## ARTICLE OPEN



# Comparison of the rate of healthcare encounters for influenza from source-specific $PM_{2.5}$ before and after tier 3 vehicle standards in New York state

Daniel P. Croft Mark J. Utell<sup>1,2</sup>, Philip K. Hopke<sup>3,4</sup>, Han Liu<sup>5</sup>, Shao Lin<sup>6</sup>, Sally W. Thurston<sup>2,7</sup>, Sathvik Thandra<sup>6</sup>, Yunle Chen<sup>3</sup>, Md Rayhanul Islam<sup>3</sup>, Kelly Thevenet-Morrison<sup>3</sup>, Carl J. Johnston<sup>8</sup>, Tianming Zhao<sup>3</sup>, Catherine Yount<sup>3</sup> and David Q. Rich<sup>1,2,3</sup>

© The Author(s) 2024

**BACKGROUND:** Influenza healthcare encounters in adults associated with specific sources of  $PM_{2.5}$  is an area of active research. **OBJECTIVE:** Following 2017 legislation requiring reductions in emissions from light-duty vehicles, we hypothesized a reduced rate of influenza healthcare encounters would be associated with concentrations of  $PM_{2.5}$  from traffic sources in the early implementation period of this regulation (2017–2019).

**METHODS:** We used the Statewide Planning and Research Cooperative System (SPARCS) to study adult patients hospitalized (N = 5328) or treated in the emergency department (N = 18,247) for influenza in New York State. Using a modified case-crossover design, we estimated the excess rate (ER) of influenza hospitalizations and emergency department visits associated with interquartile range increases in source-specific PM<sub>2.5</sub> concentrations (e.g., spark-ignition emissions [GAS], biomass burning [BB], diesel [DIE]) in lag day(s) 0, 0–3 and 0–6. We then evaluated whether ERs differed after Tier 3 implementation (2017–2019) compared to the period prior to implementation (2014–2016).

**RESULTS:** Each interquartile range increase in DIE in lag days 0–6 was associated with a 21.3% increased rate of influenza hospitalization (95% CI: 6.9, 37.6) in the 2014–2016 period, and a 6.3% decreased rate (95% CI: —12.7, 0.5) in the 2017–2019 period. The GAS/influenza excess rates were larger in the 2017–2019 period than the 2014–2016 period for emergency department visits. We also observed a larger ER associated with increased BB in the 2017–2019 period compared to the 2014–2016 period. **IMPACT STATEMENT:** We present an accountability study on the impact of the early implementation period of the Tier 3 vehicle emission standards on the association between specific sources of PM<sub>2.5</sub> air pollution on influenza healthcare encounters in New York State. We found that the association between gasoline emissions and influenza healthcare encounters did not lessen in magnitude between periods, possibly because the emissions standards were not yet fully implemented. The reduction in the rates of influenza healthcare encounters associated with diesel emissions may be reflective of past policies to reduce the toxicity of diesel emissions. Accountability studies can help policy makers and environmental scientists better understand the timing of pollution changes and associated health effects.

**KEYWORDS:** Source-specific PM<sub>2.5</sub>; Influenza; Traffic related air pollution; Accountability study; Air pollution; Biomass burning *Journal of Exposure Science & Environmental Epidemiology* (2025) 35:205–213; https://doi.org/10.1038/s41370-024-00710-w

### **Graphical Abstract**

# Diesel Lag 0-5 Diesel Lag 0-5 Diesel Lag 0-6 Diesel Lag 0-6 Diesel Lag 0-7 Diesel Lag 0-7 Diesel Lag 0-8 Diesel

Excess rate (%) of influenza hospitalization

**Before and After Tier 3 Vehicle Emission Standards** 

Received: 20 December 2023 Revised: 24 July 2024 Accepted: 26 July 2024

Published online: 10 August 2024

### INTRODUCTION

Influenza continues to be a concerning cause of adult morbidity and mortality each year in the United States since it represents the 13<sup>th</sup> highest cause of death [1]. Although influenza rates were much lower during COVID-19, influenza remains a relevant seasonal respiratory infection as influenza rates have returned to pre-COVID rates after the acute phase of the COVID pandemic [2]. While the association between influenza and increased concentrations of PM25 has been reported by several groups previously [3-6], the association between constituent sources of PM<sub>2.5</sub> and influenza is an area of active research. In our prior accountability study in New York State, we observed an increase in the relative rate of emergency department visits for influenza associated with an increased concentration of gasoline and diesel PM<sub>2.5</sub> concentrations. This potential increase in the toxicity per unit mass in the setting of fuel regulations and decreasing particulate pollution highlighted the need for ongoing evaluation of complex health responses to atmospheric exposures [7].

As described previously [3, 8], a series of energy and vehicle related regulations were enacted including spanning coal-fired power plant closures, sulfur reduction in diesel for trucks, and now the more recent 2017 implementation of the tier 3 vehicle standards [9]. These Tier 3 standards, will reduce the average sulfur content of gasoline from 30 parts per million (ppm) to 10 ppm and also when fully phased in, will lead to a 70% reduction in the per-vehicle particulate matter standard for light to medium duty vehicle tailpipe emissions [10]. The updated Non-methane Organic Gases (NMOG) and NOx tailpipe emissions were set at 30 milligrams per mile (mg/mi) for model year 2025. Overall, the Tier 3 standards will match the California's Super Ultra Low Emission Vehicle (SULEV) program so that the same cars can be sold in all 50 states in the U.S. [10].

In this study we again examined associations between influenza hospitalizations and emergency department visits associated with source-specific PM<sub>2.5</sub> concentrations, specifically comparing these associations before and after Tier 3 vehicle emissions implementation on January 1, 2017.

While many PM<sub>2,5</sub> constituents decreased from 2005 to 2013, secondary organic carbon (SOC) increased in the 2014-2016 period and has remained at relatively constant concentrations in the 2017–2019 period across all sites in New York State (NYS) [11]. Therefore, we also examined two forms of organic carbon, primary organic carbon (POC) and SOC. POC is directly emitted by a source and is commonly a combustion product while SOC is an atmospheric oxidation product of volatile organic compounds. The oxidative potential (ability to generate oxidative stress in the body) of SOC is greater than POC [12, 13]. The majority of PM<sub>2.5</sub> is a combination of secondary inorganic aerosol (SIA; secondary nitrate [SN] and secondary sulfate [SS]) and secondary organic aerosols (SOA; e.g. secondary organic carbon). The fact that SOC concentrations did not fall in the 2017-2019 period raises the possibility of a sustained risk of influenza across periods due to SOC's ability to disrupt the innate immune response to infection [14]. In our prior study of the difference in rates of influenza healthcare encounters associated with source-specific PM2.5 from 2005 to 2016 in adult New York State residents we observed an increase in the rate of healthcare encounters associated with influenza associated with interguartile range (IQR) increases in multiple source specific PM<sub>2.5</sub> constituents including diesel, sparkignition (gasoline) and biomass burning [7]. While SOC and POC were not included in this prior study the increased concentration of SOC and its potential health risk warranted the inclusion of both SOC and POC in the current study.

In this study, we examined whether the association between short term increases in ambient source-specific PM<sub>2.5</sub> concentrations and influenza healthcare visits differed between the period prior to Tier 3 implementation (2014–2016) and the initial years of Tier 3 implementation (2017–2019). We hypothesized that the Tier

3 standards could result in a lower relative rate of hospitalization or emergency department visits for influenza associated with increased light-duty vehicle (gasoline), compression-ignition, heavy-duty diesel vehicle, and SOC PM<sub>2.5</sub> concentrations in the 2017–2019 period compared to 2014–2016.

### **METHODS**

### Study population

Respiratory infection hospital admissions and emergency department visits for adult New York residents (age > 18) were obtained from the Statewide Planning and Research Cooperative System (SPARCS) database. In total, N = 135,236 hospitalizations and N = 549,528 emergency department visits of adults living within 15 miles of the Buffalo, Rochester, Albany, Bronx, Manhattan, or Queens PM<sub>2.5</sub> monitoring sites from January 1, 2014, to December 31, 2019, were retained. These monitoring sites were selected since they collect the PM<sub>2,5</sub> samples for speciation analysis. We included participants with a primary diagnosis (at time of healthcare encounter) of influenza (ICD9 = 4870, 4871, 48811, 48812, 48881, 48882; ICD10 = J09X1, J09X2, J1000, J1001, J1008, J101, J1100, J1108 and J111. In the analysis, we paired pollution to the healthcare encounter in the same day (lag 0), prior 4 days (lag day 0–3) and prior 7 days (lag days 0–6). Given the vast majority (95%) of patients with influenza will become symptomatic within the first 2 days of infection [15], and then present for a healthcare encounter sometime in the next week (if at all), the 0-6 day lag period (7 days prior to presentation) may represent the exposure closest to the start of the influenza infection. While there is variability in when patients present for care after the start of symptoms, lag day 0 (same day) and lag days 0-3 are likely representative of exposure several days into influenza infection. For this reason, we focus our discussion on the associations at the 0-6 lag days (closest to the start of infection) and less so on the shorter lags. This study was approved by the Institutional Review Board at the University at Albany, State University of New York.

# Air pollution and meteorology measurements

We obtained PM<sub>2.5</sub> compositional data from the U.S. Environmental Protection Agency (EPA) Chemical Speciation Network (www.epa.gov/aqs). In all six urban sites (Buffalo, Rochester, Albany, Manhattan, Bronx and Queens) daily samples were collected and analyzed every third day. Organic carbon, including primary organic carbon (POC) and secondary organic carbon (SOC), was measured every third day using thermo-optical analysis. The PM<sub>2.5</sub> sources were identified using EPA positive matrix factorization (PMF) version 5. Further details on this approach can be found in our prior source-specific analyses by Squizzato et al. [16]. The seven PM<sub>2.5</sub> sources included in this study included spark-ignition emissions (GAS), diesel (DIE), biomass burning (BB), road dust (RD), secondary nitrate (SN), secondary sulfate (SS), and pyrolyzed organic rich (OP). Daily ambient temperature and relative humidity were obtained from the National Weather Service.

# Statistical analysis

A time-stratified, case-crossover design [17, 18] was used to estimate the rates of respiratory infection hospital admissions and emergency department visits associated with each interquartile range increase in GAS concentrations on the same day (lag day 0). Assuming a common slope across sites for influenza hospital admissions for all sites, we fit a conditional logistic regression model stratified on each respiratory infection hospital admission matched set (1 case and 3-4 control periods per subject). This conditional logistic regression model regressed case–control status (i.e., case = 1, control = 0) against the mean GAS concentration on case and control days. Since the case-crossover approach controls for non-time-varying confounders, variables such as comorbidities, socioeconomic factors and seasons are all controlled for, by design. However, we included natural splines for temperature and relative humidity (4 degrees of freedom determined using Akaike's information criterion) [19], and PM<sub>2.5</sub> mass of all other sources (i.e., rPM2.5 (residual value) =  $PM_{2.5}$  - GAS). This same model was run for the mean GAS concentrations on lag days 0-3 and 0-6). From each model, we estimated the rate of hospitalizations or emergency department visits associated with each interquartile range increase in GAS concentration. Since we examined 3 lag times for GAS, statistical significance was defined as p < 0.017 (0.05/ 3), though we focused mainly on the magnitude and direction of the effect estimates rather than the statistical significance. Next, we examined

**Table 1.** Characteristics of respiratory infectious disease hospital admissions and emergency department visits (2014–2019), by study site/city.

	Hospitaliza (N = 20,20		Emergen departm visits (N = 70,8	ent
Characteristic	n	%	n	%
Female	11,764	58	42,722	60
AGE in years: mean (standard deviation)	69 (18)		42 (18)	
≥18–64	7201	36	61,987	87
≥65	13,004	64	8868	13
Race/ethnicity				
White	9351	62	20,316	44
Black	4588	30	22,348	48
Native American	53	0	261	1
Asian	5122	6	7420	5
Native Hawaiian	42	0	165	0
Hispanic	3816	19	19,061	27
Year				
2014	2445	12	7809	11
2015	2150	11	4918	7
2016	2554	13	8360	12
2017	3015	15	7560	11
2018	5761	29	18,049	25
2019	4280	21	24,159	34
Season				
Spring	5978	30	18,398	26
Summer	215	1	626	1
Fall	609	3	2900	4
Winter	13403	66	48,931	69
Length of stay in days mean $\pm$ standard deviation	54.6 ± 7.0		$0.2 \pm 0.7$	

whether the rates of respiratory infection admission associated with each interquartile range increase in GAS concentrations differed by period (2014–2016 and 2017–2019), by adding an interaction term (Period \* GAS) to the model. We then re-ran these models for each of the other PM<sub>2.5</sub> sources including biomass burning (BB), diesel (DIE), organic-rich phosphate (OP), road dust (RD), secondary nitrate (SN), secondary sulfate (SS) as well as SOC and POC concentrations. For each of these models a rPM<sub>2.5</sub> was recalculated and included in the model. For example, the SOC rPM<sub>2.5</sub> = PM<sub>2.5</sub> – SOC and the RD rPM<sub>2.5</sub> = PM<sub>2.5</sub> – RD. By computing the residual for all sources, the rate of influenza healthcare encounters associated with a specific PM<sub>2.5</sub> source (e.g. GAS) is thus independent of other of the PM<sub>2.5</sub> mass from other specific sources (e.g. DIE, BB, SN) [20]. Similarly, the effect of SOC is independent of POC (but not the source specific PM<sub>2.5</sub> such as GAS and BB as these sources can contain SOC). All analyses were completed using R version 3.0.1 (https://www.r-project.org/).

### **RESULTS**

Patients hospitalized for influenza were older (69 years old [SD of 18]) than patients treated and sent home from the emergency department (42 years old [SD of 18] (Table 1). The majority of patients hospitalized or treated in the emergency department were female (58% and 60% respectively) and sought care during the winter months (66% and 69% respectively). While the majority of hospitalized patients were White (62%), the majority of patients treated and sent home from the emergency department were Black (48%).

### Pollution data

Using Theil-Sen nonparametric estimator and piecewise linear regression, our prior study [21] observed reduction in PM<sub>2.5</sub> values across New York State ranging from 5.24% reduction in PM<sub>2.5</sub> in Queens (95% CI: -5.61%, -4.39%) to a reduction of 2.51% in Albany (95% CI: -3.43%, -1.32%) from 2010 to 2019. When comparing the 2017-2019 period to the 2014-2016 period in our current study, the median concentration of GAS (2014-2016: 1.04  $\mu$ g/m<sup>3</sup>; 2017–2019: 1.7  $\mu$ g/m<sup>3</sup>) and SN (2014–2016: 1.3  $\mu$ g/m<sup>3</sup>; 2017–2019: 1.8 μg/m<sup>3</sup>) showed the largest increases (Table 2). DIE (2014–2016:  $0.49 \,\mu g/m^3$ ; 2017–2019:  $0.54 \,\mu g/m^3$ ) and POC (2014–2016:  $0.5 \,\mu g/m^3$ ; 2017–2019:  $0.9 \,\mu g/m^3$ ) had smaller increases in concentrations between periods. We observed a decreased concentration of OP (2014–2016: 0.3 µg/m<sup>3</sup>; 2017–2019:  $0.6 \,\mu\text{g/m}^3$ ), RD (2014–2016:  $0.14 \,\mu\text{g/m}^3$ ; 2017–2019:  $0.08 \,\mu\text{g/m}^3$ ) and SS (2014–2016:  $0.9\,\mu g/m^3$ ; 2017–2019:  $0.3\,\mu g/m^3$ ) in 2017-2019 compared to 2014-2016 and no appreciable change in BB or SOC.

As hypothesized, the relative rates of hospital admission from influenza associated with interquartile range increases in DIE were smaller in the 2017–2019 period than the 2014–2016 period at all lag times. Specifically, each  $0.3 \, \mu g/m^3$  increase in DIE concentration in the prior 6 days was associated with a 21.3% increased rate of influenza hospitalization (95% Cl: 6.9, 37.6) in the 2014–2016 period, but a protective effect (-6.3% rate; 95% Cl: -12.7, 0.5) in the 2017–2019 period (Table 3, Fig. 1). A similar pattern was observed for influenza emergency department visits and DIE in the prior 6 days. Specifically, each  $0.4 \, \mu g/m^3$  increase in DIE concentration was associated with a 24.6% increased rate of emergency department visits for influenza (95% Cl: 14.5, 35.5) in the 2014–2016 period, and a -1.2% rate (95% Cl: -4.9, 2.7) in the 2017–2019 period (Table 4, Fig. 2).

The pattern for the association between GAS concentrations and influenza emergency department visits was different from DIE. Each 1.5  $\mu$ g/m³ increase in GAS concentration was associated with a larger excess rate of influenza emergency department visits in the 2017–2019 period (ER = 23.9%; 95% CI: 17.6, 30.6) than in the 2014–2016 period (ER = 7.7%; 95% CI: 0.1, 15.8) for lag days 0–6, but not for lag day(s) 0 or 0–3 (where no differences were observed between periods) (Table 4, Fig. 2). The rate of influenza hospitalizations associated with each IQR increase in GAS concentration in lag days 0–6 was also larger in the 2017–2019 period (ER = 16.4%; 95% CI: 5.5%, 28.4%) than in the 2014–2016 period (ER = 11.7%; 95% CI: -1.1%, 26.1%).

The pattern of ER of biomass burning source-specific PM was the inverse of the DIE results, with no association between BB and influenza hospitalizations in the 2014–2016 period and increased relative rates in the 2017–2019 period at all lag periods. Specifically, each  $0.4\,\mu\text{g/m}^3$  increase in BB concentration in lag days 0–6 was associated with a 2.3% increased rate (95% CI: -6.8, 12.3) in the 2014–2016 period, and a 18.7% increased rate of hospitalizations for influenza (95% CI: 12.2, 25.5) in the 2017–2019 period (Table 3, Fig. 1). The pattern for BB/influenza emergency department visits was the same as hospitalizations. We observed a 1.2% increase in emergency department visits (95% CI: -3.3, 5.9) associated with each  $0.4\,\mu\text{g/m}^3$  increase in BB in the 2014–2016 period and a 16.8% increase (95% CI: 14.0, 19.7) in the 2017–2019 period (Table 4, Fig. 2).

The rate of influenza healthcare encounters associated with increased concentrations of SOC and POC were larger in the 2017–2019 compared to 2014–2016. Each 0.4 μg/m³ increase in POC concentrations in lag days 0–6 was associated with an 11.8% increase in emergency department visits for influenza (95% Cl: 8.0, 15.6) in the 2017–2019 period, but only a 1.5% decrease in 2014–2016 (95% Cl: –6.2, 3.3) (Table 4). Each 0.7 μg/m³ increase in SOC concentration in lag days 0–6 was associated with a 13.4% increase in emergency department visits for influenza (95% Cl: 7.7, 19.3) in the 2017–2019 period and a 1.6% decrease (–6.6, 3.7) in

13.10 13.10 11.94 14.67 14.67 4.09 5.68 5.68 14.55 14.55 12.47 10.66 4.59 33.09 32.38 33.09 99.90 92.06 3.24 2.47 13.64 4.81 4.21 3.72 4.21 7.05 95<sup>th</sup> %tile 17.10 17.80 1.19 1.37 1.50 4.53 4.10 4.73 1.39 1.98 1.06 0.69 0.88 6.57 6.52 6.73 3.30 5.14 2.27 1.70 1.60 2.17 2.15 16.14 1.75 1.77 1.75 0.61 1.74 2.31 18.11 90.65 4.31 0.73 2.30 2.36 0.50 0.42 0.25 0.34 0.22 2.90 2.23 0.10 1.16 1.00 1.20 1.13 1.08 79.39 9.47 0.93 0.95 0.91 0.80 0.84 1.97 0.81 2.81 2.81 1.27 7.60 6.86 0.10 0.50 0.50 0.52 0.49 0.54 1.51 1.00 1.66 0.14 0.31 0.08 0.09 0.14 1.53 1.20 1.59 0.40 0.90 0.24 0.72 0.50 0.90 99.0 0.62 0.51 4.07 4.87 -0.07 0.30 1.16 -0.06 0.03 0.06 0.02 0.59 0.40 -0.11 0.28 0.33 -0.01 0.73 0.06 0.47 0.33 0.31 0.31 0.41 -0.170.37 0.21 Controls -0.02 -0.12 -0.03 -0.13-0.03 -0.61 -0.11 -0.02 -0.06 -0.12-0.12-0.03-0.01 0.05 -0.01 0.09 -0.46-0.64 0.08 0.10 -0.01 -6.70 -6.60 40.04 2.80 2.65 0.01 0.17 0.10 0.01 0.01 -7.010.7 5.28 2.35 11.94 11.94 13.05 13.05 3.49 5.68 5.68 14.55 14.55 11.33 11.88 11.88 3.72 32.38 5.52 2.69 2.69 3.72 3.38 5.52 8.69 4.81 7.01 4.60 Distribution of source-specific PM<sub>2.5</sub> concentrations (µg/m³), by time period, for case and control periods 18.60 1.26 1.42 5.08 4.53 5.72 1.40 1.98 0.73 0.89 6.73 9.60 6.73 3.54 2.00 2.17 19.01 1.35 1.22 0.61 5.24 2.57 1.63 2.24 19.01 11.33 20.14 1.88 2.47 2.22 14.07 10.89 0.80 0.42 0.36 1.10 1.19 82.14 75<sup>th</sup> %tile 1.00 1.00 0.80 0.75 0.82 2.46 2.08 2.67 0.50 0.27 0.23 3.41 2.94 3.52 1.40 2.20 1.02 1.24 1.20 1.12 7.58 10.33 80.59 74.11 10.66 6.90 0.52 0.49 0.55 1.62 1.04 1.74 0.29 90.0 0.10 0.12 0.08 1.63 1.80 0.45 0.78 0.50 0.70 7.50 0.54 0.52 0.13 1.31 0.85 0.31 0.94 0.67 7.25 0.54 0.67 4.30 4.03 -0.16 -0.07 -0.02 -0.080.03 0.46 0.90 -0.01 0.39 0.29 0.34 0.28 0.30 1.02 0.45 1.25 0.05 0.02 0.72 0.07 0.50 0.36 0.37 0.43 0.21 0.22 0.21 0.31 0.31 0.70 5.1 0.16 -0.12 -0.00 -0.64 0.10 -0.02-0.12 -0.02 -0.03 -0.05 -0.14 -0.15-0.04 -0.07 -0.46 -0.70 -6.60 %tile 2.73 0.03 0.68 0.06 -0.01 0.11 0.08 0.11 0.01 -6.20 0.11 0.01 0.01 -6.01 **Fime Period** 2014-2016 2017-2019 2014-2016 2017-2019 2017-2019 2014-2016 2017-2019 2014-2016 2014-2016 2017-2019 2014-2016 2017-2019 2014-2016 2017-2019 2014-2016 2017-2019 2014-2016 2017-2019 2014-2016 2017-2019 2017-2019 2014-2016 2017-2019 2014-2016 Overall Organic-rich phosphate Primary organic carbon Secondary organic carbon Secondary nitrate Secondary sulfate Relative humidity **Biomass burning Temperature** Total PM<sub>2.5</sub> Road dust Table 2. Sasoline Diesel

<0.001 2000 0.54 0.01 0.55 0.51 0.36 0.04 0.11 ŧ 8.9 (-17.0, -0.0) 13.7 (-23.9, -2.2) Excess rate (95% 2.3 (-6.8, 12.3) 11.7 (-1.1, 26.1) 4.1 (-5.7, 14.9) 8.1 (-0.6, 17.5) 6.2 (-17.6, 6.7) 1.2 (-10.8, 9.4) 18.7 (12.2, 25.5) 1.5 (-3.5, 6.8) 0.7 (-6.3, 5.1) 1.6 (-4.5, 8.0) 21.3 (6.9, 37.6) 16.4 (5.5, 28.4) 14.8 (0.3, 31.4) 9.3 (3.7, 15.2) QR (µg/m³) 9.0 0.4 0.7 0.7 6.1 6. 7. 0.3 0.3 0.7 0.1 0.1 2009 3335 1721 3335 3335 1721 3335 3335 3449 2009 1721 1721 1721 1721 p-value 0.001 60.0 0.87 0.77 0.28 0.83 0.21 0.26 0.88 Excess rate (95% 8.3 (-2.8, 20.7) 4.6 (-12.7, 4.3) 2.0 (-6.6, 11.4) 5.7 (-1.3, 13.1) 1.2 (-7.0, 10.1) 7.8 (-3.8, 20.7) 8.8 (-0.4, 18.9) 0.8 (-4.2, 6.1) 2.3 (-7.8, 3.5) 1.2 (-4.7, 7.4) 2.9 (-7.6, 2.0) 0.2 (-6.3, 7.0) 1.0 (-6.8, 9.3) 1.3 (-7.3, 5.0) 2.6 (-2.4, 7.8) 1.8 (5.8, 18.2) 6. 0.3 6: 3 <u>..</u> 0.3 0.7 0.1 1513 3064 1513 1513 3064 1513 3064 1513 3064 1513 3064 1852 3174 1852 p-valu 0.04 0.16 0.18 0.23 ਹ -7.5 (-13.8, -0.7) Excess rate (95% 4.7 (-5.5, 16.1) 7.4 (-0.6, 16.1) 4.0 (-10.5, 3.0) 3.4 (-1.7, 8.8) 2.3 (-5.2, 10.4) 0.5 (-5.1, 6.5) 1.9 (-7.8, 4.3) 1.5 (-2.2, 5.2) 2.5 (-3.4, 8.7) 11.6 (3.1, 20.7) 3.0 (-0.6, 6.8) 0.1 (-3.6, 3.9) -0.9 (-5.7, 4.2) 0.4 (-5.8, 7.0) -0.7 (-4.9, 3.6) p-value for interaction between periods (2014–2016 and 2017–2019). 0.5 2.2 2.2 0.7 0.7 0.2 0.2 0.5 0.5 2075 853 853 3588 2075 Time period 2014-16 2014-16 2017-19 2017-19 2014-16 2017-19 2014-16 2017-19 Lag day 2014-16 2017-19 2014-16 2014-16 2017-19 2014-16 Spark-ignition emissions (GAS) Pyrolyzed organic rich (OP) Secondary nitrate (SN) Secondary sulfate (SS) Siomass burning (BB) Road dust (RD) Diesel (DIE) 900

Excess rate of acute hospitalizations for influenza associated with each interquartile range increase in source-specific PM $_{2.5}$ 

Table 3.

specific constituent (Lag days) Hospitalizations 2014-2016 Diesel Lag 0-Hospitalizati 2017-2019 Diesel Lag 0-3 Diesel Lag 0-6 Biomass Lag 0 Biomass Lag 0-3 Biomass Lag 0-6 Gasoline Lag 0 Source Gasoline Lag 0-3 Gasoline Lag 0-6 40 -20 n 20 Excess rate (%) of influenza hospitalization

**Fig. 1** Excess rate of hospitalizations for influenza. Excess rate of hospitalizations for influenza associated with IQR increases in diesel (DIE), biomass burning (BB) and gasoline (GAS) at multiple lag times.

2014–2016. The difference in the rates between periods was slightly smaller for the hospitalizations for influenza compared to emergency department visits.

Similar to SOC and POC, the rate of influenza emergency department visits associated with SN and SS at the 0–6 lag time were larger in the 2017–2019 period compared to 2014–2016 period. One unique finding was the decreased rate of influenza hospitalizations associated with a 1.7  $\mu$ g/m³ increase in SS at the 0–6 lag period (ER -13.7%; 95% CI: -23.9, -2.2). We observed larger rates of influenza hospitalizations associated with each 0.7  $\mu$ g/m³ increase in OP concentration in lag days 0–6 in the 2017–2019 period (ER 14.8%; 95% CI: 0.3, 31.4) than the 2014–2016 period (ER of -8.9%; 95% CI: (-17.0, 0.0) at the 0–6 lag days (Table 3). There was no clear pattern of association for road dust and influenza healthcare encounters.

### DISCUSSION

In this study of source-specific PM<sub>2.5</sub> concentrations and influenza hospitalizations and emergency department visits in Rochester, NY, we estimated and compared the relative rate of influenza healthcare encounters associated with increased source-specific PM<sub>2.5</sub>, SOC, and POC concentrations both before (2014–2016) and after (2017-2019) Tier 3 vehicle emissions standards were implemented. As hypothesized, we observed a decrease in the rate of influenza hospitalizations and emergency department visits associated with increased DIE concentrations at all lag times in the 2017-2019 period compared to the 2014-2016 period. Unexpectedly, in the 2017–2019 period, we observed an increase in the rate of emergency department visits associated with spark-ignition/ gasoline vehicle emissions (GAS) in the prior one week (lag days 0-6; compared to 2014-2016 period) and no change between periods at other lag times. A similar increase in the rate of healthcare encounters for influenza in the 2017–2019 period was observed for biomass burning, SS, POC, and SOC, but not other PM<sub>2.5</sub> sources. The relative toxicity per unit mass of some of these PM<sub>2.5</sub> sources may have been affected by implementation of the Tier 3 emissions standards and other prior emissions standards.

The need to examine the health effects of sources of PM<sub>2.5</sub> is supported by our recent study on the effect of total PM<sub>2.5</sub> on the rate of influenza healthcare encounters [22]. Despite reductions in overall PM<sub>2.5</sub> concentrations during the early implementation period of the Tier 3 standards, the rates of influenza healthcare encounters associated with IQR increases in PM<sub>2.5</sub> concentrations

Table 4.         Excess rate of acute emergency department visits for influenza associated with each interquartile range increase in source-specific PM <sub>2.5</sub> .	cute emergen	cy depart	ment visits fo	ır influenza associate	d with ea	ch interq	luartile range	increase in source-s	pecific PM	2.5•			
	Lag day	0				0-3				9-0			
	Time period	2	IQR (µg/m³)	Excess rate (95% CI)	p-value*	>	IQR (µg/m³)	Excess rate (95% CI)	p-value*	2	IQR (µg/m³)	Excess rate (95% CI)	p-value*
Diesel (DIE)	2014–16	4910	0.4	7.1 (1.7, 12.9)	0.05	3842	0.3	9.7 (3.4, 16.3)	<0.001	4653	0.4	24.6 (14.5, 35.5)	<0.001
	2017–19	13,337	0.4	1.1 (-1.5, 3.7)		11,485	0.3	-1.7 (-4.2, 0.8)		12,861	0.4	-1.2 (-4.9, 2.7)	
Biomass burning (BB)	2014-16	4910	0.4	0.2 (-2.9, 3.4)	<0.001	3842	0.4	0.6 (-4.2, 5.6)	<0.001	4653	0.4	1.2 (-3.3, 5.9)	<0.001
	2017–19	13,337	0.4	7.7 (6.0, 9.4)		11,485	0.4	11.6 (9.0, 14.4)		12,861	0.4	16.8 (14.0, 19.7)	
Spark-ignition emissions (GAS)	2014-16	4910	2.0	11.1 (4.9, 17.7)	90.0	3842	1.5	8.0 (0.8, 15.7)	95.0	4653	1.5	7.7 (0.1, 15.8)	<0.001
	2017–19	13,337	2.0	4.9 (1.4, 8.4)		11,485	1.5	10.3 (5.7, 15.0)		12,861	1.5	23.9 (17.6, 30.6)	
Pyrolyzed organic rich (OP)	2014–16	4910	0.7	-3.5 (-6.2, -0.7)	<0.001	3842	9.0	0.2 (-3.5, 4.0)	90.0	4653	0.7	-4.3 (-8.9, 0.6)	0.16
	2017–19	13,337	0.7	6.9 (3.4, 10.5)		11,485	9.0	6.5 (1.1, 12.1)		12,861	0.7	1.1 (-4.7, 7.3)	
Road dust (RD)	2014–16	4910	0.2	2.7 (0.5, 5.0)	0.00	3842	0.1	2.1 (-0.2, 4.4)	0.14	4653	0.1	0.9 (-1.7, 3.7)	0.40
	2017–19	13,337	0.2	-1.6 (-3.3, 0.2)		11,485	0.1	-0.0 (-1.8, 1.8)		12,861	0.1	-0.6 (-3.0, 1.9)	
Secondary nitrate (SN)	2014–16	4910	2.1	-2.9 (-6.6, 1.0)	0.01	3842	1.9	-3.4 (-8.4, 1.8)	0.04	4653	1.9	-3.4 (-9.2, 2.7)	<0.001
	2017-19	13,337	2.1	2.7 (0.1, 5.4)		11,485	1.9	2.3 (-1.2, 5.9)		12,861	1.9	12.9 (8.3, 17.8)	
Secondary sulfate (SS)	2014-16	4910	1.9	-2.6 (-6.5, 1.4)	<0.001	3842	1.3	-0.7 (-5.1, 4.0)	<0.001	4653	1.7	-11.3 (-17.7, -4.4)	<0.001
	2017-19	13,337	1.9	20.8 (16.1, 25.8)		11,485	1.3	15.4 (10.7, 20.3)		12,861	1.7	8.9 (1.7, 16.6)	
POC	2014-16	5644	0.4	0.7 (-2.6, 4.2)	0.07	4819	0.4	-2.9 (-6.8, 1.2)	<0.001	5509	0.4	-1.5 (-6.2, 3.3)	<0.001
	2017–19	13,879	0.4	4.1 (1.9, 6.3)		11,965	0.4	5.0 (2.1, 7.9)		13,295	0.4	11.8 (8.0, 15.6)	
SOC	2014-16	5644	0.8	-1.3 (-4.6, 2.1)	0.15	4819	9.0	-0.7 (-4.7, 3.4)	0.01	5509	0.7	-1.6 (-6.6, 3.7)	<0.001
	2017-19	13,879		1.4 (-1.5, 4.5)		11,965		5.0 (1.2, 8.9)		13,295		13.4 (7.7, 19.3)	
$^*$ p-value for interaction between periods (2014–2016 and 2017–2019).	ween periods (	2014–201	6 and 2017–20	019).									

Source specific constituent (Lag days) Diesel Lag 0 ☐ ED Visits 2014-2016 ED Visits 2017-2019 Diesel Lag 0-3 Diesel Lag 0-6 Biomass Lag 0 Biomass Lag 0-3 Biomass Lag 0-6 Gasoline Lag 0 Gasoline Lag 0-3 Gasoline Lag 0-6 -10 10 30 40 20 Excess rate (%) of Influenza ED visits

Fig. 2 Excess rate of emergency department visits for influenza. Excess rate of emergency department visits for influenza associated with IQR increases in diesel (DIE), biomass burning (BB) and gasoline (GAS) at multiple lag times.

were not lower in the 2017–2019 period compared to the 2014–2016 period. As policy makers consider interventions to address specific sources of air pollution, the health effects of the remaining  $PM_{2.5}$  (which has a new relative composition) will require continued assessment.

Since the time of our prior study of source-specific PM<sub>2.5</sub> and influenza [7] that reported increased rates of influenza hospitalization and emergency department visits associated with increased PM<sub>2.5</sub>, GAS, and DIE, the global COVID-19 pandemic redirected some of the environmental science community to focus environmental research on COVID-19 infection [23]. The uniqueness of the SARS-CoV-2 infection and its changing pandemic dynamics limited the applicability of the air pollution/COVID-19 studies to other seasonal respiratory viruses [24, 25]. Although there have been several studies estimating influenza/PM<sub>2.5</sub> associations [4, 26], there have been few previous studies estimating associations between source-specific PM<sub>2.5</sub> or PM<sub>2.5</sub> constituents and influenza health care encounters [27]. In the interim from our prior source-specific study, a recent study in Guangdong Province, China observed a 2.8% increase in the relative risk of influenza infection associated with a 10 µg/m<sup>3</sup> increase in PM<sub>2.5</sub> concentrations in the 8 days prior to symptom onset [6]. Though only indirectly related to respiratory infection, an accountability study in Los Angeles, CA observed a decrease in risks of emergency department visits for asthma associated with increased PM<sub>2.5</sub> constituents (organic carbon, sulfate, and nitrate) concentrations over time (2005–2016) [28]. Not all source-specific PM<sub>2.5</sub> findings in our current study are consistent with these studies.

Our current study employed an improved PMF analysis to better apportion the aerosol resulting in differences between the effect estimates in the 2014–2016 period when comparing with our prior study source-specific/influenza study [7]. In this prior study we examined the association between increased source-specific PM<sub>2.5</sub> (using the prior version of the PMF analysis) and influenza healthcare encounters in New York State from the same SPARCS database. Given the volume of data given the inclusion of three periods from 2005 to 2016 and the inclusion of culture negative pneumonia, we did not present the period-specific data in that study (only the total period). While the absolute values are not directly comparable, the trends in excess rates from 2014 to 2019 are important to compare. For example, the excess rates for GAS/influenza healthcare encounters decreased from 2005 to 2007 (ER

17%; 95% CI: 2.7, 33.3) to 2014–2016 (ER 6.1%; 95% CI: -2.4, 9.8) in lag days 0–6 (and similar pattern for lag days 0–3) (unpublished data). In the current study, rather than a trend of smaller excess rates over time, the excess rates for the GAS/influenza emergency department visit associations appeared similar (or larger) in the 2017–2019 period (ER = 23.9%; 95% CI: 17.6, 30.6) compared to the 2014–2016 period (ER 7.7%; 95% CI: 0.1, 15.8) for lag days 0–6.

The observation of a similar (or increased) relative rate for healthcare encounters for influenza associated with GAS in the 2017-2019 period compared to 2014-2016, despite the early implementation of the 2017 Tier 3 vehicle standards, is inconsistent with the hypothesis that Tier 3 vehicle standards could have reduced the toxicity per unit mass of the GAS component. Specifically, if the early implementation of the Tier 3 emissions standards decreased the secondary organic carbon (SOC) component of PM<sub>2.5</sub>, thereby reducing the oxidative potential of PM<sub>2.5</sub>, we would also expect to see a decrease in the rate of healthcare encounters for influenza associated with increased PM<sub>2.5</sub> and GAS concentrations. The lack of reduction in the GAS/influenza association may be related to the incomplete fleet penetration of Tier 3 vehicles. While 36% of the current light duty vehicle fleet was purchased after 2017, we do not yet have data on what proportion of those vehicles met the Tier 3 standards (so the actual Tier 3 market penetration must be roughly 36% or lower) [29]. There may be other explanations including a change in the particle size distribution within GAS. The reduction in the mass of relatively less toxic particles such as secondary sulfate may have led to a relative increase in the toxicity per unit mass of the remaining sources. While the proportion of GAS consisting of UFP increased related in part to the 50% penetrance of gasoline direct injection in 2016, the change in toxicity related to this change will require further study.

In our prior study [7], the ERs for the DIE/Influenza emergency department visits were similar in the 2005-2007 period (ER 5.8% [95% CI: 2.6%, 9.0%]) compared to the 2014–2016 period (ER of 6.3% [95% CI 1.3%, 11.6%]) at the 0-6 lag days (unpublished data). In contrast, we observed a substantial decrease in the ER of the DIE/ Influenza from 2014–2016 to the 2017–2019 period in the current study. The observed decrease in excess rates of influenza healthcare encounters associated with each 0.3–0.5 µg/m<sup>3</sup> increase in DIE may be related to changes prior to the implementation of the 2017 Tier 3 vehicle standard. As a result of multiple diesel fuel related policies targeting the sulfur content of diesel, the concentrations of SO<sub>2</sub> (and SS) have decreased [8]. Though NO<sub>2</sub> decreased in part due to selective catalytic reduction (SCR) systems (required in new vehicles in 2010), the lack of reduction in SN concentrations may be related to an increase in ammonia that escapes the SCR system [30]. Since the Tier 3 vehicle emission standard predominantly addressed gasoline vehicles, it is unclear to what degree the early implementation of this regulation had the potential to reduce the toxicity per unit mass of DIE. Though not guaranteed, the toxicity of both DIE and GAS would be expected to continue to fall, as the emissions from both diesel and gasoline vehicles come below the recommended Tier 3 standard.

When considering non-traffic related PM<sub>2.5</sub> sources, the increase in the rate of influenza healthcare encounters associated with BB concentrations in the 2017–2019 period, compared to the 2014–2016 period, may be related to a change in the characteristics of wildfire smoke or residential wood-burning smoke and/or their interaction with other pollutants in the outdoor atmosphere. While wildfire smoke has been suggested to comprise up to 25% of PM<sub>2.5</sub> in the United States (and up to 50% in the Western U.S.) [31], the majority of BB comes from residential wood burning in the Rochester, NY area [32, 33]. The respiratory health effects of residential biomass burning are particularly difficult to study given the complexity of the factors that determine its emission profile including the condition of the fuel sources, efficiency of burning and ventilation of the indoor space

[34]. It remains unclear which, if any, of these variables have changed in NYS to explain a change in relative toxicity in BB from residential wood burning. For wildfire smoke, climate change driven increases in outdoor temperature and reductions in humidity, combined with an increase in large diameter solid fuels (felled trees) due to high winds and draught or flooding is expected to lead to an increase in frequency, intensity and duration of wildfires [35]. Since the proportion of PM<sub>2.5</sub> comprised of wildfire/biomass burning increases with further climate change, the reactive potential and toxicity per unit mass of PM<sub>2.5</sub> may also increase [36, 37]. However, due to lower average temperature in New York State (NYS) compared to California, where the majority of the U.S. based literature focuses, we would expect less atmospheric reactivity, and potentially less toxicity, during studies similar to our current study in the winter months. Though wildfire smoke is not currently the dominant biomass source in NYS, the proportion of BB in NYS from wildfire smoke may increase in the future due to an expected increase in the frequency of dynamic weather patterns similar to the lingering low pressure system which led to significant biomass exposure in NYS from Canadian wildfires during the summer of 2023 [38]. Similar to the increased risk of hospitalization from asthma related to wildfire smoke, we may expect to see an increased risk of influenza associated with PM<sub>2.5</sub> since the proportion of BB in NYS that comes from wildfire smoke increases [37].

Though the continued penetrance of Tier 3 vehicles into the New York State vehicle fleet is expected to result in a decrease in SOC [8], it is possible that the increase in organic carbon from wildfire activity over time will contribute more SOC to PM<sub>2.5</sub> [39], thereby stabilizing SOC concentrations. The increase in rates of influenza healthcare encounters associated with increased SOC concentrations in 2017-2019 compared to 2014-2016 may indicate an increase in toxicity per unit mass of SOC. This pattern is the same as the main contributors to SOC, gasoline vehicle emissions (GAS) at lag days 0-6, and biomass burning. While we also observed an increased rate of influenza hospitalizations and emergency department visits with secondary sulfate (SS) (except for the 0-6 time period for hospitalization), due to the role of SS as a vector for a heterogeneous collection of condensates, the health effect of specific secondary sulfates remains uncertain [40]. Since the majority of PM<sub>2.5</sub> is from inorganic and organic secondary aerosols (i.e. SIA and SOA), the reactivity of each constituent PM<sub>2.5</sub> source such as traffic related air pollution (DIE and GAS) and biomass burning have been found to be more similar to the higher oxidative potential of SOC than the less reactive POC [12, 41]. For a viral infection like influenza, PM<sub>2.5</sub> constituents with high oxidate potential (e.g. DIE, GAS and BB) have the potential to enhance viral entry into cells, impair pathogen recognition, immune signaling, and immune cell function [14]. We will need to continue to focus future studies on the potential pathophysiologic mechanisms of the association between PM<sub>2.5</sub> and influenza healthcare encounters and what collection of sources appear to be the main drivers of this association.

Inequity in both exposure to PM<sub>2.5</sub> and treatment for influenza exist in the U.S. Our study observed that a higher proportion of Black and Hispanic patients with influenza were treated and released from the ED (48% and 27% respectively) compared to those who were admitted (30% and 19% respectively). Our patient proportions are not consistent with national data observing the odds of hospitalization (including intensive care) and death for influenza are higher in Black, Hispanic, and Asian patients when compared to white patients [42]. It is also well known that Black and Hispanic communities suffer a disproportionate burden of traffic related air pollution [43]. These two facts combine to increase the risk of a poor outcome from influenza infection in Black and Hispanic communities when compared to White communities. In a future analysis, we will need to consider how the difference in our local healthcare practice patterns and

geographic distribution of source-specific PM<sub>2.5</sub> may affect the association between PM/influenza in different racial and ethnic communities.

Although there were several strengths of this study, including a large sample size and inclusion of the 4 largest cities in New York State, there are several limitations that should be considered when making inference. First, there is likely exposure misclassification and downward bias resulting from assignment of the same air pollutant concentrations for everyone within 15 miles of an individual monitor (e.g., Buffalo), no matter how far they live from the monitoring station. Second, this study was a wintertime study given the seasonality of influenza infection, limiting the generalizability to year-round respiratory infections or bacterial respiratory infections. Last, to be consistent with our previous analyses, we used a case-crossover design and conditional logistic regression analyses to estimate the rate of influenza hospitalizations and emergency department visits associated with sourcespecific PM<sub>2.5</sub>, SOC, and POC concentrations in the previous 7 days. However, as described in our previous analysis of influenza/PM25 associations in these same cities, we cannot assess lag periods longer than 7 days without overlap of case and control period dates and thus case and control period pollutant concentrations [22]. Thus, we are limited to assessment of pollutant concentrations in just the previous 1–7 days. Last, the study population is only adult residents, and not children, and only those adults living in urban centers of the state (i.e., within 15 miles). Thus, generalizability of results to children or to adult residents in rural locations is less certain.

### **CONCLUSION**

While there did not appear to be any clear reduction in influenza healthcare encounters from gasoline emissions during the early implementation period of the Tier 3 emissions standards, we did see a reduction in the rate of healthcare encounters associated with diesel emissions. It is possible that we are seeing the effect of prior diesel regulations to lower sulfur content and that the ability to fully evaluate the health effects of the gasoline emission regulation may come only later once the Tier 3 vehicle penetration into the entire New York State vehicle fleet increases. The finding of a potential increase in the toxicity per unit mass of biomass burning (including wildfires) is concerning due to the expectation that this component of PM<sub>2.5</sub> is expected to continue to increase as a proportion of PM<sub>2.5</sub> due to climate change. Furthermore, addressing the burden of both influenza and air pollution on disproportionately affected communities will be an important research focus in the future, especially as climate change serves to worsen air quality and increase the frequency and intensity of natural disasters. Given the complexity of the atmospheric chemistry of a changing PM<sub>2.5</sub> composition, future studies will need to evaluate effect modification of the rate of respiratory infection healthcare encounters associated with PM<sub>2,5</sub> by heat waves and cold waves or implementation of air quality policies.

# DATA AVAILABILITY

The data that support the findings of this study are available from the Statewide Planning and Research Cooperative System (SPARCS) in New York State, but restrictions apply to the availability of these data, which were used under an agreement for the current study, and so are not publicly available. Data can be made available from SPARCS after an application process <a href="https://www.health.ny.gov/statistics/sparcs/">https://www.health.ny.gov/statistics/sparcs/</a>.

### **REFERENCES**

- 1. CDC. Influenza. 2023. https://www.cdc.gov/nchs/fastats/flu.htm.
- Chow EJ, Uyeki TM, Chu HY. The effects of the COVID-19 pandemic on community respiratory virus activity. Nat Rev Microbiol. 2023;21:195–210.

- Croft DP, Zhang WJ, Lin S, Thurston SW, Hopke PK, Masiol M, et al. The Association between Respiratory Infection and Air Pollution in the Setting of Air Quality Policy and Economic Change. Ann Am Thorac Soc. 2019;16:321–30.
- Horne BD, Joy EA, Hofmann MG, Gesteland PH, Cannon JB, Lefler JS, et al. Short-Term Elevation of Fine Particulate Matter Air Pollution and Acute Lower Respiratory Infection. Am J Respir Crit Care Med. 2018;198:759–66.
- EPA. Integrated Science Assessment (ISA) for Particulate Matter. 2019. https:// www.epa.gov/isa/integrated-science-assessment-isa-particulate-matter.
- Zhang R, Lai KY, Liu W, Liu Y, Ma X, Webster C, et al. Associations between shortterm exposure to ambient air pollution and influenza: an individual-level casecrossover study in Guangzhou, China. Environ Health Perspect. 2023;131:127009.
- Croft DP, Zhang W, Lin S, Thurston SW, Hopke PK, van Wijngaarden E, et al. Associations between Source-Specific Particulate Matter and Respiratory Infections in New York State Adults. Environ Sci Technol. 2020;54:975–84.
- Chen Y, Rich DQ, Masiol M, Hopke PK. Changes in ambient air pollutants in New York State from 2005 to 2019: Effects of policy implementations and economic and technological changes. Atmos Environ. 2023;311:119996.
- Hopke PK, Hidy G. Changing emissions results in changed PM2. 5 composition and health impacts. Atmosphere. 2022;13:193.
- EPA. Final Rule for Control of Air Pollution from Motor Vehicles: Tier 3 Motor Vehicle Emission and Fuel Standards. 2023. Available from: https://www.epa.gov/regulations-emissions-vehicles-and-engines/final-rule-control-air-pollution-motor-vehicles-tier-3.
- 11. Lin SXY, Thandra S, Qi Q, Hopke PK, Thurston SW, Croft DP, et al. PM2.5 and its components and respiratory disease healthcare encounters unanticipated increased exposure-response relationships in recent years after environmental policies. Environ Pollut. 2024;360:124585.
- Lovett C, Baasiri M, Atwi K, Sowlat MH, Shirmohammadi F, Shihadeh AL, et al. Comparison of the oxidative potential of primary (POA) and secondary (SOA) organic aerosols derived from α-pinene and gasoline engine exhaust precursors. F1000Research. 2018;7:1031.
- Wang S, Takhar M, Zhao Y, Al Rashdi LNS, Chan AW. Dynamic oxidative potential of organic aerosol from heated cooking oil. ACS Earth Space Chem. 2021;5:1150–62.
- Rebuli ME, Brocke SA, Jaspers I. Impact of inhaled pollutants on response to viral infection in controlled exposures. J Allergy Clin Immunol. 2021;148:1420–9.
- Lessler J, Reich NG, Brookmeyer R, Perl TM, Nelson KE, Cummings DAT. Incubation periods of acute respiratory viral infections: a systematic review. Lancet Infect Dis. 2009;9:291–300.
- Squizzato S, Masiol M, Rich DQ, Hopke PK. A long-term source apportionment of PM2. 5 in New York State during 2005–2016. Atmos Environ. 2018;192:35–47.
- 17. Maclure M. The case-crossover design: a method for studying transient effects on the risk of acute events. Am J Epidemiol. 1991;133:144–53.
- Levy D, Lumley T, Sheppard L, Kaufman J, Checkoway H. Referent selection in case-crossover analyses of acute health effects of air pollution. Epidemiology. 2001;12:186–92.
- Aho K, Derryberry D, Peterson T. Model selection for ecologists: the worldviews of AIC and BIC. Ecology. 2014;95:631–6.
- Mostofsky E, Schwartz J, Coull BA, Koutrakis P, Wellenius GA, Suh HH, et al. Modeling the association between particle constituents of air pollution and health outcomes. Am J Epidemiol. 2012;176:317–26.
- Chen Y, Rich DQ, Hopke PK. Changes in source specific PM2.5 from 2010 to 2019 in New York and New Jersey identified by dispersion normalized PMF. Atmos Res. 2024;304:107353.
- Croft DP, Utell MJ, Liu H, Lin S, Hopke PK, Thurston SW, et al. Change in rate of healthcare encounters for respiratory infection from air pollution exposure after improved vehicle emissions standards in New York State. Air Quality Atmos Health. 2024;17:1267–80.
- Monoson A, Schott E, Ard K, Kilburg-Basnyat B, Tighe RM, Pannu S, et al. Air pollution and respiratory infections: the past, present, and future. Toxicological Sci. 2023;192;3–14
- Wu X, Nethery RC, Sabath MB, Braun D, Dominici F. Air pollution and COVID-19 mortality in the United States: Strengths and limitations of an ecological regression analysis. Sci Adv. 2020;6:eabd4049.
- Barceló MA, Saez M. Methodological limitations in studies assessing the effects of environmental and socioeconomic variables on the spread of COVID-19: a systematic review. Environ Sci Eur. 2021;33:1–18.
- Pirozzi CS, Jones BE, VanDerslice JA, Zhang Y, Paine IIIR, Dean NC. Short-Term Air Pollution and Incident Pneumonia. A Case-Crossover Study. Ann Am Thorac Soc. 2018;15:449–59.
- Krall JR, Mulholland JA, Russell AG, Balachandran S, Winquist A, Tolbert PE, et al. Associations between Source-Specific Fine Particulate Matter and Emergency Department Visits for Respiratory Disease in Four U.S. Cities. Environ Health Perspect. 2017;125:97–103.

- Bi J, D'Souza RR, Rich DQ, Hopke PK, Russell AG, Liu Y, et al. Temporal changes in short-term associations between cardiorespiratory emergency department visits and PM2. 5 in Los Angeles, 2005 to 2016. Environ Res. 2020;190:109967.
- Yount CS, Utell MJ, Hopke PK, Thurston SW, Lin S, Ling FS, et al. Triggering of STelevation myocardial infarction by ultrafine particles in New York: Changes following Tier 3 vehicle introduction. Environ Res. 2023;216:114445.
- 30. Hopke PK, Querol X. Is Improved Vehicular NOx Control Leading to Increased Urban NH3 Emissions? Environ Sci Technol. 2022;56:11926–7.
- 31. Burke M, Driscoll A, Heft-Neal S, Xue J, Burney J, Wara M. The changing risk and burden of wildfire in the United States. Proc Natl Acad Sci. 2021;118:e2011048118.
- 32. Wang Y, Hopke PK, Rattigan OV, Chalupa DC, Utell MJ. Multiple-year black carbon measurements and source apportionment using delta-C in Rochester, New York. J Air Waste Manag Assoc. 2012;62:880–7.
- 33. Wang YG, Hopke PK, Xia XY, Rattigan OV, Chalupa DC, Utell MJ. Source apportionment of airborne particulate matter using inorganic and organic species as tracers. Atmos Environ. 2012;55:525–32.
- Sigsgaard T, Forsberg B, Annesi-Maesano I, Blomberg A, Bølling A, Boman C, et al. Health impacts of anthropogenic biomass burning in the developed world. Eur Respiratory J. 2015;46:1577–88.
- 35. Odwuor A, Yañez C, Chen Y, Hopkins F, Moreno A, Xu X, et al. Evidence for multidecadal fuel buildup in a large California wildfire from smoke radiocarbon measurements. Environ Res Lett. 2023;18:094030.
- Rice MB, Henderson SB, Lambert AA, Cromar KR, Hall JA, Cascio WE, et al. Respiratory impacts of wildland fire smoke: future challenges and policy opportunities. An official American Thoracic Society workshop report. Ann Am Thorac Soc. 2021:18:921–30.
- DeFlorio-Barker S, Crooks J, Reyes J, Rappold AG. Cardiopulmonary effects of fine particulate matter exposure among older adults, during wildfire and non-wildfire periods, in the United States 2008–2010. Environ health Perspect. 2019:127:037006.
- Thurston G, Yu W, Luglio D. An Evaluation of the Asthma Impact of the June 2023 New York City Wildfire Air Pollution Episode. Am J Respiratory Crit Care Med. 2023;208:898–900.
- 39. Requia WJ, Coull BA, Koutrakis P. The impact of wildfires on particulate carbon in the western USA. Atmos Environ. 2019;213:1–10.
- 40. Reiss R, Anderson EL, Cross CE, Hidy G, Hoel D, McClellan R, et al. Evidence of health impacts of sulfate-and nitrate-containing particles in ambient air. Inhalation Toxicol. 2007;19:419–49.
- Daellenbach KR, Uzu G, Jiang J, Cassagnes L-E, Leni Z, Vlachou A, et al. Sources of particulate-matter air pollution and its oxidative potential in Europe. Nature. 2020:587:414–9.
- 42. O'Halloran AC, Holstein R, Cummings C, Kirley PD, Alden NB, Yousey-Hindes K, et al. Rates of influenza-associated hospitalization, intensive care unit admission, and in-hospital death by race and ethnicity in the United States from 2009 to 2019. JAMA Netw Open. 2021;4:e2121880.
- 43. Jbaily A, Zhou X, Liu J, Lee T-H, Kamareddine L, Verguet S, et al. Air pollution exposure disparities across US population and income groups. Nature. 2022;601:228–33.

### **AUTHOR CONTRIBUTIONS**

The conceptualization of the study was performed by DPC, MJU, SL, SWT, PKH and DQR. The team that participated in data curation included HL, SL, YC, PKH, DQR, KTM, CJJ, CY and MRI. The formal analysis was performed by DPC, HL, ST, SL, SWT and DQR. The methodology for the study was led by DQR, SWT, DPC, SL and PKH. The supervisors of the project included DQR, PKH and SL. The original draft was completed by DPC, DQR, PKH and MJU, while all authors contributed to the subsequent writing, review & editing. The team responsible for acquiring funding included: DQR, PKH, SL, SWT and MJU.

### **FUNDING**

Funding for this project was provided by New York State Energy Research and Development Authority (NSYERDA) contracts #156226 and #125993 as well as the NIH environmental health science center P30 center grant #P30 ES001247. Dr. Croft was funded by the NIEHS mentored career development grant #K23 ES032459.

### **COMPETING INTERESTS**

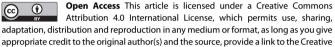
The authors declare no competing interests.

### ADDITIONAL INFORMATION

Correspondence and requests for materials should be addressed to Daniel P. Croft.

Reprints and permission information is available at http://www.nature.com/reprints

**Publisher's note** Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.



Commons licence, and indicate if changes were made. The images or other third party material in this article are included in the article's Creative Commons licence, unless indicated otherwise in a credit line to the material. If material is not included in the article's Creative Commons licence and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this licence, visit <a href="http://creativecommons.org/licenses/by/4.0/">http://creativecommons.org/licenses/by/4.0/</a>.

© The Author(s) 2024

<sup>1</sup>Pulmonary and Critical Care Division, Department of Medicine, University of Rochester Medical Center, Rochester, NY, USA. <sup>2</sup>Department of Environmental Medicine, University of Rochester Medical Center, Rochester, NY, USA. <sup>3</sup>Department of Public Health Sciences, University of Rochester Medical Center, Rochester, NY, USA. <sup>4</sup>Institute for a Sustainable Environment, Clarkson University, Potsdam, NY, USA. <sup>5</sup>Population Studies and Training Center, Brown University, Providence, RI, USA. <sup>6</sup>Department of Environmental Health Sciences. University at Albany, the State University of New York, Albany, NY, USA. <sup>7</sup>Department of Biostatistics and Computational Biology, University of Rochester Medical Center, Rochester, NY, USA. <sup>8</sup>Department of Pediatrics, University of Rochester, Rochester, NY, USA. <sup>8</sup>Department of Pediatrics, University of Rochester, Rochester, NY, USA. <sup>8</sup>Department of Pediatrics, University of Rochester, Rochester, NY, USA. <sup>8</sup>Department of Pediatrics, University of Rochester, Rochester, NY, USA. <sup>8</sup>Department of Pediatrics, University of Rochester, Rochester, NY, USA. <sup>8</sup>Department of Pediatrics, University of Rochester, NY, USA.