

Introduction

- Ethylene Oxide (EtO) is a colorless, flammable gas at room temperature, produced by the catalytic oxidation of ethylene^[1]
- EtO has alkylating properties used to sterilize medical devices and equipment^[2]
- In 2016, epidemiological studies led the EPA to update the inhalation hazard unit for EtO due to more potent carcinogenic findings
- Chronic EtO exposure is associated with an increased risk of white blood cells related cancers, including non-Hodgkin lymphoma, myeloma, and leukemia^[3]
- The Salt Lake City geography and meteorological conditions may prevent EtO from being diluted in the ambient air after emitted
- Background environmental sources of EtO are unknown, but the EPA proposed automobile combustion and biological metabolic by-product as possible source emitters^[4]

Purpose

- To investigate how the method of measuring ambient EtO using the EPA's dispersion modeling techniques (AERMOD) compares to canister-based measurements
- If a discrepancy is found, a correction factor could be derived to aid in subsequent cancer risk assessment of the nearby community using human exposure modeling (HEM-4)

Methods

- A local sterilization facility was identified as point source of ambient EtO emission in Salt Lake City
 - Fugitive and stack emission information was obtained
- New, whole-air passive canisters lined with silonite collected 24-hour samples of ambient EtO concentrations
 - 8 sites near the sterilization facility
 - 5 background sites
- Lab analyzed with the EPA's TO-15 Method using gas chromatography and mass spectrometry techniques
- Summer and Winter background values were computed and added to modeled estimates using Mean-of-the-Median values calculated from EtO concentrations measured at background canister sites

Methods

- The American Meteorological Society – U.S. EPA Regulatory Model (AERMOD) used dispersion modeling to estimate EtO concentration outputs from the point sources' stack and fugitive emissions
- Measured and estimated concentrations of ambient EtO were compared with a Wilcoxon Signed-ranked test for paired, non-parametric datasets
 - 16 one-sided tests to determine directionality

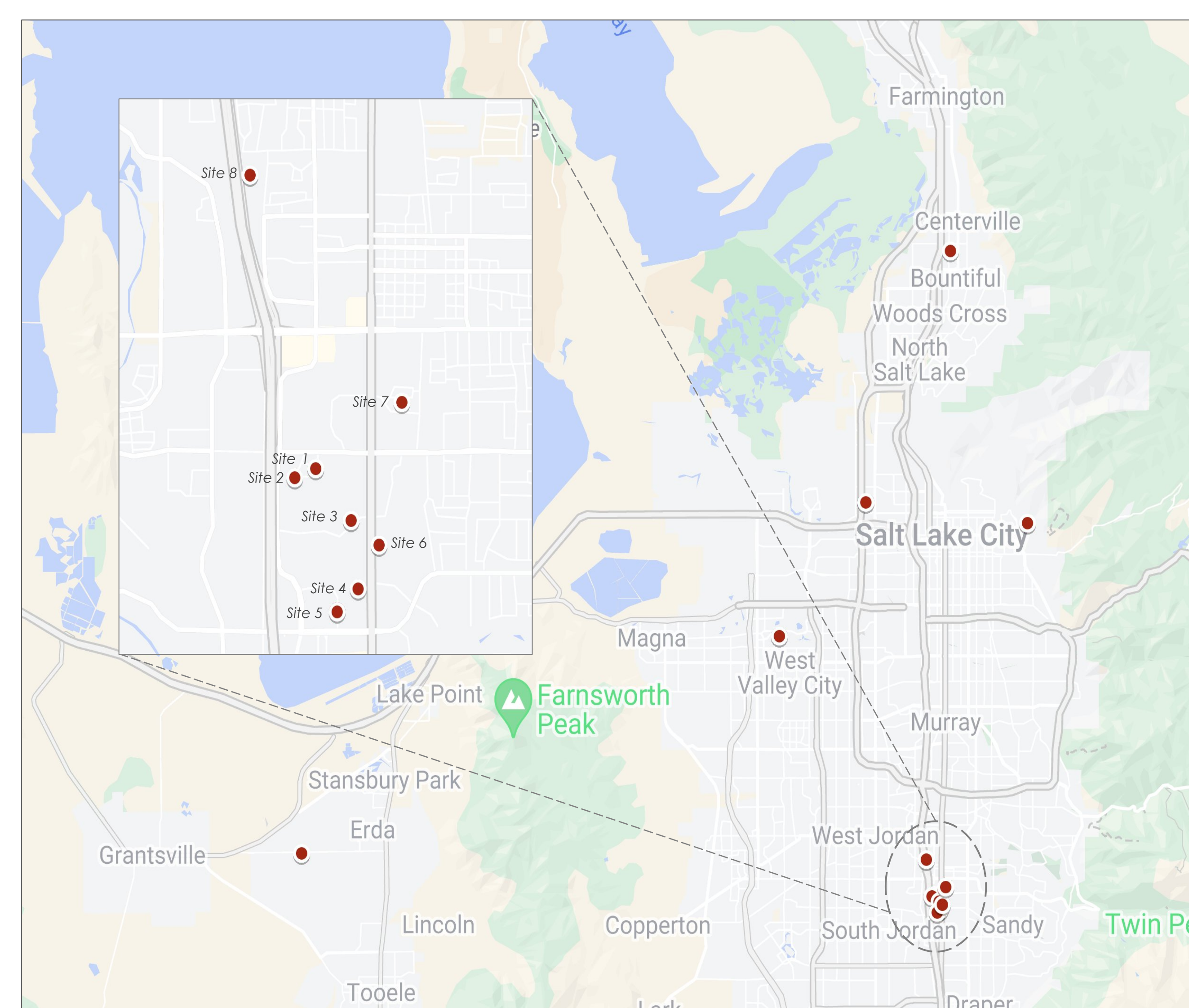


Figure 1. Map showing where the canister-based measurements were taken across Utah's Salt Lake City. There are 8 sampling sites near the medical sterilization facility (see blown up image) and 5 background sites locations.

Results

- The background EtO value was approximately 2.5x greater for measured concentrations in summer compared to winter
- Background site 3, a residentially and industrially mixed location, showed the greatest EtO median concentration for both seasons

Table 1. Background site medians and Mean-of-median values for winter and summer

Site	n	Winter (µg/m ³)		Summer (µg/m ³)		
		Median	Mean-of-medians	n	Median	Mean-of-Medians
1	19	0.0575		19	0.2015	
2	18	0.0621		14	0.1543	
3	19	0.1070	0.0779	15	0.2254	0.1962
4	16	0.0958		19	0.1987	
5	15	0.0667		0		

Results

- As the sites increase in distance from the point source, the median EtO concentrations generally decreased
- Of the 8 sites sampled around the point source, 6 of the AERMOD estimates were statistically different from the canister-based measurements
 - AERMOD estimates > canister measurements at sites 1, 3, and 4
 - AERMOD estimates < canister measurements at sites 2, 5, and 7
 - Sites 6 and 8 were not statistically significant in either direction

Table 2. Wilcoxon Signed-ranked test results from comparing median concentrations of EtO estimated with AERMOD modeled to (less than or greater than) canister-based measurements. Significant p-values are noted.

Site	n	Modeled Estimates	Canister Measurements	Wilcoxon p-values	
		Median (µg/m ³)	Median (µg/m ³)	Less than	Greater than
1	29	1.0195	0.5296	0.99	< 0.001*
2	38	0.2343	0.2765	0.03*	0.97
3	32	0.3420	0.2812	0.99	0.01*
4	35	0.2131	0.1295	0.99	< 0.01*
5	37	0.2023	0.1855	0.01*	0.99
6	36	0.2211	0.1919	0.22	0.78
7	37	0.1348	0.1665	0.05*	0.95
8	34	0.1540	0.1536	0.24	0.77

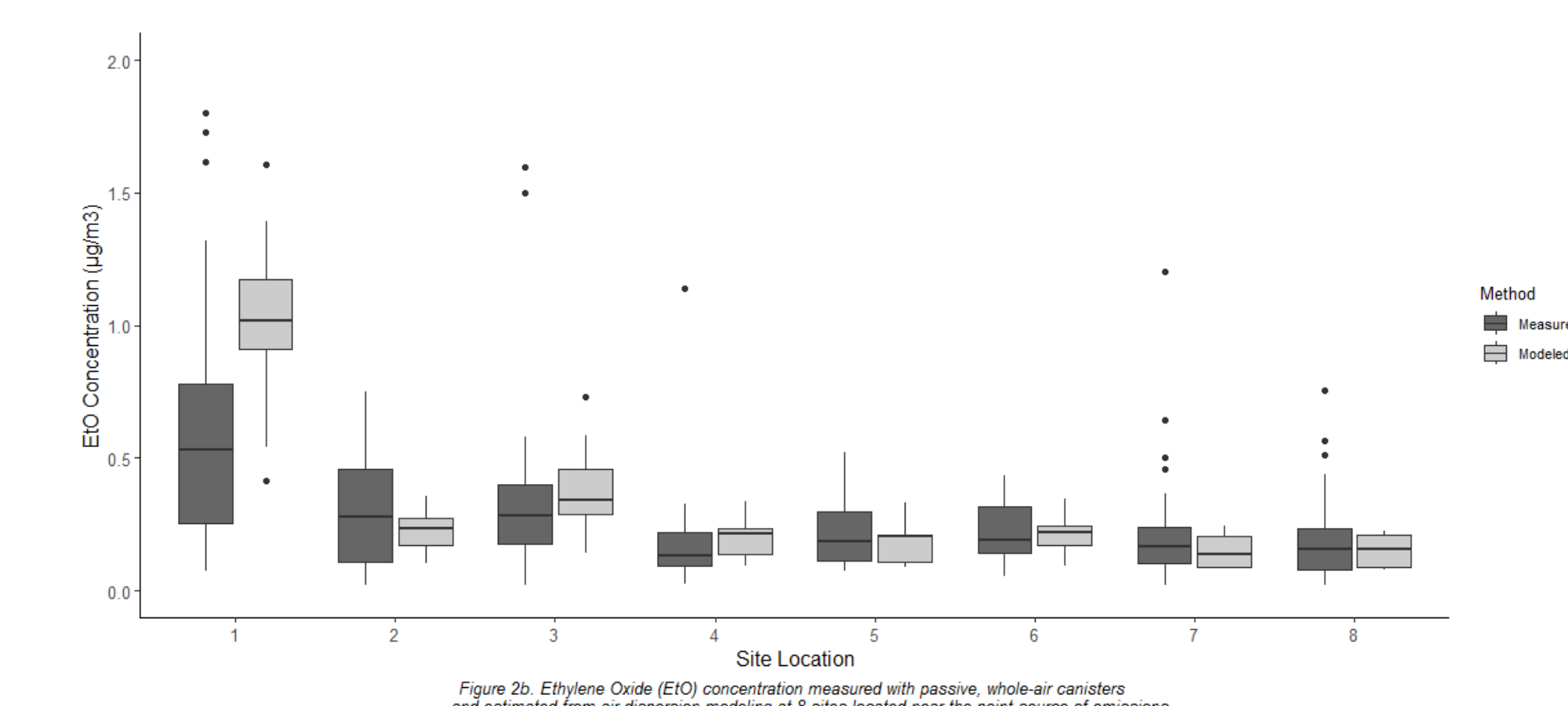


Figure 2. Ethylene Oxide (EtO) concentration measured with passive, whole-air canisters and estimated from air dispersion modeling at 8 sites located near the point source of emissions.

Conclusion

- After comparing the estimated EtO concentrations from dispersion modeling to canister-based measurements, 5 of the closest sites, and site 7, were statistically different.
- Because neither method indicated being more conservative than the other, a correction factor could not be derived for future cancer risk assessment modeling with the EPA's human exposure modeling
- A vast seasonal difference was found between background data, which deserves further investigation

Acknowledgments

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References

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