Remediation Operational Period involved localizing air monitoring resources closer to the Site fenceline and nearby residential and commercial receptors, with all analytes measured at a height consistent with the typical human breathing zone. This approach incorporated two air monitoring and sampling strategies: 1) roaming (and/or mobile) real-time air monitoring using direct-reading instruments at the fenceline and nearby residential locations upwind, downwind, and crosswind from the Site, and 2) stationary air monitoring and sampling using a combination of real-time and analytical sampling methodologies at five discrete locations along, or directly across the street from, the fenceline (DU01-05), and at two schools, as listed in **Table 1**. All analytes, air sampling methodologies, and locations for stationary monitoring and sampling were selected in coordination with South Coast AQMD.

Table 1: Stationary Air Monitoring and Sampling Locations

Operational Period	Location Code	Location Description
Remediation	DU01	Northwest Corner Facility Fenceline
	DU02	North Side of Warehouse
	DU03	Northeast Corner Facility Fenceline
	DU04	East Side of Facility Fenceline
	DU05	Southeast Side of Facility Fenceline
	AS03	Maywood Elementary School
	AS07	Eastman Elementary School

2.1 Real-Time Air Monitoring Methods

Real-time air monitoring was conducted by Onterris in the community surrounding the Site and at the fenceline in coordination with US EPA and South Coast AQMD. Two air monitoring strategies were utilized and are described below. All instrumentation was calibrated daily or in accordance with manufacturer's recommendations.

Roaming Community/Fenceline Air Monitoring

Onterris personnel conducted roaming real-time community and fenceline monitoring for volatile organic compounds (VOCs), ammonia (NH₃), hydrogen sulfide (H₂S), chlorine (Cl₂), carbon monoxide (CO), atmospheric flammability measured as a percentage of the lower explosive limit (%LEL), oxygen (O₂), mercaptans, and formaldehyde. Monitoring was conducted using handheld instruments including Honeywell MultiRAEs with chemical-specific sensors and photoionization detectors (PIDs) and Gastec piston pumps with colorimetric detector tubes.

Onterris personnel also used a mobile AreaRAE instrument to continuously assess air at proposed community monitoring locations adjacent to the Site. The AreaRAE was equipped with sensors capable of measuring VOCs, %LEL, NH₃, H₂S, and O₂ concentrations in real-time at approximately 30-second intervals while recording GPS coordinates for each data point. VOC, %LEL, H₂S, and NH₃ detections at individual sensor limits of detection identified during mobile monitoring prompt field personnel to safely stop the vehicle and further assess conditions at the location.

Stationary Air Monitoring

During the Remediation Operational Period, stationary real-time telemetering air monitoring was conducted using DustTraks to assess particulates (PM_{2.5} and PM₁₀) at DU01-05 and AS07, while AreaRAEs were used to assess VOCs, %LEL, NH₃, CO, and O₂ along the Site fenceline at DU01-05. Additionally, datalogging SPM Flex devices were deployed and utilized at DU02, DU03, and AS07 for H₂S monitoring at concentrations as low as 0.001 ppb. All data were collected over 24-hour periods, from 0:00 PDT to 23:59 PDT, with respective real-time readings at each location yielding a daily average. For chemical concentrations yielding no detection, the sensor detection limit was used to derive conservative averages per analytes and respective locations.

2.2 Analytical Air Sampling Methods

To complement real-time air monitoring data collected at stationary air monitoring locations, analytical sampling was conducted for VOCs, sulfur compounds, and metal particulates over 24-hour periods in the community at DU02, DU03, AS03, and AS07 (**Table 1** and **Appendix C**). Analytical air sampling refers to the collection of discrete quantities of air using containers or chemical-specific media that are sent to an off-site laboratory for chemical analysis under chain of custody. All laboratories used for analytical air sampling are accredited through the National Environmental Laboratory Accreditation Conference (NELAC), California Environmental Laboratory Accreditation Program (CA ELAP), and/or American Industrial Hygiene Association (AIHA). Condition of analytical sampling containers collected by Onterris field personnel may dictate if samples are suitable for accessioning and processing by respective laboratories. Because of this, deployed analytical samples that are tampered with by non-field personnel or stolen, will be unavailable for laboratory processing. Laboratory results are reported as they become available.

VOCs and Sulfur

Following transition from the Recovery Operational Period to the Remediation Operational Period, Onterris personnel deployed air samples at DU02, DU03, AS03, and AS07 (**Appendix C**). Integrated air sampling was conducted using 1.4-L evacuated canisters to enable off-site laboratory analysis of VOCs via US EPA Method TO-15 and sulfur compounds via SCAQMD 307-91 or equivalent. Analytical air samples were collected for a duration of approximately 24 hours and were deployed at a height representative of the human breathing zone.

Metal Particulate Sampling

Air sampling for metals was conducted at DU02, DU03, AS03, and AS07 (**Appendix C**) using SKC sampling pumps calibrated to a flow rate of 3 liters per minute (L/min), and sampling pumps were set up to draw air through 0.8- μ m mixed cellulose ester membrane filter cassettes. Samples were collected over a 24-hour period (approximately 1,440 minutes) at a height representative of the human breathing zone and sent to an off-site laboratory under chain of custody for analysis of target metals via NIOSH Method 7303. Sample volume recommendations by the laboratory were followed for achieving appropriate lower detection limits for respective metal analysis.

3.0 Air Monitoring and Sampling Results

Results of real-time air monitoring are summarized in **Tables 2** through **5**. Maps of the incident site and real-time air monitoring locations are provided in **Appendix B**. Results of VOC analysis in air samples collected August 1, August 3-4, and August 25, 2026, and results of sulfur analysis in air samples collected on August 1 and August 3-4, 2026, are summarized in **Table 6**. Results of air samples for metal particulates collected August 6 and August 24, 2026, are summarized in **Table 7**.

The map of stationary air monitoring and analytical air sampling locations is provided in **Appendix C**. Meteorological data during the August 26, 2026, reporting period were acquired from the Hawthorne Municipal Airport and are provided in **Appendix D**.

Table 2: Community Handheld Air Monitoring Results
Collected August 26, 2026

Analyte	Instrument	Number of Readings	Number of Detections	Range of Detections*
Chlorine	Gastec #8La	6	0	< 0.05 ppm
CO	Gastec #1LC	26	0	< 0.5 ppm
Formaldehyde	Gastec #91LL	4	0	< 0.03 ppm
H ₂ S	MultiRAE	32	0	< 0.1 ppm
% LEL	MultiRAE	32	0	< 1. %
Mercaptans	Gastec #70L	29	0	< 0.05 ppm
NH ₃	Gastec #3L	21	0	< 0.2 ppm
NH ₃	MultiRAE	5	0	< 1. ppm
O ₂	MultiRAE	11	11	20.9 %
VOCs	MultiRAE	32	0	< 0.1 ppm

Abbreviations: ppm = parts per million

Note: This is a preliminary data summary, indicating that the data provided have not undergone the full quality assurance and quality control (QA/QC) process and should be considered preliminary at this time.

*Analytes with no detections are reported as less than (“<”) the instrument’s Limit of Detection (LOD)

Table 3: Fenceline Handheld Air Monitoring Results
Collected August 26, 2026

Analyte	Instrument	Number of Readings	Number of Detections	Range of Detections*
Chlorine	Gastec #8La	20	0	< 0.05 ppm
CO	Gastec #1LC	8	0	< 0.5 ppm
CO	MultiRAE	14	0	< 1. ppm
Formaldehyde	Gastec #91LL	14	0	< 0.03 ppm
H ₂ S	MultiRAE	40	0	< 0.1 ppm
% LEL	MultiRAE	60	0	< 1. %
Mercaptans	Gastec #70L	25	0	< 0.05 ppm
NH ₃	Gastec #3L	7	0	< 0.2 ppm
NH ₃	MultiRAE	55	0	< 1. ppm
O ₂	MultiRAE	60	60	20.5-20.9 %
VOCs	MultiRAE	61	2	0.1 ppm

Abbreviations: ppm = parts per million

Note: This is a preliminary data summary, indicating that the data provided have not undergone the full quality assurance and quality control (QA/QC) process and should be considered preliminary at this time.

*Analytes with no detections are reported as less than (“<”) the instrument’s Limit of Detection (LOD)

Table 4: Mobile AreaRAE Air Monitoring Results
Collected August 26, 2026

Date	Analyte	Number of Readings	Number of Detections	Range of Detections*
August 26, 2026	% LEL	5,651	0	< 1 %
	H ₂ S	5,651	0	< 0.1 ppm
	NH ₃	5,651	0	< 1 ppm
	O ₂	5,651	5,651	20.9 %
	VOCs	5,651	0	< 0.1 ppm

Abbreviations: ppm = parts per million

Note: This is a preliminary data summary, indicating that the data provided have not undergone the full quality assurance and quality control (QA/QC) process and should be considered preliminary at this time.

*Analytes with no detections are reported as less than (“<”) the instrument’s Limit of Detection (LOD)

Table 5: Fixed Station Air Monitoring Results
Collected August 26, 2026

Location Code	Instrument	Analyte	Number of Readings	Number of Detections	Average Concentration	Range of Detections*
DU01	AreaRAE	% LEL	5,412	0	ND	< 1 %
		CO	5,412	0	ND	< 1 ppm
		NH ₃	5,412	0	ND	< 1 ppm
		O ₂	5,412	5,412	20.9	20.9 %
		VOCs	5,412	0	ND	< 0.1 ppm
	DustTrak	PM ₁₀	8,600	8,600	0.015	0.002 - 0.409 mg/m ³
		PM _{2.5}	8,600	8,600	0.01	0.002 - 0.171 mg/m ³
DU02	AreaRAE	% LEL	5,414	0	ND	< 1 %
		CO	5,414	39	1	2 - 22 ppm
		NH ₃	5,414	0	ND	< 1 ppm
		O ₂	5,414	5,414	20.9	20.9 %
		VOCs	5,414	2	0.1	0.1 - 0.3 ppm
	DustTrak	PM ₁₀	8,414	8,414	0.025	0.003 - 0.341 mg/m ³
		PM _{2.5}	8,414	8,414	0.015	0.002 - 0.103 mg/m ³
	SPM Flex	H ₂ S	8,319	2,340	0.001	0.001 - 0.010 ppm
DU03	AreaRAE	% LEL	5,416	0	ND	< 1 %
		CO	5,416	1	1	3 ppm
		NH ₃	5,416	0	ND	< 1 ppm
		O ₂	5,416	5,416	20.9	20.9 %
		VOCs	5,416	0	ND	< 0.1 ppm
	DustTrak	PM ₁₀	8,475	8,475	0.014	0.002 - 0.452 mg/m ³
		PM _{2.5}	8,475	8,475	0.012	0.002 - 0.126 mg/m ³
	SPM Flex	H ₂ S	8,380	850	0.001	0.001 - 0.003 ppm

Location Code	Instrument	Analyte	Number of Readings	Number of Detections	Average Concentration	Range of Detections*
DU04	AreaRAE	% LEL	5,154	0	ND	< 1 %
		CO	5,151	0	ND	< 1 ppm
		NH ₃	5,154	0	ND	< 1 ppm
		O ₂	5,154	5,154	21	20.9 - 21.4 %
		VOCs	5,151	0	ND	< 0.1 ppm
	DustTrak	PM ₁₀	8,598	8,598	0.023	0.008 - 2.500 mg/m ³
		PM _{2.5}	8,598	8,598	0.016	0.008 - 0.915 mg/m ³
DU05	AreaRAE	% LEL	5,383	0	ND	< 1 %
		CO	5,383	1	1	12 ppm
		NH ₃	5,383	0	ND	< 1 ppm
		O ₂	5,383	5,383	20.9	20.9 %
		VOCs	5,383	0	ND	< 0.1 ppm
	DustTrak	PM ₁₀	8,606	8,606	0.022	0.006 - 0.696 mg/m ³
		PM _{2.5}	8,606	8,606	0.013	0.006 - 0.207 mg/m ³
AS07	DustTrak	PM ₁₀	8,583	8,583	0.01	0.004 - 0.053 mg/m ³
		PM _{2.5}	8,583	8,583	0.008	0.004 - 0.021 mg/m ³
	SPM Flex	H ₂ S	8,480	385	0.001	0.001 ppm

Abbreviations: ppm = parts per million; mg/m³ – milligrams per cubic meter; ND = No Detection (calculated mean not applicable)

Note: This is a preliminary data summary, indicating that the data provided have not undergone the full quality assurance and quality control (QA/QC) process and should be considered preliminary at this time.

*Analytes with no detections are reported as less than (“<”) the instrument’s Limit of Detection (LOD).

Table 6: Summary of Air Sampling Results for VOCs
Samples Collected August 1, August 3-4, and August 25, 2026
COCs 566622 (VOCs) and 564993, 565080, 565202 (VOCs and Sulfur)

Analytical Method	Analyte	Number of Results	Number of Detections	Concentration Range*
EPA TO-15	1,1-Dichloroethane	16	0	< 0.033 ppbv
	1,1-Dichloroethene	16	0	< 0.036 ppbv
	1,1-Difluoroethane	16	15	0.2 (J) ppbv - 0.59 (J) ppbv
	1,1,1-Trichloroethane	16	0	< 0.022 ppbv
	1,1,1,2-Tetrachloroethane	16	0	< 0.017 ppbv
	1,1,2-Trichloroethane	16	0	< 0.031 ppbv
	1,1,2,2-Tetrachloroethane	16	0	< 0.027 ppbv
	1,2-Dibromo-3-Chloropropane	16	0	< 0.036 ppbv
	1,2-Dibromoethane	16	0	< 0.029 ppbv
	1,2-Dichlorobenzene	16	0	< 0.021 ppbv
	1,2-Dichloroethane	16	0	< 0.035 ppbv
	1,2-Dichloropropane	16	0	< 0.039 ppbv
	1,2,3-Trichloropropane	16	0	< 0.015 ppbv
	1,2,4-Trichlorobenzene	16	1	0.064 (J) ppbv

Analytical Method	Analyte	Number of Results	Number of Detections	Concentration Range*
EPA TO-15	1,2,4-Trimethylbenzene	16	11	0.032 (J) ppbv - 0.062 (J)
	1,3-Butadiene	16	0	< 0.033 ppbv
	1,3-Dichlorobenzene	16	0	< 0.045 ppbv
	1,3-Dichloropropane	16	0	< 0.025 ppbv
	1,3,5-Trimethylbenzene	16	0	< 0.031 ppbv
	1,4-Dichlorobenzene	16	0	< 0.054 ppbv
	1,4-Dioxane	16	0	< 0.026 ppbv
	2-Butanone	16	16	0.75 (J) ppbv - 2.2 (J) ppbv
	2-Chlorotoluene	16	0	< 0.022 ppbv
	2-Hexanone	16	11	0.044 (J) ppbv - 0.11 (J)
	2,2,4-Trimethylpentane	16	16	0.17 (J) ppbv - 0.43 ppbv
	3-Chloropropene	16	0	< 0.12 ppbv
	4-Ethyltoluene	16	0	< 0.019 ppbv
	4-Methyl-2-Pentanone	16	16	0.074 (J) ppbv - 6.9 ppbv
	Acetaldehyde	16	16	6.6 ppbv - 23 ppbv
	Acetone	16	16	120 ppbv - 650 ppbv
	Acetonitrile	16	16	0.19 (J) ppbv - 1.1 ppbv
	Acrolein	16	16	0.26 (J) ppbv - 0.62 ppbv
	Acrylonitrile	16	16	0.15 (J) ppbv - 87 ppbv
	Benzene	16	16	0.14 (J) ppbv - 0.36 (J) ppbv
	Benzyl chloride	16	0	< 0.043 ppbv
	Bromodichloromethane	16	0	< 0.032 ppbv
	Bromoform	16	0	< 0.021 ppbv
	Bromomethane	16	0	< 0.032 ppbv
	Butane	16	16	0.72 ppbv - 1.7 ppbv
	Carbon Disulfide	16	16	0.043 (J) ppbv - 0.32 (J)
	Carbon Tetrachloride	16	16	0.066 (J) ppbv - 0.087 (J)
	Chlorobenzene	16	0	< 0.029 ppbv
	Chlorodifluoromethane	16	16	0.29 (J) ppbv - 0.7 ppbv
	Chloroethane	16	16	0.047 (J) ppbv - 1.4 ppbv
	Chloroform	16	16	0.021 (J) ppbv - 0.06 (J)
	Chloromethane	16	16	0.46 ppbv - 0.73 ppbv
	cis-1,2-Dichloroethene	16	0	< 0.028 ppbv
	cis-1,3-Dichloropropene	16	0	< 0.024 ppbv
	Cyclohexane	16	15	0.047 (J) ppbv - 0.12 (J)
	Dibromochloromethane	16	0	< 0.018 ppbv
	Ethanol	16	16	35 ppbv - 720 ppbv
	Ethyl Acetate	16	16	0.7 ppbv - 7.1 ppbv
	Ethyl tert-Butyl Ether (ETBE)	16	0	< 0.023 ppbv
	Ethylbenzene	16	16	0.035 (J) ppbv - 0.087 (J)
	Hexachlorobutadiene	16	0	< 0.031 ppbv
	Hexachloroethane	16	0	< 0.035 ppbv
	Isopropanol (IPA)	16	16	3.6 ppbv - 29 ppbv

Analytical Method	Analyte	Number of Results	Number of Detections	Concentration Range*
EPA TO-15	Isopropyl Ether (DIPE)	16	0	< 0.048 ppbv
	Isopropylbenzene	16	0	< 0.023 ppbv
	m,p-Xylenes	16	16	0.1 (J) ppbv - 0.23 (J) ppbv
	Methanol	16	16	15 ppbv - 51 ppbv
	Methyl methacrylate	16	1	0.06 (J) ppbv
	Methyl tert-Amyl Ether (TAME)	16	0	< 0.022 ppbv
	Methylene Chloride	16	16	0.13 (J) ppbv - 0.39 ppbv
	MTBE	16	0	< 0.033 ppbv
	n-Butylbenzene	16	0	< 0.023 ppbv
	n-Heptane	16	16	0.083 (J) ppbv - 0.18 (J)
	n-Hexane	16	16	0.15 (J) ppbv - 0.34 (J) ppbv
	n-Nonane	16	2	0.047 (J) ppbv - 0.049 (J)
	n-Pentane	16	16	0.42 ppbv - 0.76 ppbv
	Naphthalene	16	0	< 0.097 ppbv
	o-Xylene	16	15	0.042 (J) ppbv - 0.088 (J)
	Propylbenzene	16	0	< 0.027 ppbv
	Propylene	16	16	0.056 (J) ppbv - 0.54 (J)
	sec-Butylbenzene	16	0	< 0.018 ppbv
	Styrene	16	15	0.04 (J) ppbv - 0.16 (J) ppbv
	tert-Butyl Alcohol (TBA)	16	16	0.21 (J) ppbv - 4.5 ppbv
	tert-Butylbenzene	16	0	< 0.021 ppbv
	Tetrachloroethene	16	4	0.03 (J) ppbv - 0.061 (J)
	Tetrahydrofuran	16	0	< 0.078 ppbv
	Toluene	16	16	0.36 ppbv - 0.92 ppbv
	trans-1,2-Dichloroethene	16	0	< 0.027 ppbv
	trans-1,3-Dichloropropene	16	0	< 0.019 ppbv
	Trichloroethene	16	0	< 0.029 ppbv
	Trichlorofluoromethane	16	16	0.18 (J) ppbv - 0.21 (J) ppbv
	Vinyl Acetate	16	0	< 0.29 ppbv
	Vinyl bromide	16	0	< 0.022 ppbv
	Vinyl Chloride	16	0	< 0.022 ppbv
	Xylene (total)	16	16	0.1 (J) ppbv - 0.32 (J) ppbv
	Freon 12	16	16	0.42 ppbv - 0.49 ppbv
	Freon 113	16	15	0.05 (J) ppbv - 0.068 (J)
	Freon 114	16	0	< 0.02 ppbv
SCAQMD 307-91	2 & 3-Ethyl Thiophene	12	0	< 7.94 (H) ppbv
	2 & 3-Methyl Thiophene	12	0	< 7.94 (H) ppbv
	2,5 Dimethylthiophene	12	0	< 7.94 (H) ppbv
	Carbon Disulfide	12	0	< 7.94 (H) ppbv
	Carbonyl Sulfide	12	0	< 7.94 (H) ppbv
	Diethyl Disulfide	12	0	< 7.94 (H) ppbv
	Diethyl Sulfide	12	0	< 7.94 (H) ppbv
	Dimethyl Disulfide	12	0	< 7.94 (H) ppbv

Analytical Method	Analyte	Number of Results	Number of Detections	Concentration Range*
SCAQMD 307-91	Dimethyl Sulfide	12	0	< 7.94 (H) ppbv
	Ethyl Mercaptan	12	0	< 7.94 (H) ppbv
	Hydrogen Sulfide	12	0	< 7.94 (H) ppbv
	iso-Butyl Mercaptan	12	0	< 7.94 (H) ppbv
	iso-Propyl Mercaptan	12	0	< 7.94 (H) ppbv
	Methyl Mercaptan	12	0	< 7.94 (H) ppbv
	n-Butyl Mercaptan	12	0	< 7.94 (H) ppbv
	n-Propyl Mercaptan	12	0	< 7.94 (H) ppbv
	sec-Butyl Mercaptan	12	0	< 7.94 (H) ppbv
	Sulfur Dioxide	12	0	< 7.94 (H) ppbv
	tert-Butyl Mercaptan	12	0	< 7.94 (H) ppbv
	Tetrahydrothiophene	12	0	< 7.94 (H) ppbv
	Thiophene	12	0	< 7.94 (H) ppbv
	Total Non-Target as H ₂ S	12	0	< 7.94 (H) ppbv
	Total Reduced Sulfur as H ₂ S	12	0	< 7.94 (H) ppbv
	Total Sulfur as H ₂ S	12	0	< 7.94 (H) ppbv

Abbreviations: ppbv = parts per billion by volume

(J) indicates the result is an estimated value below the laboratory reporting limit

(H) indicates the hold time was exceeded.

Note: This is a preliminary data summary, indicating that the data provided have not undergone the full quality assurance and quality control (QA/QC) process and should be considered preliminary at this time.

*Analytes with no detections are reported as less than ("<") the laboratory's method detection limit (MDL)

Table 7: Summary of Air Sampling Results for Metal Particulates
Samples Collected August 6 and August 24, 2026
COC L705189

Analyte	Number of Results	Number of Detections	Concentration Range*
Arsenic	8	0	< 0.035 µg/m ³
Cadmium	8	0	< 0.0035 µg/m ³
Chromium	8	0	< 0.52 µg/m ³
Cobalt	8	0	< 0.011 µg/m ³
Copper	8	0	< 0.069 µg/m ³
Iron Oxide	8	4	2.5 µg/m ³ - 8 µg/m ³
Lead	8	0	< 0.017 µg/m ³
Lithium	8	0	< 0.035 µg/m ³
Manganese	8	4	0.068 µg/m ³ - 0.1 µg/m ³
Nickel	8	0	< 0.069 µg/m ³
Zinc Oxide	8	0	< 1.1 µg/m ³

Abbreviations: µg/m³ = micrograms per cubic meter

Note: This is a preliminary data summary, indicating that the data provided have not undergone the full quality assurance and quality control (QA/QC) process and should be considered preliminary at this time.

*Analytes with no detections are reported as less than ("<") the laboratory's method detection limit (MDL)

Appendix A Sampling and Analysis Plan

Air Sampling and Analysis Plan

Boyle Heights Warehouse Fire
Los Angeles, California

Prepared by: Onterris Response and Recovery, LLC

June 27, 2026

Version 1.2

	Name/Organization	Signature	Date Signed
Prepared by:	Max Ford, PhD / Onterris	<i>Max Ford</i>	6/23/2026
Reviewed by:			
Reviewed by:			

Air Monitoring and Sampling Strategy

Onterris is focusing on the mixtures, chemicals, and indicators of flammability chosen below because they are among the most important and readily monitored airborne hazards during a cold storage facility fire. The possible hazards may vary by the source and type of refrigerant used within the facility as well as with the combustion and environmental conditions associated with the incident. Monitoring and/or sampling for some chemicals associated with the event may be added, conducted less frequently, or discontinued as site-specific information becomes available or as monitoring and/or sampling results indicate that these chemicals and indicators do not pose a health concern.

The strategy is to utilize three broadly defined monitoring plans, namely **1) Community Monitoring**, **2) and 3) Site Assessment**. While some of these plans may not be implemented immediately, this document delineates variations in these monitoring plans, should they be implemented. Community Monitoring will occur in publicly accessible community areas, including residential and/or commercial locations surrounding the incident site, at a height representative of the human breathing zone. Community Monitoring may also include Fenceline Monitoring, which may occur at the perimeter or outer fenceline of the incident site or work area, at a height representative of the human breathing zone. If conducted, detections observed during Fenceline Monitoring are intended to be a first-line indicator of potential issues at the site and are not necessarily indicative of exposure to downwind receptors. Unlike Community Monitoring, Site Assessment may involve a variety of different monitoring tasks intended to provide information that may help to delineate the nature and extent of the release (e.g., worst case determination, container headspace, ground level) and is not necessarily representative of ambient air monitoring near breathing zone level.

Free-roaming handheld real-time air monitoring may be conducted in a variety of areas based on levels of activity, proximity to the release, and site conditions. The presence or absence of odors, including odor characteristics and description if an odor is detected, will be documented with every handheld real-time air monitoring reading.

Radio-telemetry real-time air monitoring units may be deployed at discrete (stationary) locations to allow for continuous air monitoring in multiple areas. Readings collected using stationary real-time air monitoring instruments will be received and monitored in a centralized location by field personnel to allow for recognition, communication, and rapid response to changing conditions.

Air samples may be collected and sent to an off-site laboratory for chemical analysis. These analytical air sampling techniques may be used to provide air quality data beyond the scope of real-time instruments. When necessary, air samples may be collected on individual workers (personal sampling) to provide exposure data over the course of a work shift for more direct comparison to occupational exposure values.

Site-Specific Action Levels

Onterris site-specific action levels are employed to provide information for corrective action to limit potential exposures. These values do not replace occupational or community exposure standards or guidelines but, rather, are intended to represent a concentration limit that triggers a course of action to better address worker and public safety. Action level exceedances will be communicated to Site Management and the Onterris Project Manager (PM) and/or Project Technical Director (PTD). Work practices will be assessed and adjusted as necessary. Site-specific action levels are not utilized for Site Assessment monitoring.

Plan 1: Community/Fenceline Monitoring

Objective: Report air levels before they reach those causing nuisance or health issues

Analyte	Action Level ¹	Action to be Taken	Basis ¹	Instrument	Detection Limit	Notes	Correction Factor
Total volatile organic compounds (VOCs) ²	0.5 ppm 15 min	Assess for ammonia vapor with secondary instrument; note odors for all detections; report ammonia detections to PM/PTD	1/6 of AEGL-1 for ammonia	MultiRAE PID (10.6 eV lamp)	0.1 ppm	Range: 0.1 – 5,000 ppm	Variable ⁴
Ammonia	Detection	Confirm with secondary instrument; note odors for all detections; report reading to PM/PTD.	Inform PM/PTD of potential off-site impact	MultiRAE Sensor AreaRAE Sensor	1.0 ppm	Range: 1 – 100 ppm	N/A
				Gastec #3L	0.2 ppm	Range: 1 – 30 ppm Volume: 100 mL Pull one 100-mL pump stroke	Variable ⁴
Hydrogen Sulfide	Detection	Assess for hydrogen sulfide with secondary instrument; note odors for all detections; report hydrogen sulfide detections to PM/PTD	Inform PM/PTD of potential off-site impact	MultiRAE Sensor AreaRAE Sensor	0.1 ppm	Range: 0 – 100 ppm	NA
				Gastec tube #4LL	0.1 ppm	Range: 0.25 – 120 ppm Volume: 1,000 mL (10 x 100 mL pump strokes)	Var.
Chlorine	Detection	Assess for chlorine with secondary instrument; note odors for all detections; report hydrogen sulfide detections to PM/PTD	Indicator of potential off-site impacts	Gastec tube #8La	0.05 ppm	Range: 0.1 – 16 ppm Volume: 500 mL (5 x 100-mL pump strokes)	Variable ⁴
Mercaptans	Detection	Assess for mercaptans with secondary instrument; note odors for all detections; report hydrogen sulfide detections to PM/PTD	Indicator of potential off-site impacts	Gastec tube #70L	0.1 ppm	Range: 0.1 – 8 ppm Volume: 400 mL (4 x 100-mL pump strokes)	Variable ⁴
Oxygen (O ₂)	< 19.5 % Immediate	Exit area; Report reading to PM/PTD	O ₂ -deficient atmosphere, as defined by 29 CFR 1910.134	MultiRAE Sensor	0.1 %	Range: 0 – 30 %	N/A
				AreaRAE Sensor	0.1 %	Range: 0 – 30 %	N/A

¹ Community action levels are generally set as “detection” to inform the PM/PTD of any potential off-site impact. Protective Action Criteria (PAC-1) established by the United States Department of Energy (US DOE) for the chemical analytes listed above are provided in Table 1. A PAC-1 is the airborne concentration (expressed as ppm [parts per million] or mg/m³ [milligrams per cubic meter]) of a substance above which it is predicted that the general population, including susceptible individuals, when exposed for more than one hour, could experience notable discomfort, irritation, certain asymptomatic, or nonsensory effects, or may perceive a clearly defined objectionable odor. However, these effects are not disabling and are transient and reversible upon cessation of exposure.

² 10.6 eV PID must be used for ammonia concentrations over 50 ppm to avoid ammonia sensor exhaustion/damage.

³ Readings collected using the MultiRAE PID and AreaRAE PID are recorded as uncorrected (raw) concentrations. Correction factors for various compounds are listed in Honeywell Technical Notes TN-106B (MultiRAE) and TN-106A (AreaRAE).

⁴ Readings collected using Gastec tubes are recorded as uncorrected (raw) concentrations. The correction factors for Gastec tubes are dependent on volume of air used (i.e., number of pump strokes).

Abbreviations: VOCs = volatile organic compounds; ppm = parts per million; PM = Onterris Project Manager; PTD = Onterris Project Technical Director; PID = photoionization detector; eV = electron volt; N/A = not applicable; CFR = Code of Federal Regulations

Plan 1: Community/Fenceline Monitoring – Combustion Products¹

Analyte	Action Level	Action to be Taken	Basis ²	Instrument	Detection Limit	Notes	Correction Factor
Particulate matter with a diameter of 2.5 microns or less (PM _{2.5})	0.035 mg/m ³ 1 hour	Report reading to PM/PTD	Upper-bound breakpoint for AQI category considered Moderate	TSI SidePak AM510/AM520 TSI DuskTrak	0.001 mg/m ³	Range: 0.001 – 100 mg/m ³ PM _{2.5} impactor: 50% cut-off at 2.5 microns	0.38 ³
Particulate matter with a diameter of 10 microns or less (PM ₁₀)	0.15 mg/m ³ 1 hour	Report reading to PM/PTD	Upper-bound breakpoint for AQI category considered Moderate	TSI SidePak AM510/AM520 TSI DuskTrak	0.001 mg/m ³	Range: 0.001 – 100 mg/m ³ PM ₁₀ impactor: 50% cut-off at 10 microns	0.38 ³
Carbon monoxide (CO)	Detection	Report reading to PM/PTD	Inform PM/PTD of potential off-site impact	MultiRAE Sensor AreaRAE Sensor	1 ppm	Range: 1 – 500 ppm	NA
				Gastec tube #1LC	0.5 ppm	Range: 1 – 30 ppm Volume: Pull 1 100-mL pump strokes	Variable ⁴
Sulfur dioxide (SO ₂)	Detection	Report reading to PM/PTD	Inform PM/PTD of potential off-site impact	Gastec tube #5Lb	0.01 ppm	Range: 0.05 – 10 ppm Volume: Pull 8 100-mL pump strokes	Variable ⁴
				MultiRAE Sensor AreaRAE Sensor	0.1 ppm	Range: 0.1 – 20 ppm	N/A
Nitrogen dioxide (NO ₂)	Detection	Report reading to PM/PTD	Inform PM/PTD of potential off-site impact	MultiRAE Sensor AreaRAE Sensor	0.1 ppm	Range: 0.1 – 20 ppm	N/A
				Gastec tube #9L	0.1 ppm	Range: 0.5 – 125 ppm Volume: Pull 2 100-mL pump strokes	Variable ⁴

¹ Monitoring for combustion products may be discontinued when the fire is extinguished.

² Community action levels are generally set as “detection” to inform the PM/PTD of any potential off-site impact. Protective Action Criteria (PAC-1) established by the United States Department of Energy (US DOE) for the chemical analytes listed above are provided in Table 1. A PAC-1 is the airborne concentration (expressed as ppm [parts per million] or mg/m³ [milligrams per cubic meter]) of a substance above which it is predicted that the general population, including susceptible individuals, when exposed for more than one hour, could experience notable discomfort, irritation, certain asymptomatic, or nonsensory effects, or may perceive a clearly defined objectionable odor. However, these effects are not disabling and are transient and reversible upon cessation of exposure.

³ Readings collected using the SidePak AM510/AM520 or DustTrak are recorded as uncorrected (raw) concentrations. A Photometric Calibration Factor (PCF) of 0.38 is used, which TSI indicates is appropriate for ambient/fugitive emissions monitoring applications. Photometric instruments (e.g., SidePak AM520) will overstate the mass of an aerosol compared to its gravimetric weight due to humid conditions; PM_{2.5} is especially prone to interference from high humidity. In cases of high humidity, PM₁₀ impactors may be used, which are not as sensitive to humidity.

⁴ Readings collected using Gastec tubes are recorded as uncorrected (raw) concentrations. The correction factors for Gastec tubes are dependent on volume of air used (i.e., number of pump strokes).

Abbreviations: mg/m³ = milligrams per cubic meter; PM = Onterris Project Manager; PTD = Onterris Project Technical Director; CO = carbon monoxide; ppm = parts per million; N/A = not applicable;

Plan 1: Community/Fenceline Monitoring – Flammability

Analyte	Action Level	Action to be Taken	Basis	Instrument	Detection Limit	Notes	Correction Factor
% LEL	1 % Instant	Report reading to PM/PTD	Detection of % LEL	MultiRAE Sensor AreaRAE Sensor	1 %	Range: 0 – 100 %	Variable ¹

¹ Readings collected using the % LEL sensor are recorded as uncorrected (raw) concentrations. Correction factors for various compounds are listed in Honeywell Technical Note TN-156.

Abbreviations: % LEL = atmospheric flammability measured as a percentage of the lower explosive limit; PM = Onterris Project Manager; PTD = Onterris Project Technical Director

TABLE 1: Community Health-Based Screening Values for Analytes (Air Monitoring)

Analyte	CAS RN	Protective Action Criteria 1 (PAC-1) ¹	OEHHA Acute Reference Exposure Level (REL) ²
Total volatile organic compounds (VOCs)	N/A	N/A	N/A
Ammonia	7664-41-7	30 ppm	3,200 µg/m ³ (4.59 ppm)
Carbon monoxide (CO)	630-08-0	75 ppm	23,000 µg/m ³ (20.08 ppm)
Sulfur dioxide (SO ₂)	7446-09-5	0.2 ppm	660 µg/m ³ (0.25 ppm)
Nitrogen dioxide (NO ₂)	10102-44-0	0.5 ppm	470 µg/m ³ (0.25 ppm)

¹ United States Department of Energy (US DOE): <https://www.energy.gov/ehss/protective-action-criteria-pac-aegls-erpgs-teels>. A Protective Action Criteria 1 (PAC-1) is the airborne concentration of a substance above which it is predicted that the general population, including susceptible individuals, when exposed for more than one hour, could experience notable discomfort, irritation, or certain asymptomatic, nonsensory effects, or may perceive a clearly defined objectionable odor. However, these effects are not disabling and are transient and reversible upon cessation of exposure.

² Office of Environmental Health Hazard Assessment (OEHHA) of the California Environmental Protection Agency (CalEPA): <https://oehha.ca.gov/air/general-info/oehha-acute-8-hour-and-chronic-reference-exposure-level-rel-summary>. An Acute Reference Exposure Level (REL) is the airborne concentration of a substance below which is not expected to cause non-cancer health effects for an exposure of one hour or less.

Abbreviations: CAS RN = Chemical Abstracts Service Registry Number; PAC-1 = Protective Action Criteria 1; OEHHA = Office of Environmental Health Hazard Assessment; REL = Reference Exposure Level; VOCs = volatile organic compounds; CO = carbon monoxide; ppm = parts per million; µg/m³ = micrograms per cubic meter; HCl = hydrogen chloride; HCN = hydrogen cyanide; HF = hydrogen fluoride; SO₂ = sulfur dioxide; NO₂ = nitrogen dioxide

TABLE 2: US EPA AQI Categories Associated with Airborne PM_{2.5} and PM₁₀ Levels

US EPA AQI Category	24-Hour Average PM _{2.5} Concentration (µg/m ³)	24-Hour Average PM ₁₀ Concentration (µg/m ³)
Good (0 – 50)	0.0 – 9.0	0 – 54
Moderate (51 – 100)	9.1 – 35.4	55 – 154
Unhealthy for Sensitive Groups (101-150)	35.5 – 55.4	155 – 254
Unhealthy (151 – 200)	55.5 – 125.4	255 – 354
Very Unhealthy (201-300)	125.5 – 225.4	355 – 424
Hazardous (301+)	225.5+	425+

Abbreviations: US EPA = United States Environmental Protection Agency; AQI = Air Quality Index; PM_{2.5} = particulate matter with a diameter of 2.5 microns or less; PM₁₀ = particulate matter with a diameter of 10 microns or less; µg/m³ = micrograms per cubic meter

Plan 2: Site Assessment

Objective: Characterize nature and extent of potential release

Analyte	Action Level	Action to be Taken	Basis	Instrument	Detection Limit	Notes	Correction Factor
Total volatile organic compounds (VOCs)	N/A	Report reading to PM/PTD	N/A	MultiRAE PID (10.6 eV lamp)	0.1 ppm	Range: 0.1 – 5,000 ppm	Variable ¹
				AreaRAE PID (10.6 eV lamp)	0.1 ppm	Range: 0.1 – 5,000 ppm	Variable ¹
				MultiRAE Sensor AreaRAE Sensor	1.0 ppm	Range: 1 – 100 ppm	N/A
Ammonia	N/A	Report reading to PM/PTD	N/A	Gastec #3L	0.2 ppm	Range: 1 – 30 ppm Volume: 100 mL (1 x 100 mL pump strokes)	Variable ³
				Gastec #3M	2 ppm	Range: 10 – 500 ppm Volume: 500 mL (5 x 100-mL pump strokes)	Variable ³
Hydrogen Sulfide	N/A	Report reading to PM/PTD	N/A	MultiRAE Sensor AreaRAE Sensor	0.1 ppm	Range: 0 – 100 ppm	NA
				Gastec tube #4LL	0.1 ppm	Range: 0.25 – 120 ppm Volume: 1,000 mL (10 x 100 mL pump strokes)	Var.
Chlorine	N/A	Report reading to PM/PTD	N/A	Gastec tube #8La	0.05 ppm	Range: 0.1 – 16 ppm Volume: 500 mL (5 x 100-mL pump strokes)	Variable ⁴
Mercaptans	N/A	Report reading to PM/PTD	N/A	Gastec tube #70L	0.1 ppm	Range: 0.1 – 8 ppm Volume: 400 mL (4 x 100-mL pump strokes)	Variable ⁴
Oxygen (O ₂)	N/A	Report reading to PM/PTD	N/A	MultiRAE Sensor	0.1 %	Range: 0 – 30 %	N/A
				AreaRAE Sensor	0.1 %	Range: 0 – 30 %	N/A

¹ Readings collected using the MultiRAE PID and AreaRAE PID are recorded as uncorrected (raw) concentrations. Correction factors for various compounds are listed in Honeywell Technical Notes TN-106B (MultiRAE) and TN-106A (AreaRAE).

² Readings collected using Gastec tubes are recorded as corrected concentrations. The correction factors for Gastec tubes are dependent on volume of air used (i.e., number of pump strokes).

³ 10.6 eV PID must be used for ammonia concentrations over 50 ppm to avoid ammonia sensor exhaustion/damage.

Abbreviations: VOCs = volatile organic compounds; NA = not applicable; PM = Onterris Project Manager; PTD = Onterris Project Technical Director; PID = photoionization detector; eV = electron volt; ppm = parts per million; O₂ = oxygen

Plan 2: Site Assessment – Combustion Products

Analyte	Action Level	Action to be Taken	Basis	Instrument	Detection Limit	Notes	Correction Factor
Particulate matter with a diameter of 2.5 microns or less (PM _{2.5}) or 10 microns or less (PM ₁₀)	N/A	Report reading to PM/PTD	N/A	TSI SidePak AM510/AM520	0.001 mg/m ³	Range: 0.001 – 100 mg/m ³ PM _{2.5} impactor: 50% cut-off at 2.5 microns; PM ₁₀ impactor: 50% cut-off at 10 microns	0.38 ¹
				DustTrak	0.001 mg/m ³	Range: 0.001 – 100 mg/m ³ PM _{2.5} impactor: 50% cut-off at 2.5 microns; PM ₁₀ impactor: 50% cut-off at 10 microns	0.38 ¹
Carbon monoxide (CO)	N/A	Report reading to PM/PTD	N/A	MultiRAE Sensor AreaRAE Sensor	1 ppm	Range: 1 – 500 ppm	N/A
				Gastec tube #1LC	0.5 ppm	Range: 1 – 30 ppm Volume: Pull 1 100-mL pump strokes	Variable ²
Sulfur dioxide (SO ₂)	N/A	Report reading to PM/PTD	N/A	Gastec tube #5Lb	0.01 ppm	Range: 0.05 – 10 ppm Volume: Pull 8 100-mL pump strokes	Variable ²
				MultiRAE Sensor AreaRAE Sensor	0.1 ppm	Range: 0.1 – 20 ppm	N/A
Nitrogen dioxide (NO ₂)	N/A	Report reading to PM/PTD	N/A	MultiRAE Sensor AreaRAE Sensor	0.1 ppm	Range: 0.1 – 20 ppm	N/A
				Gastec tube #9L	0.1 ppm	Range: 0.5 – 125 ppm Volume: Pull 2 100-mL pump strokes	Variable ²

¹ Readings collected using the SidePak AM510/AM520 or DustTrak are recorded as corrected concentrations. A Photometric Calibration Factor (PCF) of 0.38 is used, which TSI indicates is appropriate for ambient/fugitive emissions monitoring applications. Photometric instruments (e.g., SidePak AM520) will overstate the mass of an aerosol compared to its gravimetric weight due to humid conditions; PM_{2.5} is especially prone to interference from high humidity. In cases of high humidity, PM₁₀ impactors may be used, which are not as sensitive to humidity.

² Readings collected using Gastec tubes are recorded as uncorrected (raw) concentrations. The correction factors for Gastec tubes are dependent on volume of air used (i.e., number of pump strokes).

Abbreviations: PM_{2.5} = particulate matter with a diameter of 2.5 microns or less; PM₁₀ = particulate matter with a diameter of 10 microns or less; NA = not applicable; PM = Onterris Project Manager; PTD = Onterris Project Technical Director; mg/m³ = milligrams per cubic meter

Plan 3: Site Assessment – Flammability

Analyte	Action Level	Action to be Taken	Basis	Instrument	Detection Limit	Notes	Correction Factor
% LEL	N/A	Report reading to PM/PTD	N/A	MultiRAE Sensor	1 %	Range: 0 – 100 %	Variable ¹
				AreaRAE Sensor	1 %	Range: 0 – 100 %	Variable ¹

¹ Readings collected using the % LEL sensor are recorded as uncorrected (raw) concentrations. Correction factors for various compounds are listed in Honeywell Technical Note TN-156.

Abbreviations: % LEL = atmospheric flammability measured as a percentage of the lower explosive limit; PM = Onterris Project Manager; PTD = Onterris Project Technical Director

Air Sampling Methods - Community Monitoring

Analyte	Media/Can	Method	Notes
VOCs	1.4 L Evacuated Cannisters	USEPA TO-15	
Ammonia	Anasorb 747 (226-29)	Method 6015	LOD: ~ 0.5 µg
Metals	Mixed cellulose ester (MCE) filter (35-5,000 L; 1-4 LPM)	Mod. NIOSH 7303 (ICP/MS)	

Abbreviations: L = liters; USEPA = United States Environmental Protection Agency

Preliminary SF

General Information on Procedures (Assessment Techniques) Used

Procedure	Description
Handheld Real-Time Air Monitoring	Onterris personnel may utilize handheld real-time air monitoring instruments (e.g., MultiRAE, ChemLogic CLPx, Gastec colorimetric detector tubes) to measure airborne chemical concentrations in real time. Onterris personnel will use these handheld instruments primarily to monitor the ambient air quality at breathing zone level. Additionally, measurements may be made at grade level, as well as in elevated workspaces, as indicated by chemical properties or site conditions. Onterris personnel may also use these techniques to verify detections observed by the AreaRAE network.
Fixed Real-Time Air Monitoring Locations	Multiple locations may be identified and monitored at the same location approximately once per hour using handheld real-time air monitoring instruments. This allows the use of statistical analysis more effectively than with a random approach.
Analytical Air Sampling	Analytical air sampling may be used to validate the fixed and handheld real-time air monitoring data, or to provide data beyond the scope of real-time instruments. Analytical air samples may be collected as whole air samples in evacuated canisters or on specific collection media and sent to an off-site laboratory for further chemical analysis.
Particulate Monitoring Network	A network of data-logging particulate monitors may be set up and positioned in various locations.

Not all procedures listed in this table may be utilized.

Quality Assurance/Quality Control (QA/QC) Procedures

Method	Procedure
Real-Time Air Monitoring	All real-time air monitoring instruments will be calibrated per the manufacturer's recommendations, whenever indicated by site conditions or instrument readings, and/or at the start of every 12-hour shift. Serial numbers for all equipment used will be recorded. If Gastec tubes are used, the tube expiration date, lot number, and air volume (number of pump strokes) will be recorded.
Analytical Air Sampling	Chain-of-custody documentation will be completed for each air sample. Serial numbers for all analytical equipment used will be recorded. Co-located air samples may be conducted to assess field accuracy and precision (at least one co-located air sample per day or per 10 air samples). Level II data validation may be performed on all air samples. Level IV data validation may be performed on the first and last sample delivery groups (SDGs) and on at least 10% of all air samples. Level IV data validation may be performed on 10% of all samples.
Reporting	Daily data summaries may be provided for informational purposes using data that have not undergone complete QA/QC procedures. Comprehensive reports of real-time and/or analytical data may be generated following QA/QC and may be delivered 60 days following receipt of validated results, if applicable.

Glossary

Term	Definition
Sustained	Instrument reading above the action level continuously for the listed time period
Breathing Zone	The area within an approximate 10-inch radius of an individual's nose and mouth
Ambient Air	The portion of the atmosphere (indoor or outdoor) to which workers and the general public have access

Version Control

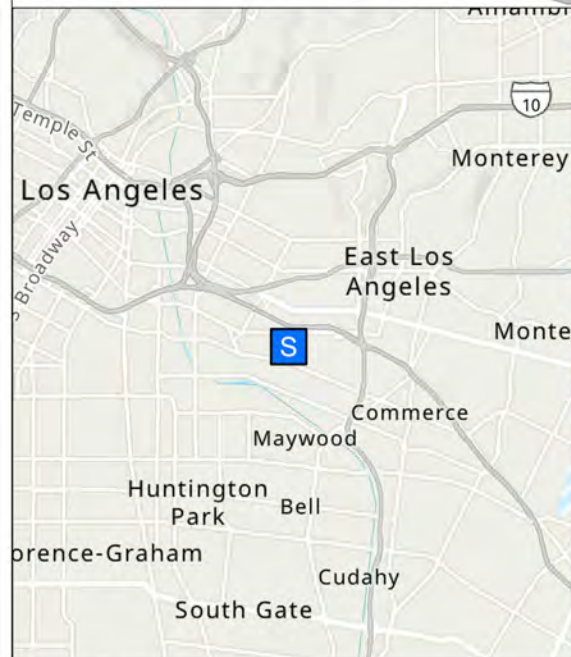
Version	Date	Author	Reviewer	Summary of Changes
1.0	6/21/26	Elysia Masters, PhD	Joseph Lucas, PhD	Initial Version
1.1	6/27/26	Max Ford, PhD		Added hydrogen sulfide to monitoring plan
1.2	7/6/26	Max Ford, PhD		Add mercaptan and chlorine monitoring to plan

Appendix B Incident Site and Real-Time Air Monitoring Locations

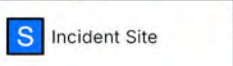


Los Palos Warehouse Fire

Incident Location
Los Angeles, CA | Los Angeles County
PROJ-069775



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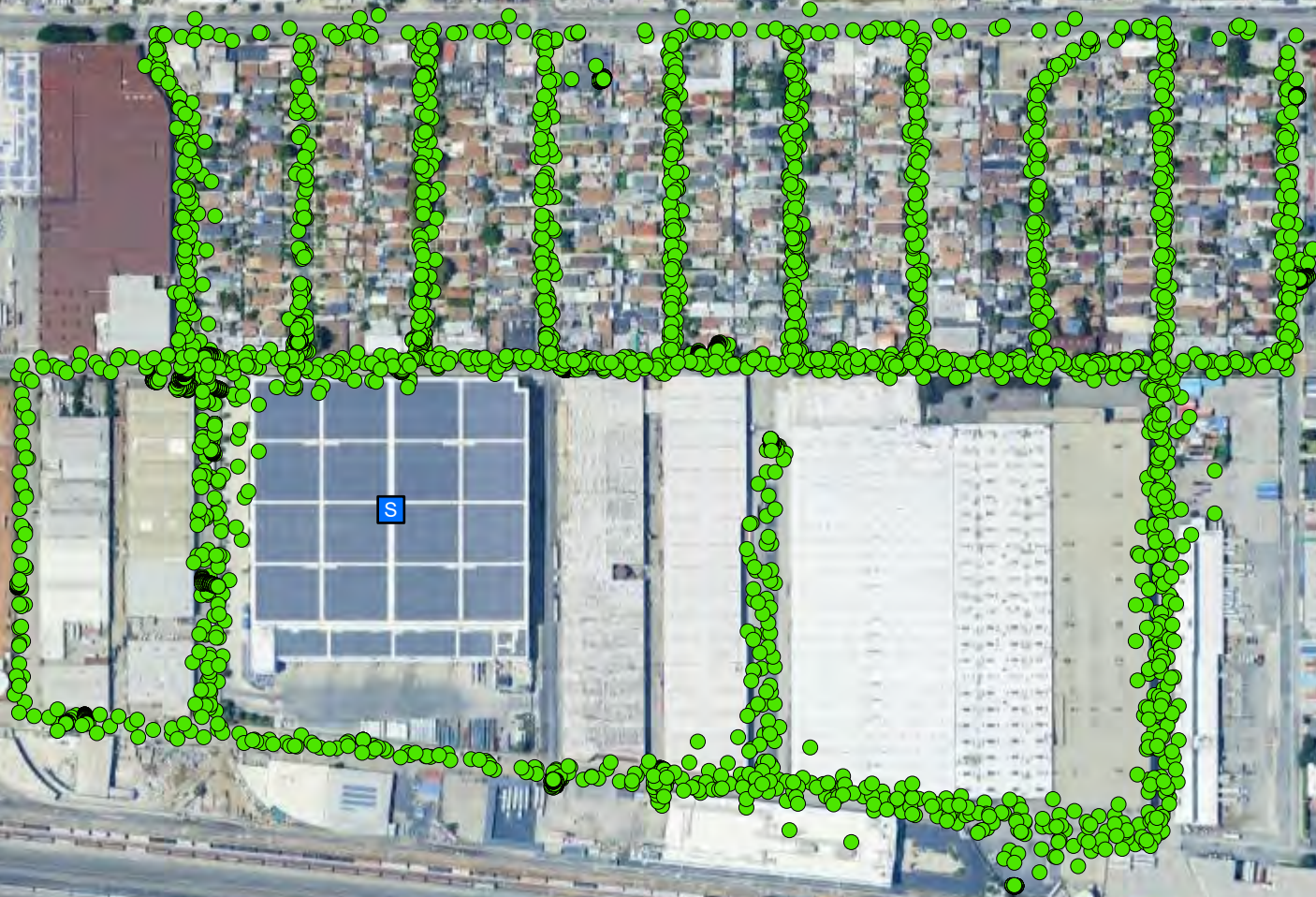




H₂S | AreaRAE

S Incident Location

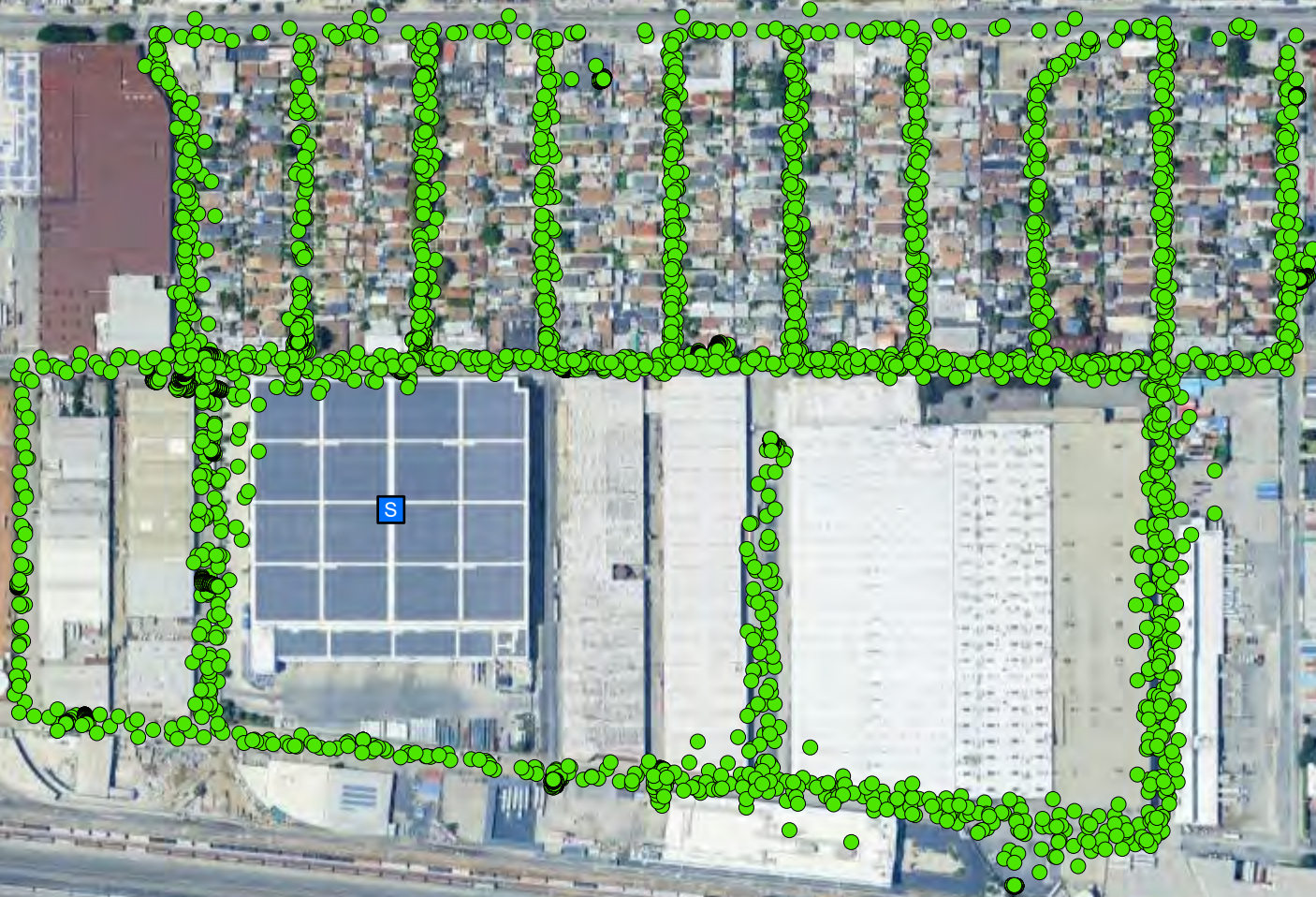
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%LEL | AreaRAE

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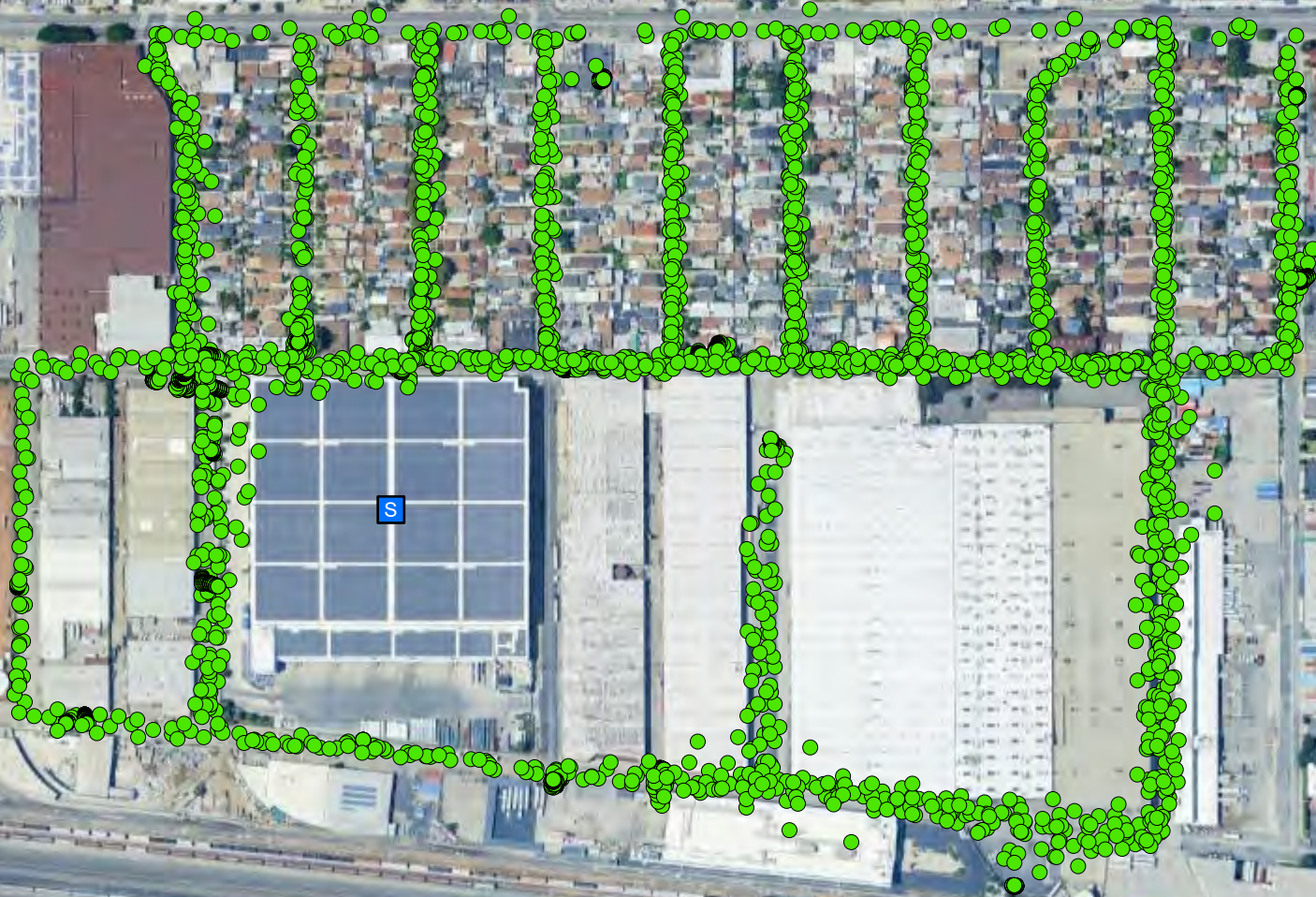
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NH₃ | AreaRAE

 Incident Location

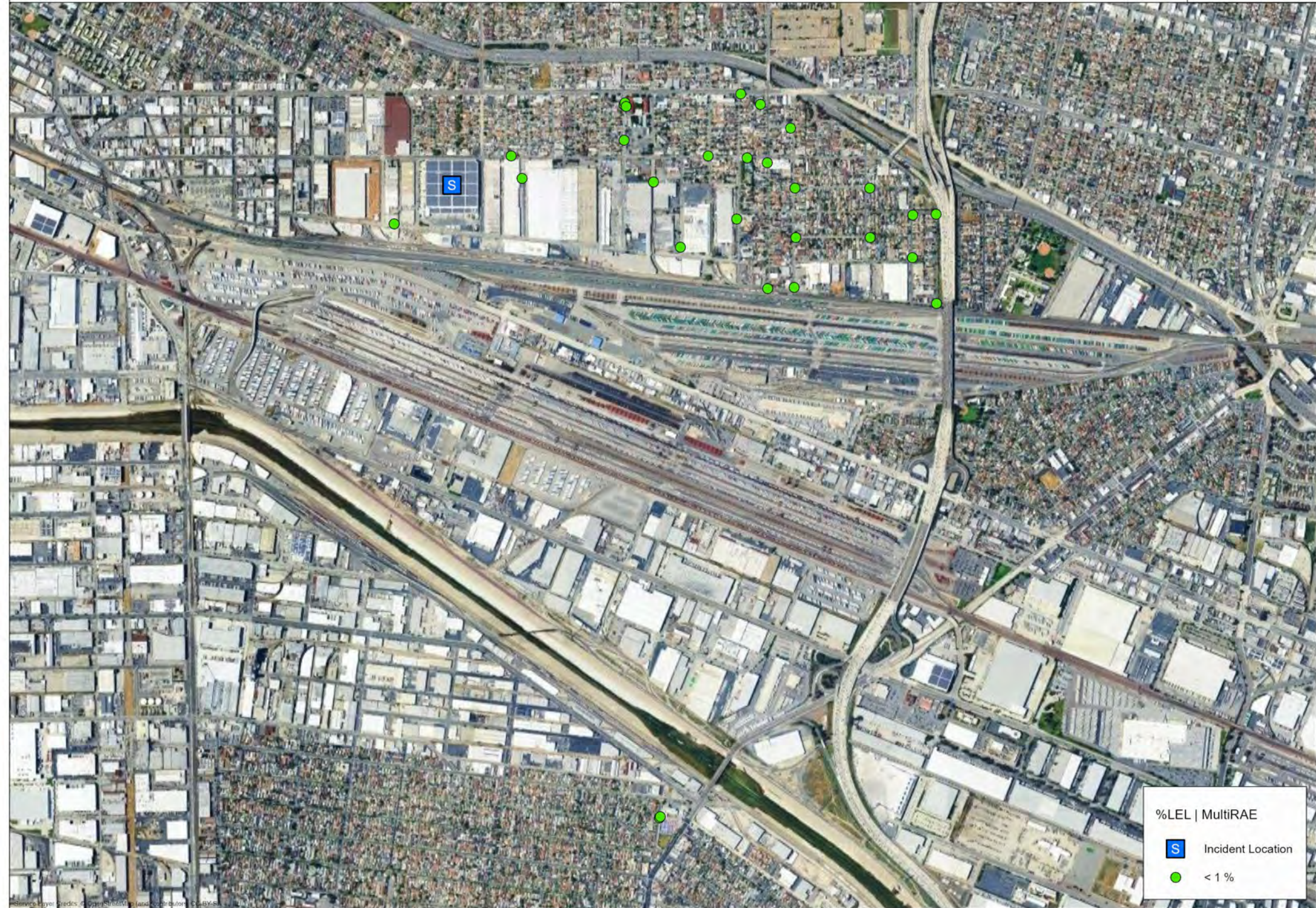
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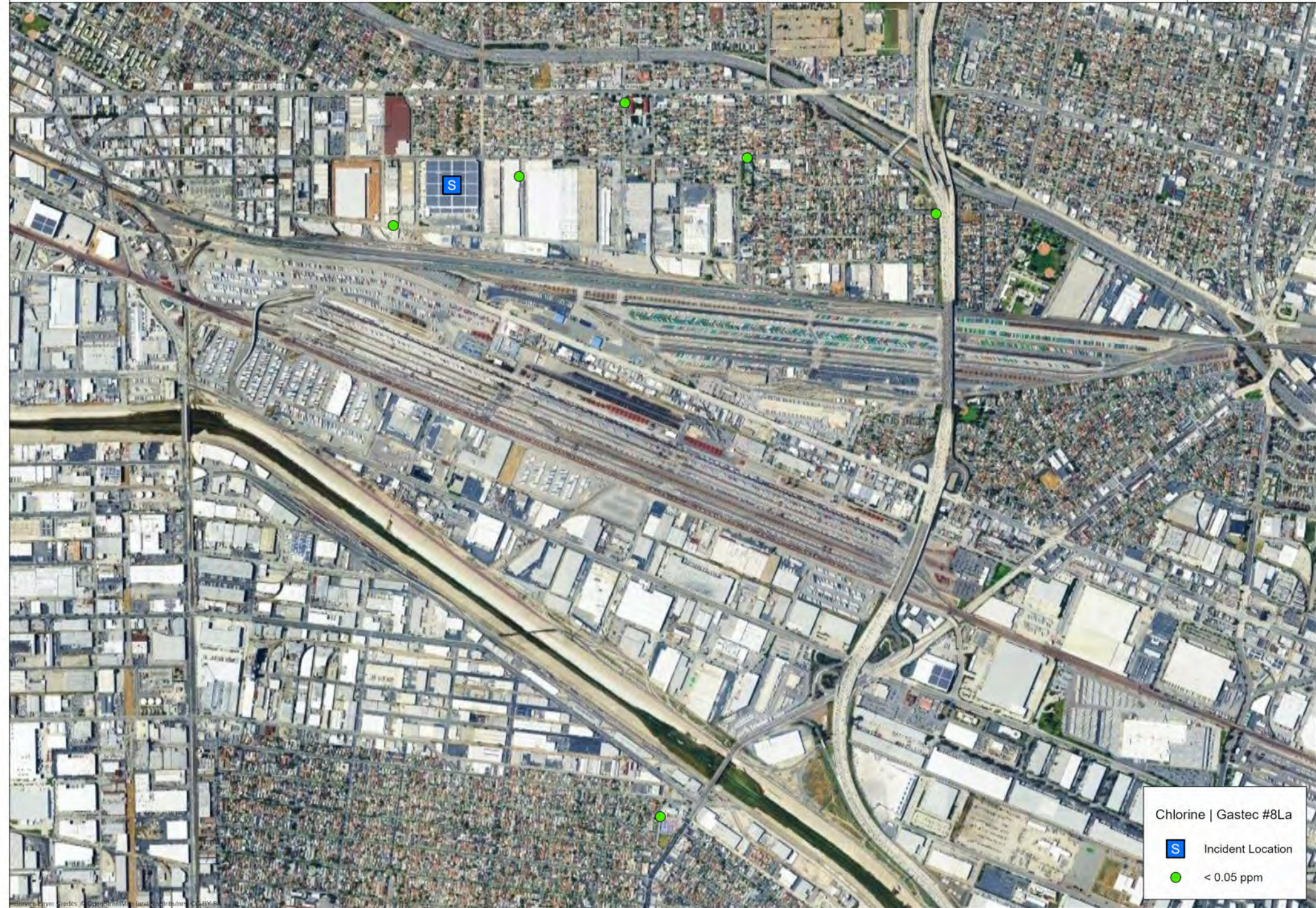


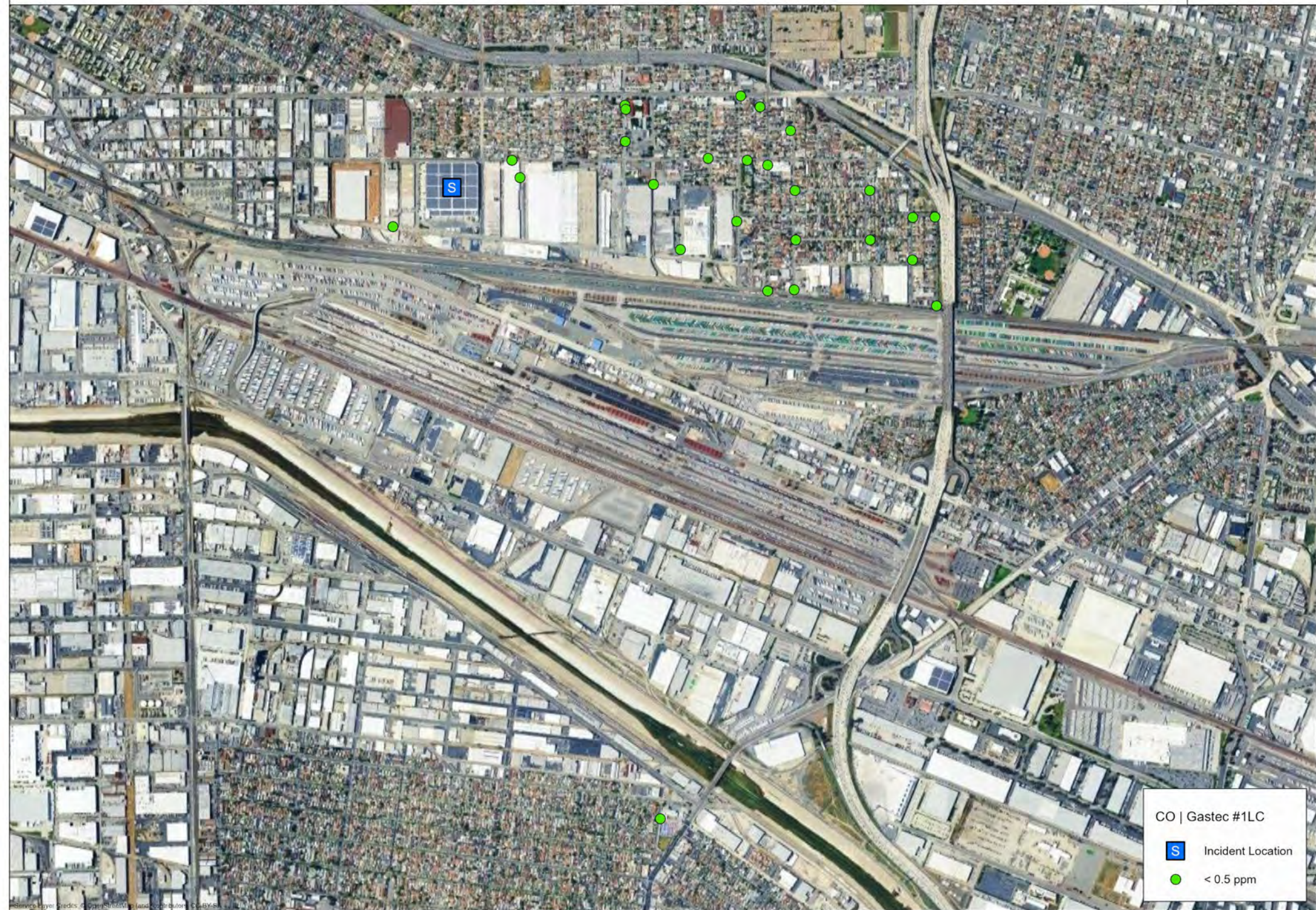
VOCs | AreaRAE

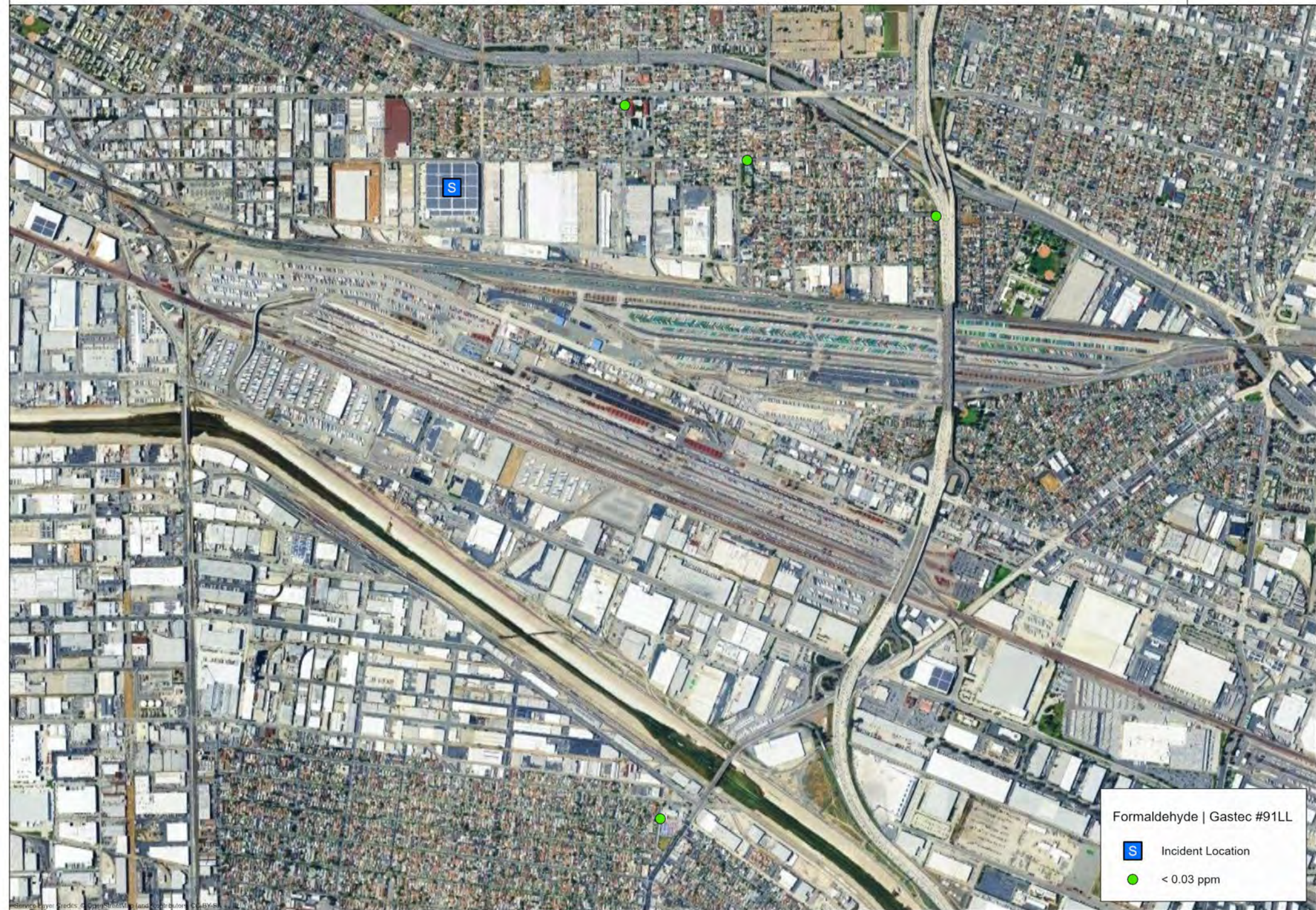
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 < 0.1 ppm









Formaldehyde | Gastec #91LL

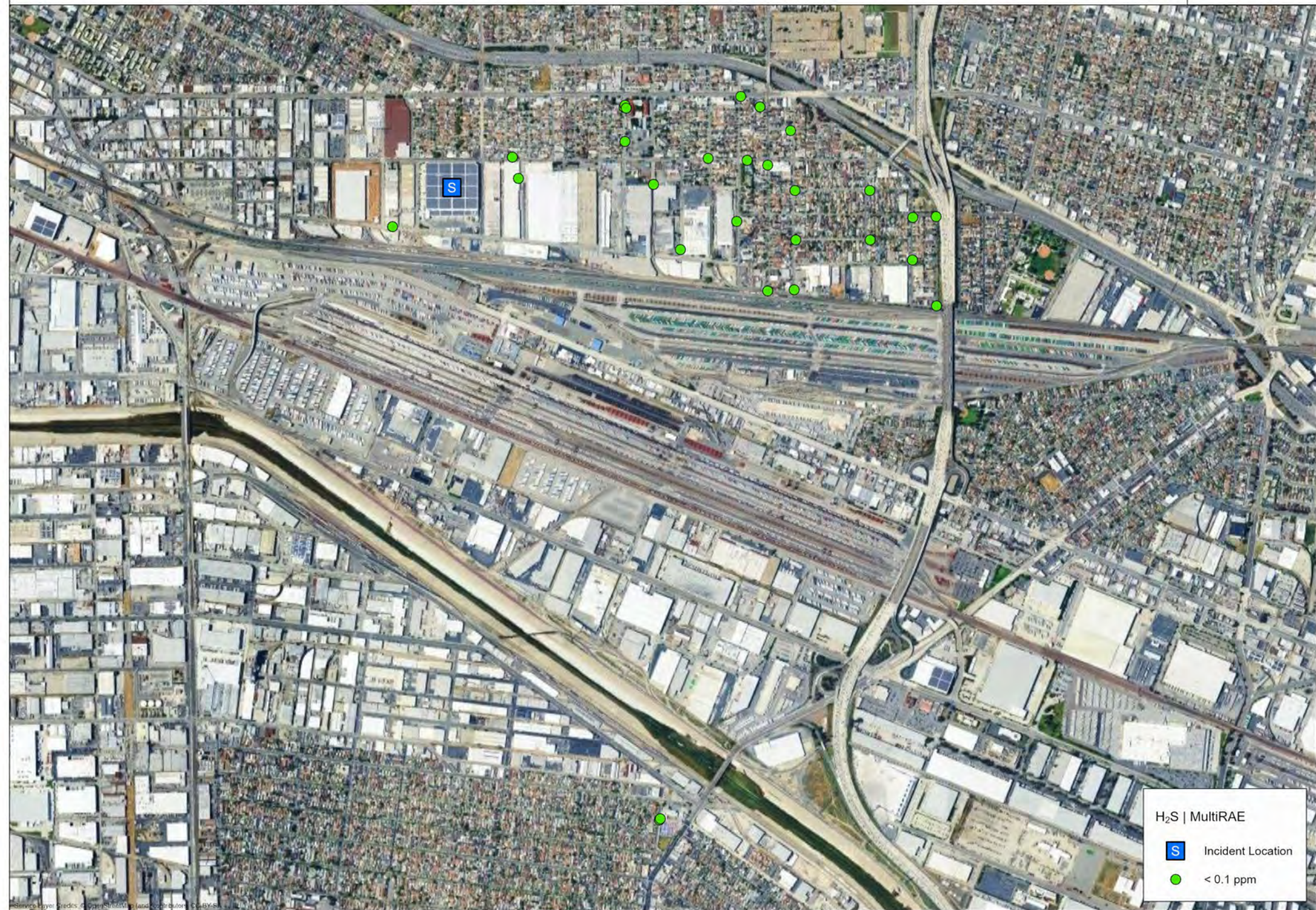
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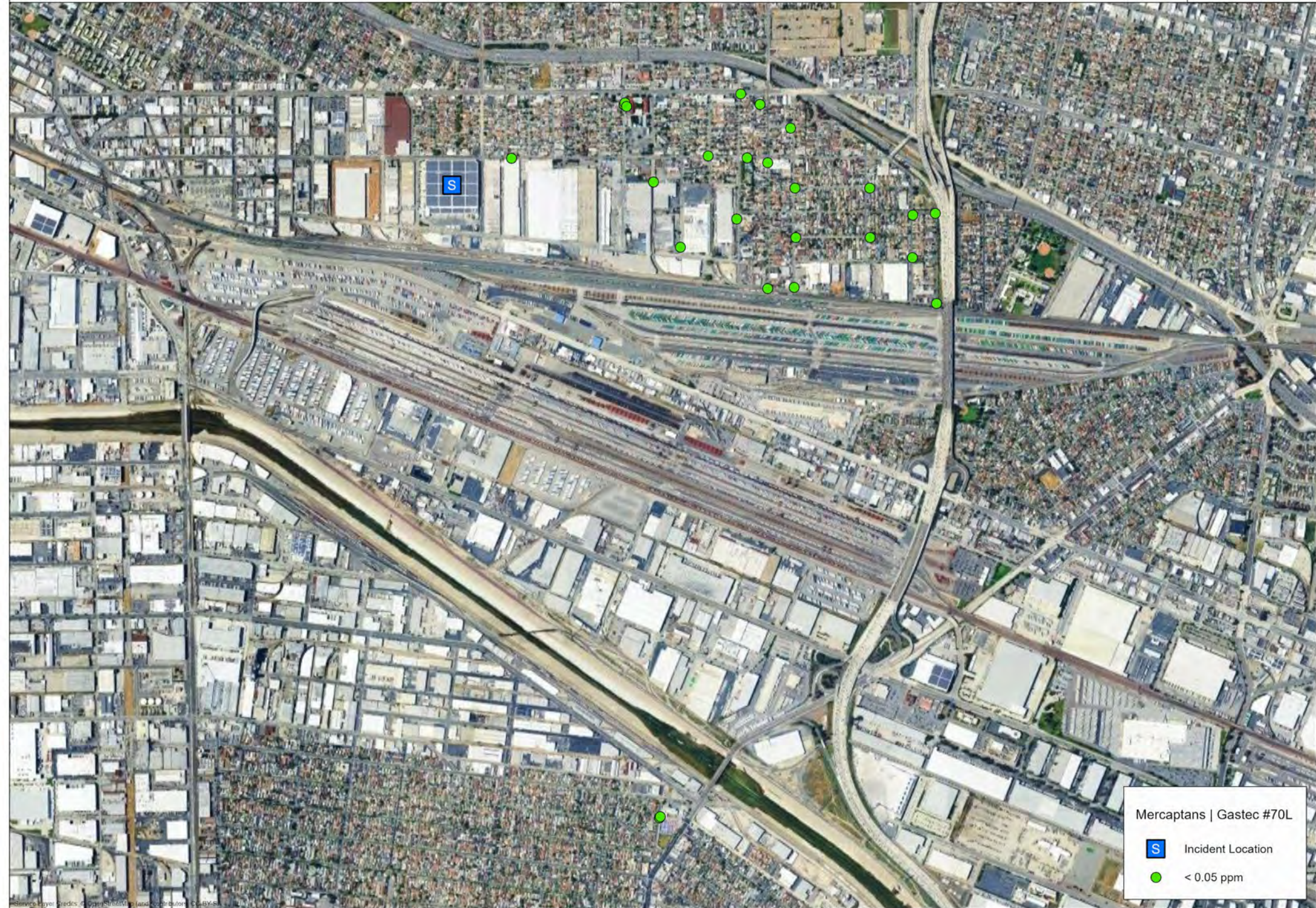
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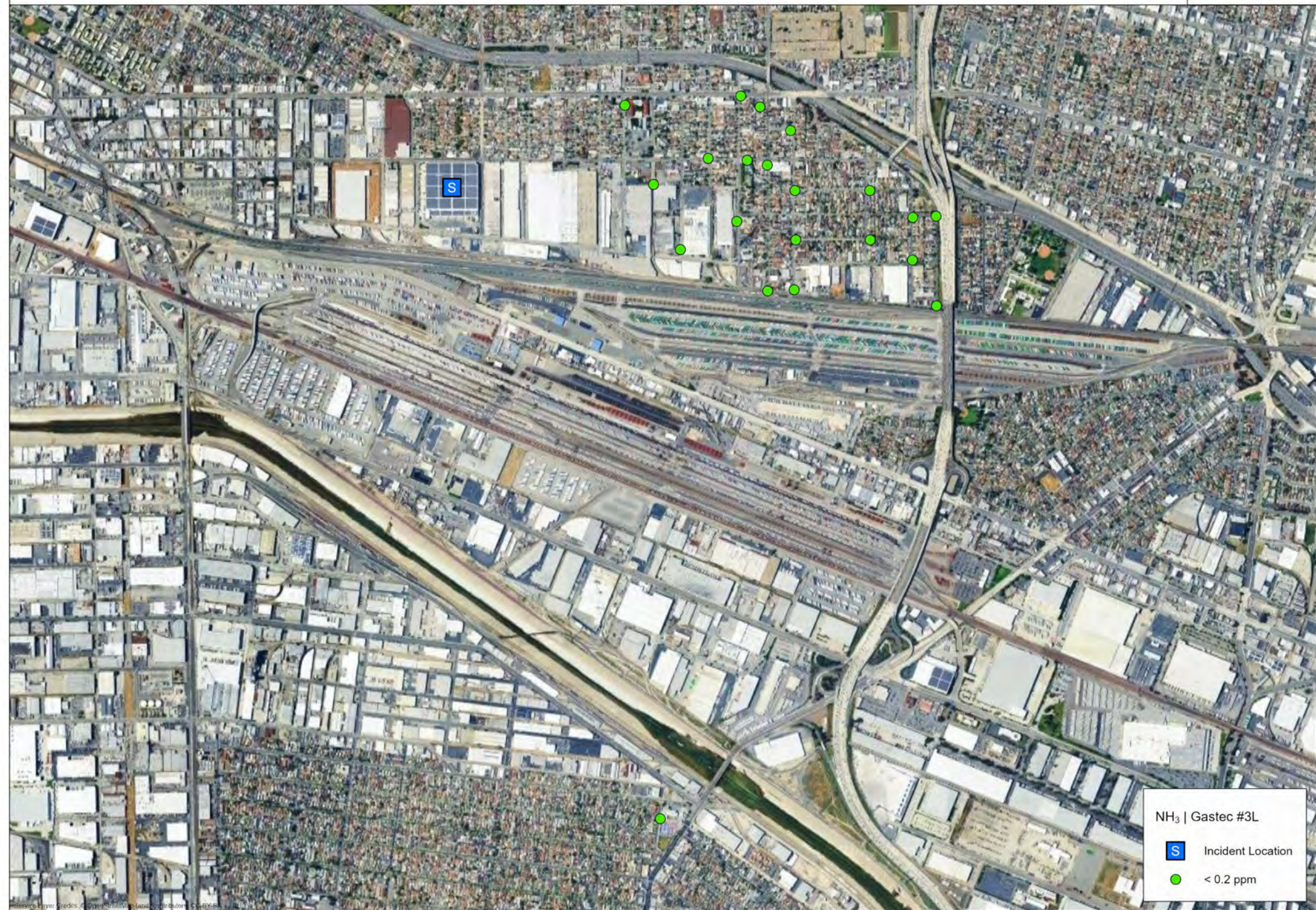
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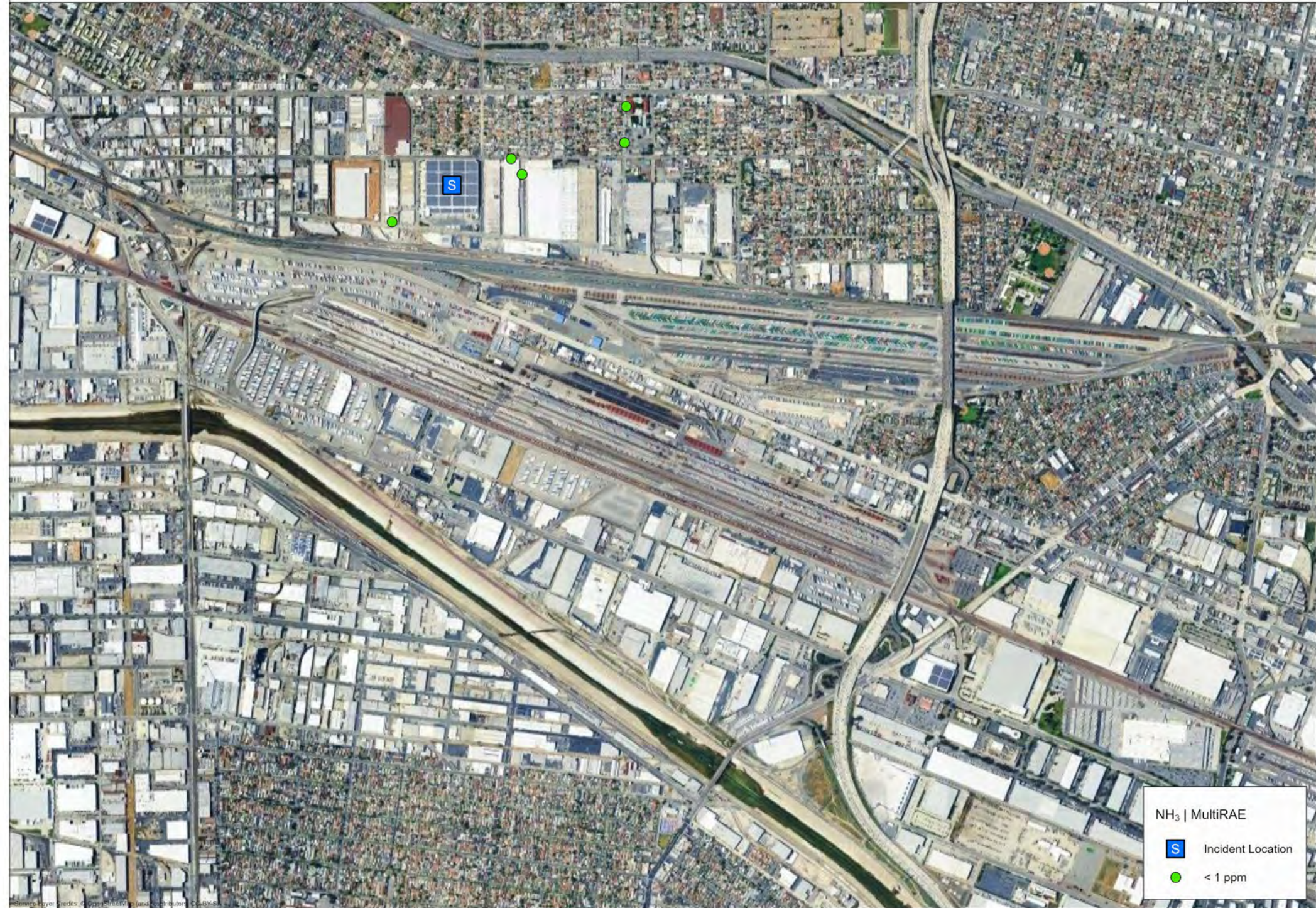


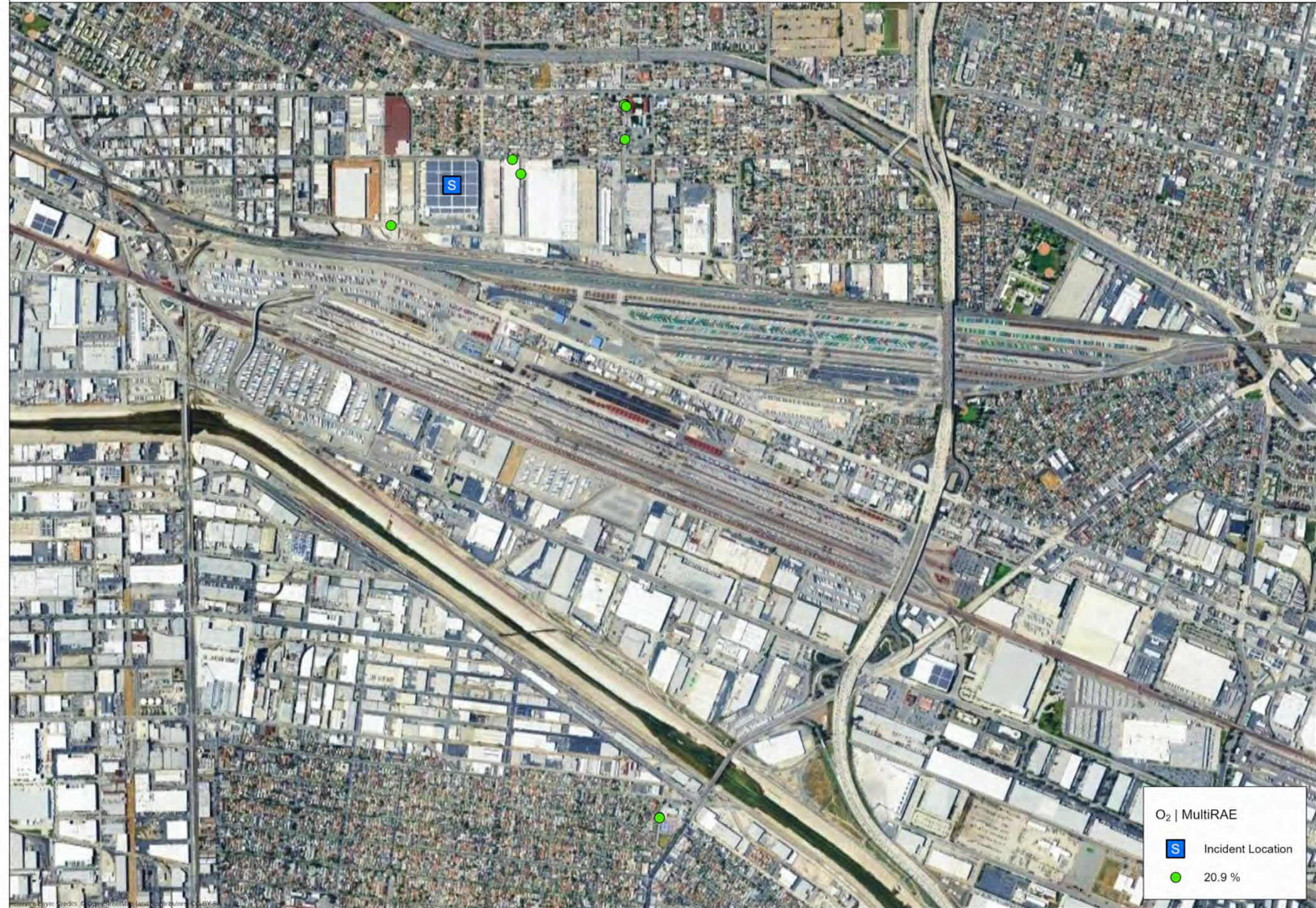
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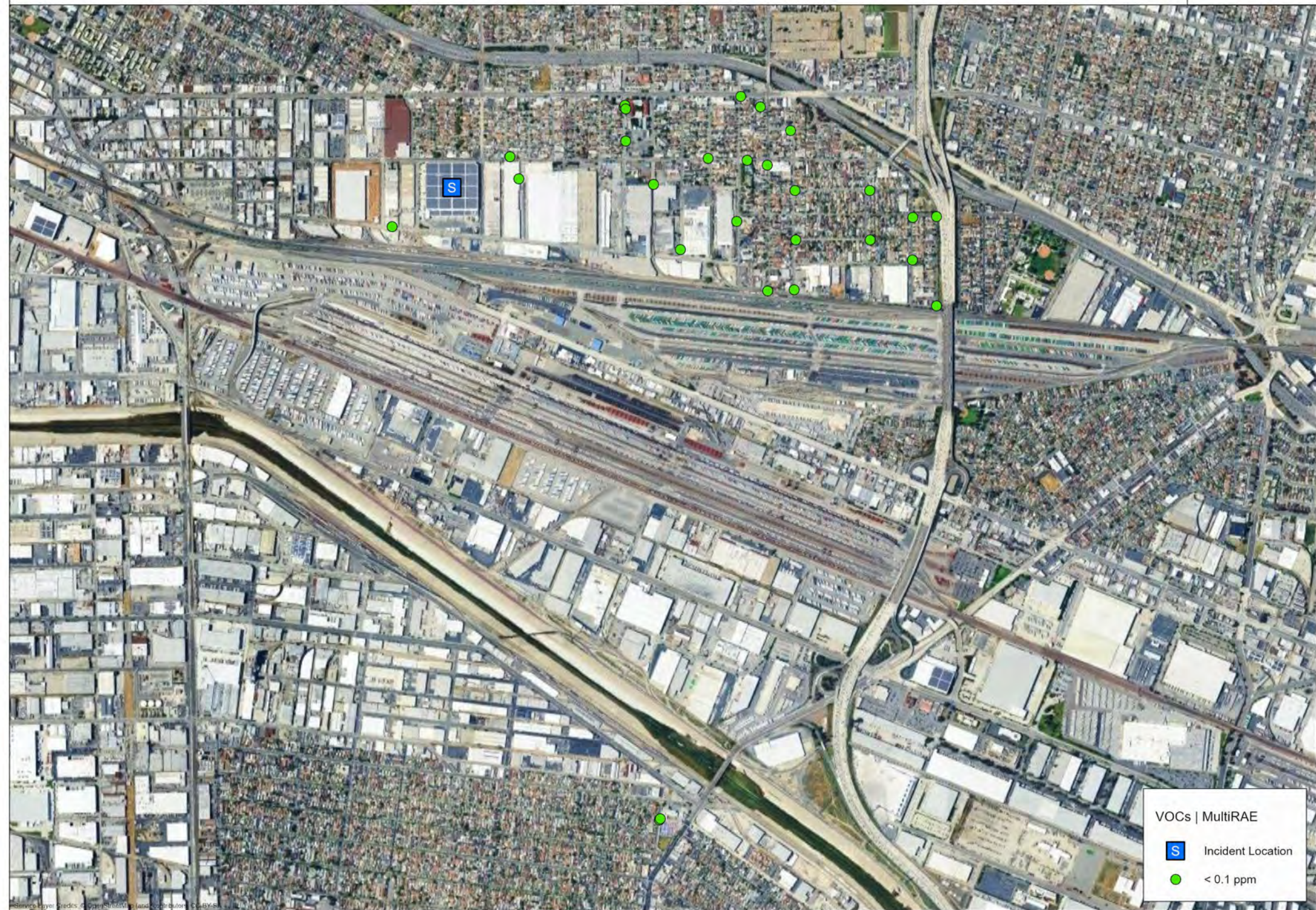








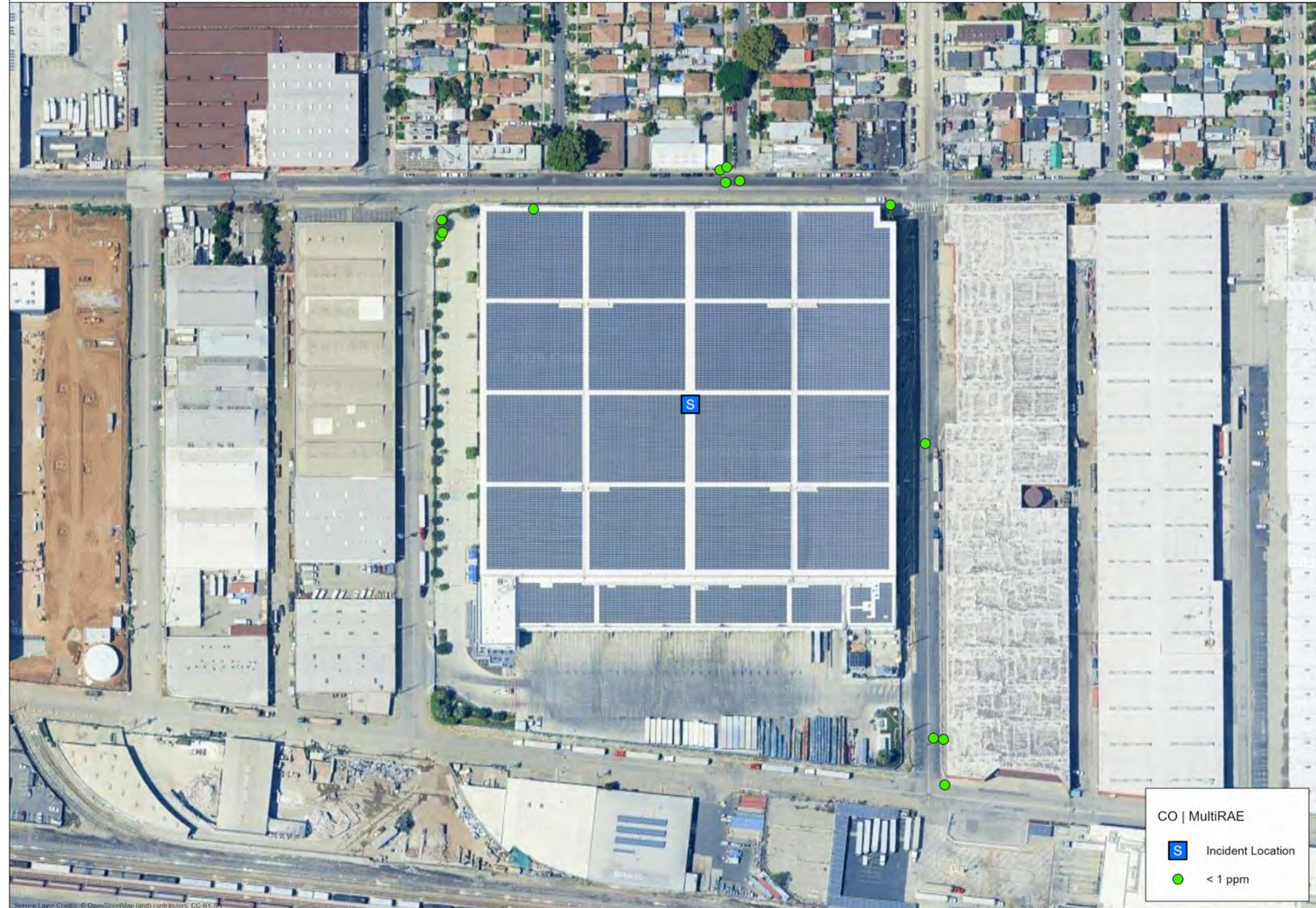








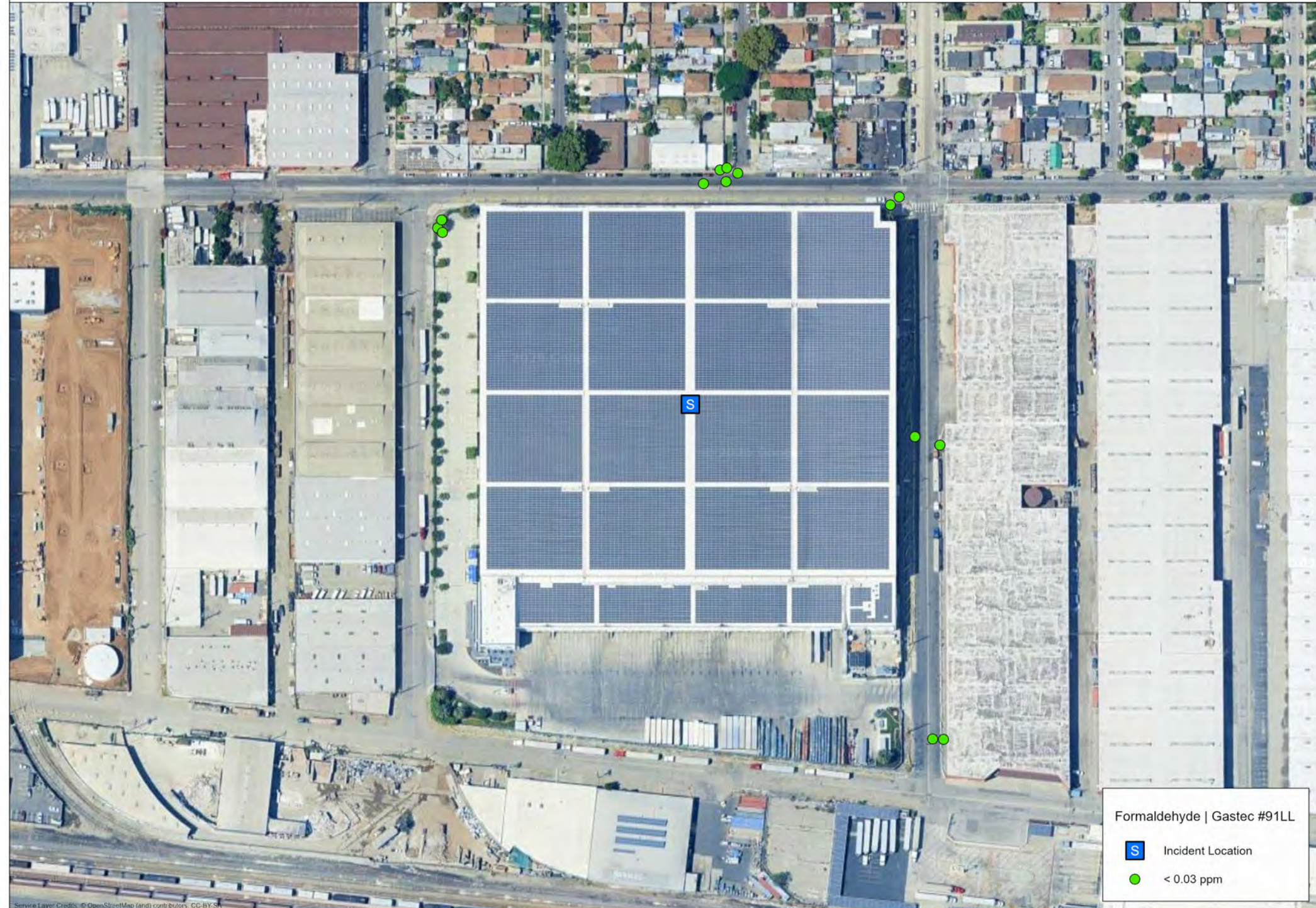




CO | MultiRAE

S Incident Location

● < 1 ppm



Formaldehyde | Gastec #91LL

S Incident Location

● < 0.03 ppm

0 50 100 Feet



Updated at 2026-08-27 10:53



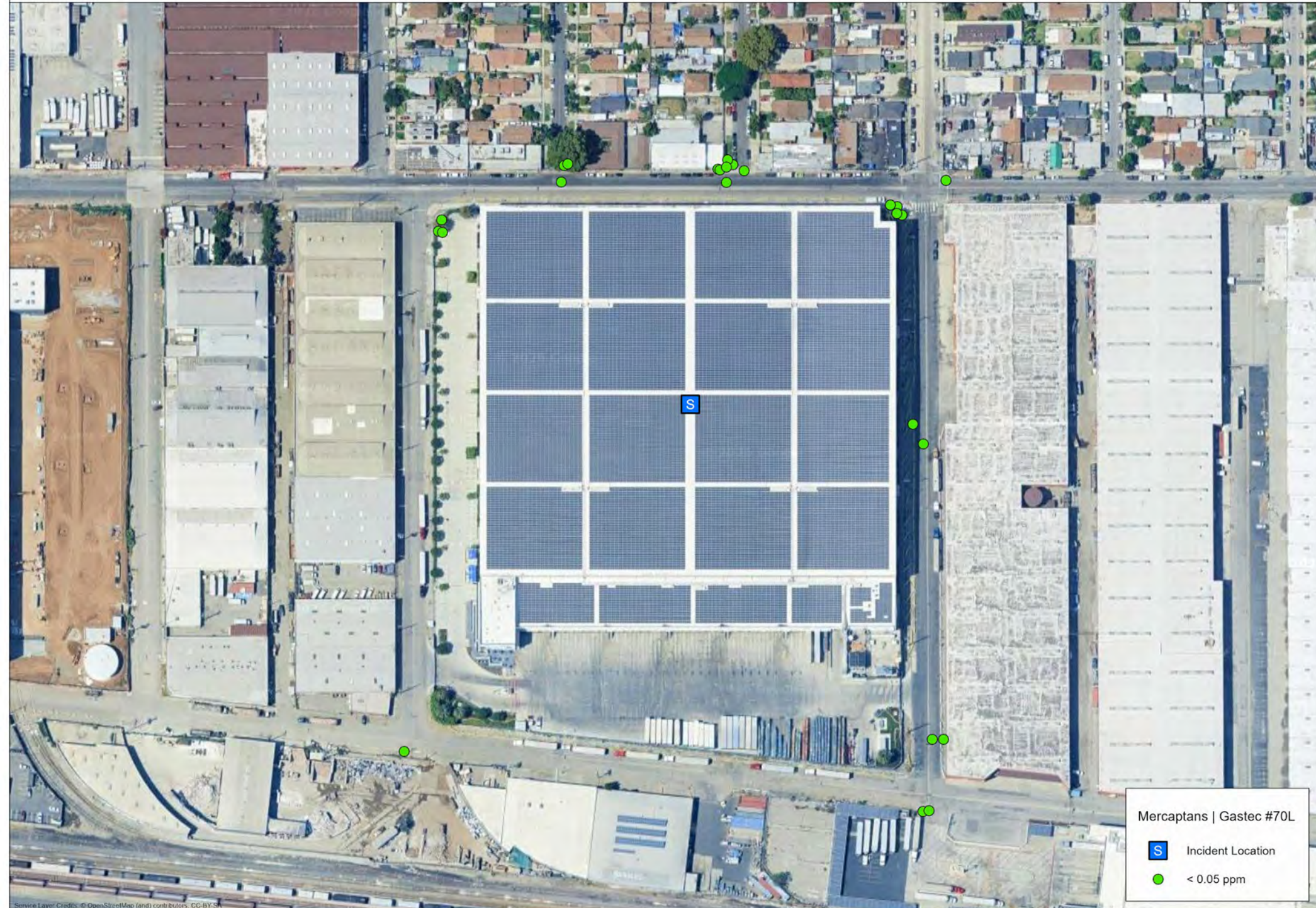
Service Layer Credits: © OpenStreetMap (and) contributors, CC-BY-SA

Datum: NAD 1983 2011
Projection: NAD 1983 2011 StatePlane California V FIPS 0405 Ft US

0 50 100 Feet



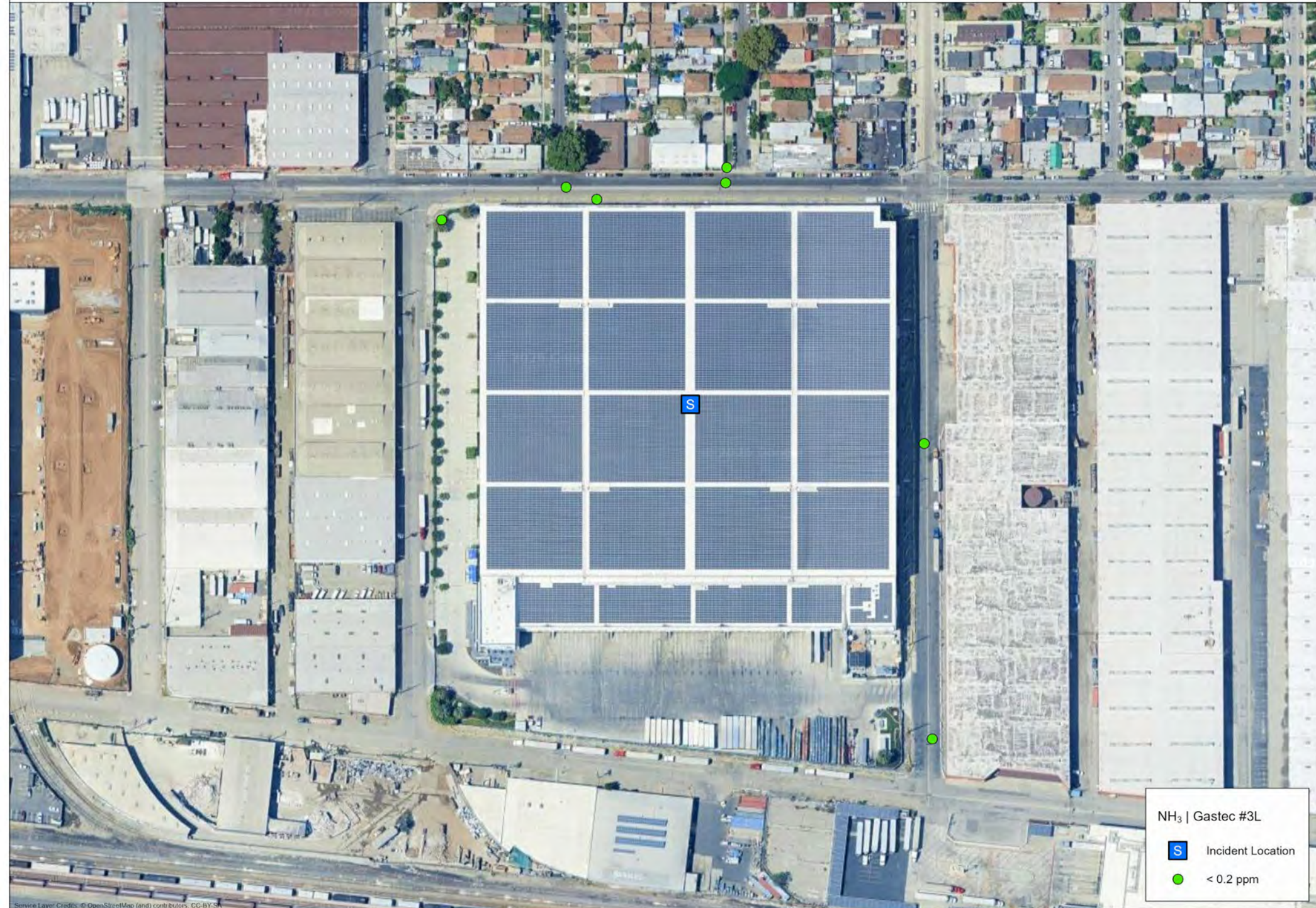
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Mercaptans | Gastec #70L

S Incident Location

● < 0.05 ppm



NH₃ | Gastec #3L

S Incident Location

● < 0.2 ppm



NH₃ | MultiRAE

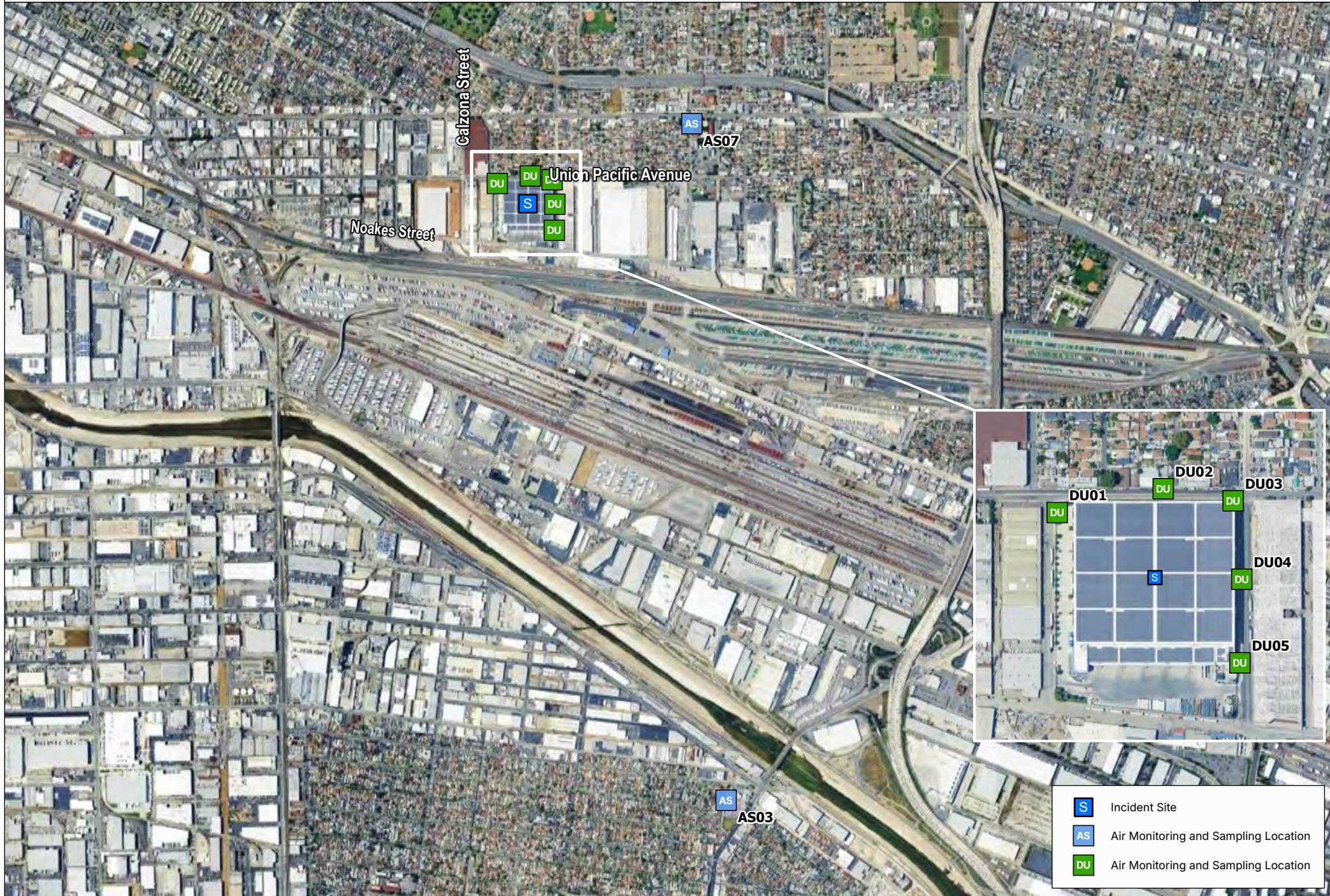
S Incident Location

● < 1 ppm





Appendix C Stationary Air Monitoring and Analytical Sampling Locations



- S Incident Site
- AS Air Monitoring and Sampling Location
- DU Air Monitoring and Sampling Location

Appendix D Meteorological Data

