



Upstream oil and gas production and ambient air pollution in California

David J.X. Gonzalez^{a,b,*}, Christina K. Francis^c, Gary M. Shaw^d, Mark R. Cullen^{e,1}, Michael Baiocchi^f, Marshall Burke^g

^a Department of Environmental Science, Policy and Management and School of Public Health, University of California, Berkeley, Berkeley, CA 94720, USA

^b Emmett Interdisciplinary Program in Environment and Resources, Stanford University, Stanford, CA, USA

^c Program in Environmental Science and Studies, Johns Hopkins University, Baltimore, MD, USA

^d Department of Pediatrics, Stanford University, Stanford, CA, USA

^e Founding Director of the Stanford Center for Population Health Sciences, USA

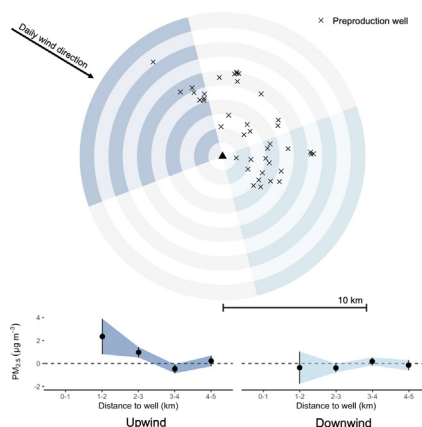
^f Department of Epidemiology and Population Health, Stanford University, Stanford, CA, USA

^g Department of Earth System Science, School of Earth, Energy and Environmental Sciences, Stanford University, Stanford, CA, USA

HIGHLIGHTS

- Oil and gas wells have been linked to adverse health, but mechanisms not well understood.
- Applied a quasi-experimental design with daily air pollution and oil production data
- We leveraged wind direction as source of exogenous variation for exposure to wells.
- Upstream oil and gas production emitted air pollutants at concentrations that may be harmful.
- Evaluated proximity as an appropriate indicator of air pollution exposure from wells

GRAPHICAL ABSTRACT



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ABSTRACT

Background: Prior studies have found that residential proximity to upstream oil and gas production is associated with increased risk of adverse health outcomes. Emissions of ambient air pollutants from oil and gas wells in the preproduction and production stages have been proposed as conferring risk of adverse health effects, but the extent of air pollutant emissions and resulting nearby pollution concentrations from wells is not clear.

Objectives: We examined the effects of upstream oil and gas preproduction (count of drilling sites) and production (total volume of oil and gas) activities on concentrations of five ambient air pollutants in California.

Methods: We obtained data on approximately 1 million daily observations from 314 monitors in the EPA Air Quality System, 2006–2019, including daily concentrations of five routinely monitored ambient air pollutants: PM_{2.5}, CO, NO₂, O₃, and VOCs. We obtained data on preproduction and production operations from Enverus and the California Geographic Energy Management Division (CalGEM) for all wells in the state. For each monitor and each day, we assessed exposure to upwind preproduction wells and total oil and gas production volume within 10 km. We used a panel regression approach in the analysis and fit adjusted fixed effects linear regression models for each pollutant, controlling for geographic, seasonal, temporal, and meteorological factors.

* Corresponding author at: 108 Hilgard Hall, University of California, Berkeley, Berkeley, CA 94720, USA.

E-mail address: djgonz@berkeley.edu (D.J.X. Gonzalez).

¹ Retired.

Results: We observed higher concentrations of PM_{2.5} and CO at monitors within 3 km of preproduction wells, NO₂ at monitors at 1–2 km, and O₃ at 2–4 km from the wells. Monitors with proximity to increased production volume observed higher concentrations of PM_{2.5}, NO₂, and VOCs within 1 km and higher O₃ concentrations at 1–2 km. Results were robust to sensitivity analyses.

Conclusion: Adjusting for geographic, meteorological, seasonal, and time-trending factors, we observed higher concentrations of ambient air pollutants at air quality monitors in proximity to preproduction wells within 4 km and producing wells within 2 km.

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1. Introduction

Recent studies have found that residing in proximity to oil and gas wells is associated with adverse cardiovascular, psychological, perinatal, and other health outcomes (Casey et al., 2015, 2018; Currie et al., 2017; Denham et al., 2021; McKenzie et al., 2014, 2018, 2019; Tang et al., 2020; Whitworth et al., 2017; Willis et al., 2021). Studies in California have found higher risk of preterm birth and low birthweight with exposure to upstream oil production, as well as impaired lung function and higher asthma prevalence (Gonzalez et al., 2020; Johnston et al., 2021; Shamasunder et al., 2018; Tran et al., 2020). Several possible mechanisms have been hypothesized for the observed associations between proximity to wells and adverse health outcomes, including emissions of ambient air contaminants during various stages of upstream oil and gas production (Adgate et al., 2014; Allshouse et al., 2019; Gonzalez et al., 2020; Johnston et al., 2019; McKenzie et al., 2012). There is a potential for widespread risk of exposure to air pollutant emissions from upstream oil and gas development, with an estimated 17.6 million U.S. residents, including 2.1 million Californians, living within 1.6 km (1 mile) of at least one active well (Czolowski et al., 2017).

Despite widespread potential exposure to wells and reported health risks, the effects of upstream oil and gas production on ambient air quality are still not well understood (Johnston et al., 2019). Under the Clean Air Act and its amendments, local regulatory agencies are responsible for maintaining networks of in situ air pollution monitors (Grainger et al., 2017). Agencies routinely monitor criteria air pollutants, which are statutorily regulated under the Clean Air Act and which include fine particulate matter with an aerodynamic diameter less than 2.5 µm (PM_{2.5}), carbon monoxide (CO), nitrogen dioxide (NO₂), and ozone (O₃). Other hazardous pollutants are also routinely monitored, including non-methane volatile organic compounds (VOCs) such as acetaldehyde, benzene, ethylbenzene formaldehyde, n-hexane, toluene and xylene. In prior studies, such as in situ monitoring campaigns conducted in California, Colorado, and Texas, investigators have reported elevated concentrations of PM_{2.5}, CO, NO₂, O₃, and VOCs near wells (Allshouse et al., 2019; Arbelaez and Baizel, 2015; Garcia-Gonzales et al., 2019a; Schade and Roest, 2016, 2018). Sources of PM_{2.5} emissions associated with upstream oil and gas production may include combustion of diesel fuel from on-site equipment and heavy trucks, dust from construction sites and unpaved roads, and secondary formation in the atmosphere (Adgate et al., 2014); emissions of CO and NO₂ may also be associated with fossil fuel combustion in vehicles and off-road equipment (Holloway et al., 2000; Jackson et al., 2014); O₃ may be formed as a secondary pollutant in photochemical reactions involving nitrous oxides (such as NO₂) and VOCs in the presence of sunlight (Mauzerall et al., 2005; Rodriguez et al., 2009).

Studies have found elevated concentrations of harmful pollutants near oil and gas wells (Garcia-Gonzales et al., 2019b). However, prior studies have been geographically and temporally constrained and often do not mirror methods applied by population health researchers. In particular, exposure characterization is often spatial in nature, whereas population health researchers often seek to exploit temporal variation to isolate the role of exposure to oil and gas wells from exposure to other spatially correlated activities that may affect pollution and health (Currie et al., 2017; Willis et al., 2021). Additionally, the unique

geological conditions of California may constrain external validity of air quality studies that investigate oil and gas production-related emissions in other settings (Garcia-Gonzales et al., 2019a). Population health studies investigating exposure to upstream oil and gas production typically use proximity to wells as the indicator of exposure without directly measuring concentrations of air pollutant emissions or other potential hazards, such as noise and water pollution (Casey et al., 2015; Currie et al., 2017; Gonzalez et al., 2020; McKenzie et al., 2014; Rasmussen et al., 2016; Tang et al., 2020; Tran et al., 2020; Willis et al., 2021). Improved understanding of pollutants emitted during upstream oil and gas production, including the classes of pollutants emitted (or secondarily produced) and the distances to which they are transported could help population health scientists more accurately parameterize exposure assessments and determine which aspects of exposure to production activities may adversely affect human health.

In our prior study (Gonzalez et al., 2020), we found that proximity to wells was associated with higher preterm birth risk, but we were not able to measure specific chemical pollutants parents were potentially exposed to during their pregnancy, or to separate proximity to wells from other activities that may also affect preterm birth risk. Our objectives in the current study were to examine how upstream oil preproduction and production activities affected ambient air quality in California from 2006 to 2019, with the aim of informing population health studies of exposure to upstream oil and gas production. We investigated whether marginal changes in preproduction and production activities resulted in increased concentrations of PM_{2.5}, CO, NO₂, O₃, and VOCs. Where we observed marginal increases in pollutant concentrations with proximity to wells, we also aimed to determine the distance at which elevated concentrations decay to background levels. To address these objectives, we applied a quasi-experimental design using a panel of publicly available air quality monitoring data.

2. Methods

2.1. Study design

We constructed a panel dataset with repeated daily measures of ambient air pollutant concentrations as well as upstream oil and gas production across California from January 1, 2006, to December 31, 2019. We made use of geospatial and temporal variation in oil and gas extraction activities, including well preproduction (defined as the interval between spudding, or initiation of drilling, and completion) and production (total monthly volume of oil and gas produced), and leveraged daily variation in wind direction as a source of exogenous variation. The type and magnitude of emissions may vary by stage due to differences in activities related to preproduction and production, and the intensity of well pad activity varies within each stage (Allshouse et al., 2017). For each monitor, we assessed daily exposure to upwind wells in preproduction and production during the study period. In the current study, we did not assess exposure of any human populations; rather, we assessed exposure of air quality monitors as a surrogate receptor. Then we used a fixed effects regression approach to assess the effect of exposure to preproduction and producing wells on the concentrations of each pollutant, accounting for geographic, seasonal, and time-trending, and meteorological factors.

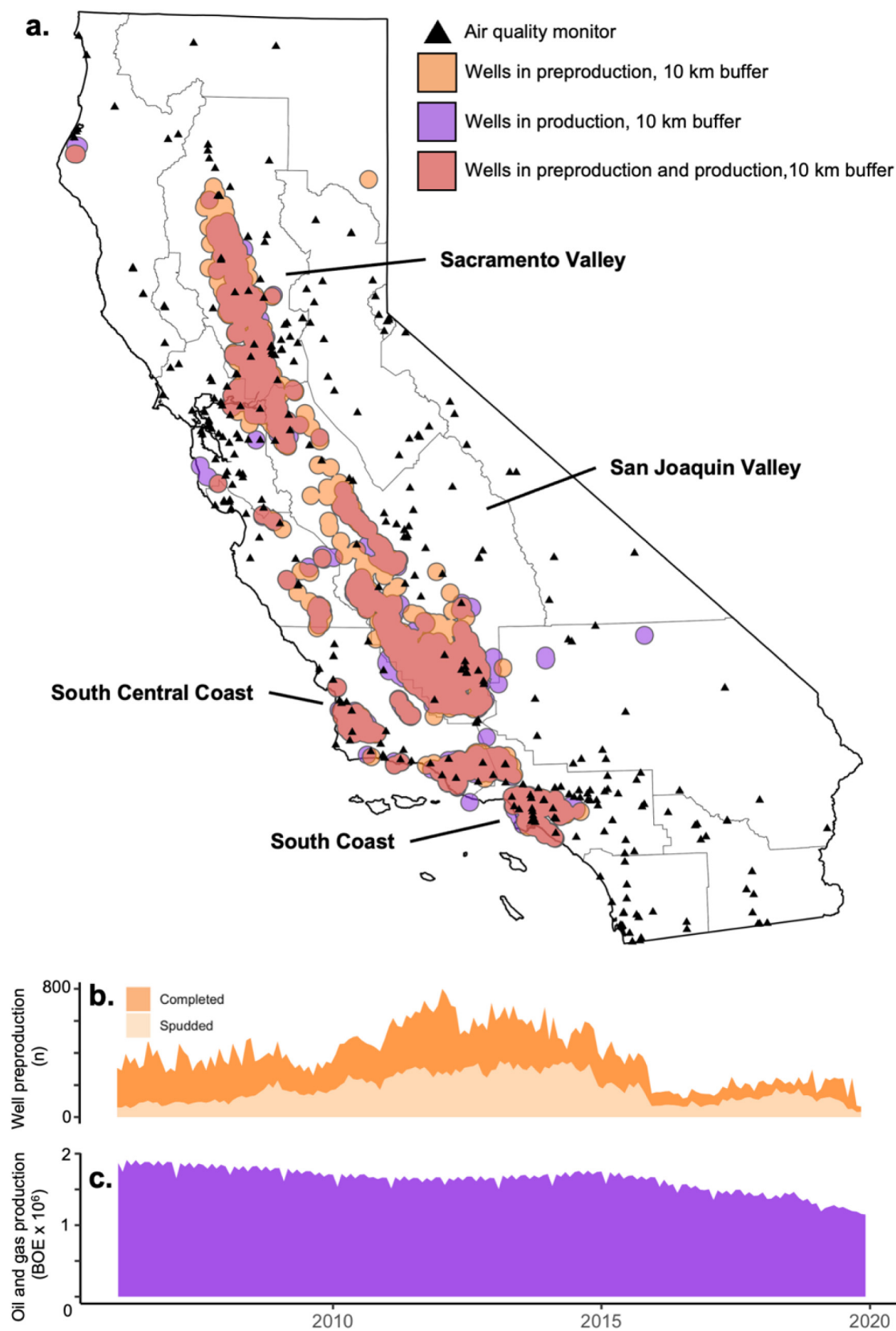


Fig. 1. (a) A map of the study region, showing air basins, air quality monitor locations, and 10 km buffers around wells in preproduction (orange) and production (purple), as well as the overlap (red). (b) Count of wells spudded and completed by month across California, including recompletions of previously drilled wells. (c) Total oil and gas production by month for all wells in California, reported as million barrels of oil equivalent (BOE).

2.2. Data

We obtained air quality data from the U.S. Environmental Protection Agency (EPA) Air Quality System (AQS). This dataset comprised daily measurements of seven air pollutants, with daily mean concentrations of PM_{2.5} ($\mu\text{g m}^{-3}$) as well as daily max concentrations of CO (ppm), NO₂ (ppb), O₃ (ppb), and non-methane VOCs (ppb C). In all analyses, the unit of observation was the pollutant concentration at each monitor for each day, or the monitor-day. We included data for all 314 AQS monitors in California that were operating during the study period and that monitored for the five pollutants of interest (Fig. 1). Missing air pollution data were omitted from the analyses; we did not impute missing air pollution data. Due to the sparse monitoring of VOCs compared to other pollutants, we included data on VOC measurements for 1999–2005; we excluded pre-2006 measurements for other pollutants because data for wildfire smoke plumes, described below, were not available before 2006. Air quality monitors detected and measured non-methane VOC concentrations via the EPA Method TO-3 for ethylbenzene, n-hexane, toluene, benzene, and ethylene using cryogenic preconcentration techniques, gas chromatography, and flame ionization detection. Xylene concentrations were estimated using preconcentration techniques, gas chromatography, and Saturn 2000 ion mass spectrometry. Acetaldehyde and formaldehyde concentrations were measured using 2,4-dinitrophenylhydrazine (DNPH) silica gel cartridges, an O₃ scrubber, and ultraviolet absorption spectroscopy.

Data on the oil and gas wells, including development dates and monthly production volume, was obtained from the California Geologic Energy Management Division (CalGEM) and Enverus, a private data aggregation service. The analytic dataset included 38,157 wells that were in the preproduction and 90,697 wells in production in California during the study period (Table S1). We defined the preproduction stage of the well as starting with the reported spud date (when drilling begins) and ending with the completion date. We assessed monitors as exposed to proximate preproduction wells on days when the well was between the dates of spudding and completion. Preproduction wells were included in the study if the preproduction interval (spudding to completion) occurred during the study period. For wells with missing data for spud date, we assumed that the preproduction interval began 30 days before completion; for wells missing completion date, we assumed the preproduction stage ended 30 days after spudding. Wells missing both spud and completion dates were assumed to have been drilled outside the study period; since the record dates to the late 19th century, we expected there to be missingness in these variables for wells drilled prior to 1999. Wells in the production stage were included for all sites with any reported oil or gas production during the study period. Because oil and gas are frequently produced from the same wells, we used a combined metric of oil and gas production reported as barrels of oil equivalent (BOE). The dataset comprised 8,064,549 well-month observations of a total of approximately 3.8 billion BOE.

We obtained meteorological data from the North American Regional Reanalysis (NARR), a product developed by the National Centers for Environmental Prediction. This dataset included modeled daily mean wind direction and speed, reported as vectors (u and v), as well as observations of mean daily surface temperature ($^{\circ}\text{C}$) and total daily precipitation (mm). There were no missing estimates for these meteorological variables. We also obtained administrative shapefiles for air basins across the state from the California Air Resources Board (CARB). We used data from the 2010 decennial census to determine whether monitors were located in urban areas (with 50,000 or more residents) or urban clusters (with 2500–50,000 residents) compared with rural areas, which comprise all other areas. To control for potential effects of wildfire smoke on daily concentrations ambient air pollutants, we used data on the daily location of wildfire smoke plumes from the Hazard Mapping System of the National Oceanic and Atmospheric Administration (NOAA), which assessed the number of overhead smoke plumes at the zip code level (Schroeder et al., 2008).

2.3. Exposure assessment

We constructed a panel dataset where, for each monitor and each day with a pollutant observation, we summed (a) the number of upwind wells in preproduction and (b) the total volume of upwind oil and gas production (BOE) in 1 km increments out to 10 km (Fig. 2). We determined the wind direction for each monitor and day from the u and v vector components from the NARR wind product. The resultant of the u and v vector components conveys wind direction and speed (magnitude). Preproduction and production wells that intersected the upwind quadrant on each day for each monitor comprised the primary exposure variables; wells outside the quadrant were excluded in the primary analyses.

As sensitivity analyses, we also assessed exposure to wells in the downwind quadrant as a placebo exposure. Additionally, we assessed exposure to all preproduction wells and production volume in 1 km annuli (or rings) radiating out from the monitor, i.e., without taking wind into account.

The receptor in our exposure assessment was the air quality monitor; this study did not consider any human receptors or health outcomes. Our aim was to use air monitors as a proxy for the residential receptors typically targeted in population health studies that assess exposure to oil and gas wells.

2.4. Identification strategy

We leveraged daily variation in wind direction as a plausibly exogenous source of variation, uncorrelated with well preproduction and production activities as well as other sources of pollution. This strategy allowed us to, by design, isolate the marginal contributions of additional preproduction wells and production volume to ambient air pollutant concentrations.

2.5. Statistical analyses

We used adjusted fixed effects linear regression models to assess how marginal changes in (a) the count of wells in preproduction or (b) the volume of oil and gas production affects concentrations of each observed pollutant (PM_{2.5}, CO, NO₂, O₃, and VOCs). For each combination of pollutants and well stage (preproduction or production), we fit the following model:

$$Y_{md} = U_{mda} + D_{mda} + O_{mda} + C_{md} + \gamma_{md} + \delta_{by} + e_{md},$$

where Y is the observed daily concentration of the pollutant at monitor m on day d ; U is a vector of either the (a) upwind count preproduction wells or (b) upwind sum oil and gas production on day d in annulus a (0–1 km, 1–2 km, ... 9–10 km) radiating from monitor m ; D is similar to U but for downwind wells; O is also similar to U , but were wells in the two quadrants orthogonal to the upwind quadrant (i.e., lateral wells); C is a vector of covariates (day of week, precipitation in mm, temperature in $^{\circ}\text{C}$, wind speed in ms^{-1} , and the count of overhead smoke plumes) at monitor m on day d ; γ is a fixed effect for monitor by month, n ; δ is a fixed effect for air basin, b , by year, y ; and e is an error term representing unmodeled sources of variation in pollution at monitor m on day d . We fit additional models with polynomial terms for each exposure bin to examine whether the response was nonlinear.

We compared the point estimates for upwind wells with downwind placebos. As sensitivity analyses, we also modified the fixed effects in the model, using monitor-by-year and air basin-by-month-by-year fixed effects in the model. Additionally, we fit models as described above in the primary analysis but using exposure assessment data that did not take wind into account (i.e., the sum of all preproduction wells or production volume within each annulus). Finally, as an additional sensitivity analysis for co-exposure to wildfire smoke, we fit models

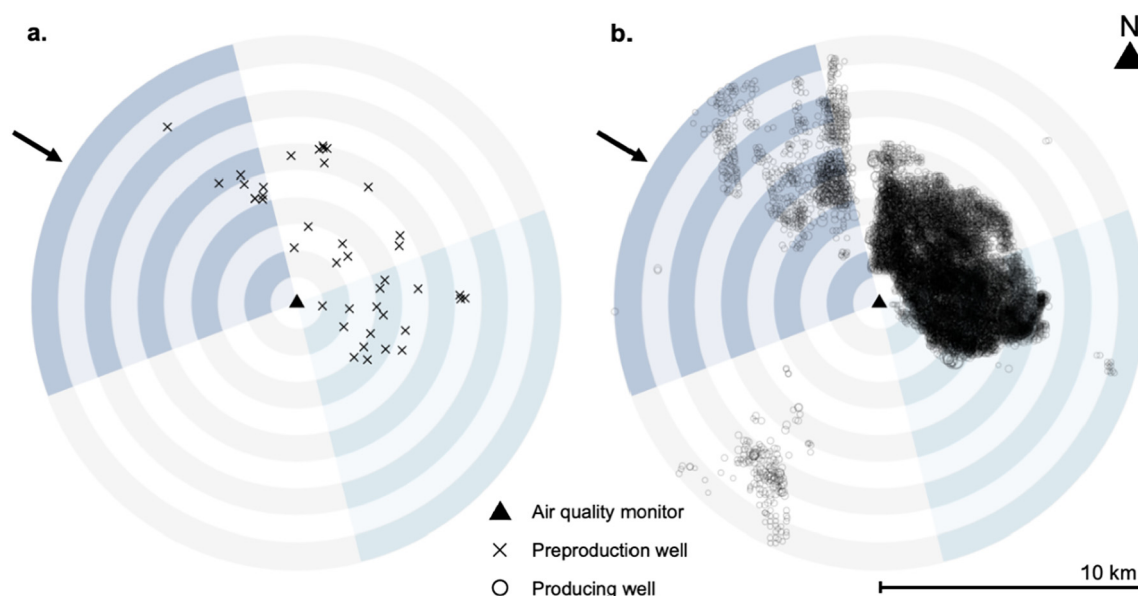


Fig. 2. A visualization of the exposure assessment method at a monitor located in Bakersfield, California, using sample data from July 1, 2009, when the wind was blowing from the northwest (arrow). For each monitor-day, we assessed exposure to (a) the count of wells in preproduction and (b) the total volume of oil and gas produced upwind (darker shaded area) of the monitor. As a placebo test, we assessed exposure to wells downwind (lighter shade) of the monitor.

for $PM_{2.5}$ where monitor-day observations that had >0 smoke plumes overhead were omitted.

In total we fit 27 models, and, as the primary analysis, we focused on the adjusted fixed effects regression models for exposure to preproduction wells and production volume. In particular, we were interested in the point estimates for exposure to upwind wells and production within 5 km of the monitor.

All data preparation and analyses were conducted using R v. 4.0 (R Core Team, 2020).

3. Results

3.1. Descriptive statistics

The analytic dataset comprised 1,058,230 daily observations of the five pollutants from 314 monitors across California collected from 2006 to 2019, with additional observations for VOCs from 1999 to 2005 (Table 1). Most (208) monitors were located in urban areas and approximately half (158) were in the four air basins with the majority of oil and gas wells (96.4%) and production (87.2%): Sacramento Valley, San Joaquin Valley, South Central Coast, and South Coast (Table S1). Not all monitors collected data for all pollutants. The majority (79.5%) of monitor-days included observations for O_3 , with 43% of monitor-days including data for NO_2 and $PM_{2.5}$. Some 31% of monitor-days included CO observations and 8.9% included observations of VOCs. Among the 94,349 monitor-days with an observation for VOCs, 39.3% were in the San Joaquin Valley and 12.8% were in the South Coast basin, both basins where most oil and gas wells were concentrated. For each pollutant, there were more observations at monitors more than 10 km from wells than monitors near wells. More observations were collected in the later years of the study period compared to earlier in the study period. The number of monitors in operation throughout the study period was relatively consistent from year to year; the minimum number of monitors in operation was 223 in 2006 and the maximum was 245 in both 2012 and 2014, with a median of 239 (Fig. S4). The number of monitors that assessed $PM_{2.5}$ concentrations increased throughout the study period. Concentrations of pollutants at monitors within 10 km of wells were similar to the concentrations at monitors further away (Table 1).

Wells in all production stages were concentrated in the San Joaquin Valley, which includes Kern County, with substantial production in the South Coast air basin, which includes Los Angeles County (Table S1). Among the 314 monitors included in the analytic dataset, 79 (25.2%) were within 10 km of at least one oil or gas well, 33 (10.5%) were within 3 km, and 11 (3.5%) were within 1 km. Of the monitor-days included in the analysis, 46,477 (4.4%) were exposed to at least one preproduction or production well within 1 km, 115,648 (10.9%) were within 3 km, and 239,764 (22.7%) were within 10 km. For monitor-days with data for $PM_{2.5}$ and VOCs, there were no preproduction wells within 1 km.

Among exposed monitor-days, the median number of preproduction wells within each upwind 1-km bin was between 1 and 4, with a maximum of 41 (Table S2). For producing wells, median upwind exposure spanned 7.2 to 166.9 BOE, with a right-skew and a maximum of 24,166.1 BOE. There was both seasonal and geographic variation in wind direction: in the San Joaquin Valley, the wind predominantly originated in the northwest; in the South Coast basin, wind predominantly came from the southwest (Fig. S1). Exposure to preproduction wells was correlated with exposure to production volume for all annuli beyond 1 km. Across producing wells, daily production volume was right-skewed, with a median of 7.3 BOE per day and mean ($\pm$ SD) of $17.1 (\pm 50.6)$ BOE per day. Exposure to preproduction wells was highly correlated for adjacent annuli and moderately correlated with further annuli; we observed a similar trend for production volume (Table S3). Exposure to preproduction wells was moderately correlated with exposure to production volume at distances greater than 1 km from wells.

3.2. Primary analyses

In the primary analysis, we observed increased concentrations of $PM_{2.5}$, CO, NO_2 , and O_3 with proximity to preproduction wells (Fig. 3). For $PM_{2.5}$, we observed an increase of $2.35 \mu g m^{-3}$ (95% CI: 0.81, 3.89) for each additional upwind preproduction well site within 2 km of the monitor, and $0.97 \mu g m^{-3}$ (0.52, 1.41) for an additional well between 2 and 3 km from the monitor. For CO, we observed an increase of 0.09 ppm (-0.0004, 0.18) with an additional upwind well within 2 km and 0.02 (0.004, 0.032) for a well at 2-3 km. Concentrations of NO_2 increased 2.27 with well at 0-1 km, 2.91 (0.99, 4.84) for a well at 1-2 km, and 0.65 (0.31, 0.99) for a well at 2-3 km upwind. For O_3 , there

Table 1

Descriptive statistics of the air monitors, pollutant concentrations, and meteorological factors during the study period, 2006–2019. The unit of observation is the monitor-day; some monitors observe multiple pollutants. VOCs in the dataset comprise non-methane volatile organic compounds.

	≤ 10 km to wells	> 10 km to wells	All
Monitors, <i>n</i> (column %)	79 (25.2)	235 (74.8)	314 (100)
Urban	57 (72.2)	151 (64.3)	208 (66.2)
Rural	22 (27.8)	84 (35.7)	106 (33.8)
Sacramento Valley	16 (20.2)	26 (11.1)	42 (26.6)
San Joaquin Valley	18 (22.8)	24 (10.2)	42 (26.6)
South Central Coast	15 (19.0)	14 (6.0)	29 (18.4)
South Coast	15 (19.0)	30 (12.8)	45 (28.5)
PM _{2.5}	43 (54.4)	155 (66.0)	198 (63.1)
CO	34 (43.0)	76 (32.3)	110 (35.0)
NO ₂	45 (57.0)	94 (40.0)	139 (44.3)
O ₃	65 (82.3)	172 (73.2)	237 (75.5)
VOCs	24 (30.4)	24 (10.2)	48 (15.3)
Observations, <i>n</i> (column %)	307,095 (29.0)	751,135 (71.0)	1,058,230 (100)
Urban	214,011 (69.7)	507,287 (67.5)	721,298 (68.2)
Rural	93,084 (30.3)	243,848 (32.5)	336,932 (31.8)
PM _{2.5}	137,657 (44.8)	317,065 (42.2)	454,722 (43.0)
CO	98,165 (32.0)	229,646 (30.6)	327,811 (31.0)
NO ₂	157,567 (51.3)	297,197 (39.6)	454,764 (43.0)
O ₃	252,572 (82.2)	588,448 (78.3)	841,020 (79.5)
VOCs ^a	44,992 (14.7)	49,357 (6.6)	94,349 (8.9)
2006–2009	77,013 (25.1)	200,404 (26.7)	277,417 (26.2)
2010–2014	104,839 (34.1)	264,066 (35.2)	368,905 (34.9)
2015–2019	107,248 (34.9)	268,876 (35.8)	376,124 (35.5)
Smoke plume overhead	21,780 (7.1)	54,299 (7.2)	76,079 (7.2)
Pollutant concentrations, daily mean ± SD			
PM _{2.5} (µg/m ³)	10.6 ± 9.5	9.9 ± 9.0	10.1 ± 9.1
CO (ppm)	0.5 ± 0.4	0.5 ± 0.4	0.5 ± 0.4
NO ₂ (ppb)	21.4 ± 14.6	22.1 ± 14.5	21.9 ± 14.5
O ₃ (ppm)	0.04 ± 0.01	0.04 ± 0.02	0.04 ± 0.02
VOCs (ppb C)	120 ± 166	104 ± 142	112 ± 155
Meteorological factors, daily mean ± SD			
Precipitation (mm)	0.9 ± 4.0	1.2 ± 5.1	1.1 ± 4.8
Temperature (°C)	18.6 ± 7.8	17.2 ± 9.1	17.6 ± 8.8
Wind speed (m/s)	3.0 ± 2.1	3.2 ± 2.0	3.1 ± 2.0

^a The data for VOCs includes observations for 1999–2019.

were no significant changes for an additional well within 2 km, an increase of 0.31 (0.20, 0.42) with an additional well at 2–3 km, and an increase of 0.14 (0.05, 0.23) with a well at 3–4 km. There were no increases in concentration with upwind exposure to VOCs, though notably there was no exposure to preproduction wells within 1 km. Across all pollutants, we did not observe any substantial increased concentrations beyond 4 km. In the placebo test, with exposure assessed to downwind wells, we did not observe any substantial increases in pollutant concentrations.

We observed increased concentrations of PM_{2.5}, NO₂, O₃, and VOCs with higher exposure to upwind production (Fig. 4). We estimated the marginal effect of exposure to an additional 100 BOE of daily total oil and gas volume within each 1-km annulus. This degree of exposure roughly corresponds with median upwind production volume within each annulus among exposed monitor-days (Table S2) and is comparable to cutoffs used in recent population health work (Tran et al., 2020). For each additional 100 BOE of total oil and gas production within 1 km, we observed an increase of 1.93 µg m⁻³ (95% CI: 1.08, 2.78) in the concentration of PM_{2.5}. For NO₂, we observed an increase of 0.62 ppb (0.37, 0.86) with an additional 100 BOE within 1 km. The concentration of O₃, increased by 0.11 ppb (0.08, 0.14) with for each 100 additional BOE at 1–2 km. There was an increase in VOC concentrations of 0.04 (0.01, 0.07) ppb C for an additional 100 BOE of production within 1 km. We did not observe any substantial changes in CO concentrations with upwind exposure to production volume. In the downwind placebo tests, we observed an increase in PM_{2.5} concentrations for exposure to increased production within 1 km, a small increase in NO₂ concentrations at 1–2 km, and an increase in O₃ at 3–4 km.

3.3. Sensitivity analyses

We performed several sensitivity analyses. Fitting models that included exposure variables for both preproduction and production did not substantially change the results; point estimates and confidence intervals were similar in models with exposure variables for both preproduction and production compared to models examining each exposure separately (Fig. S4). In models with polynomial term for exposure we did not see evidence of non-linear responses to upwind exposure. Changing model specification in the primary analysis for preproduction wells (Table S4) or for production volume did not qualitatively change findings (Table S5). In a sensitivity analysis, we fit the model as described above but omitted the 35,422 monitor-days with smoke plumes overhead, comprising 7.8% of the PM_{2.5} analytic dataset. The results were similar to the smoke-adjusted results for exposure to wells in both the preproduction and production stages (Fig. S3).

4. Discussion

We observed higher concentrations of ambient air pollutants at air monitors exposed to wells in both the preproduction and production stages. Concentrations of PM_{2.5} were substantially higher on days when a well was in preproduction within 3 km of the monitor, and also when production volume increased within 1 km of the monitor. Notably, we observed increases in PM_{2.5} within 1 km of producing wells with and without considering wind direction. There are several possible explanations for this result: it may be attributable to high volume of producing wells near monitors in San Joaquin Valley orthogonal to the upwind direction, imperfect data on wind direction,

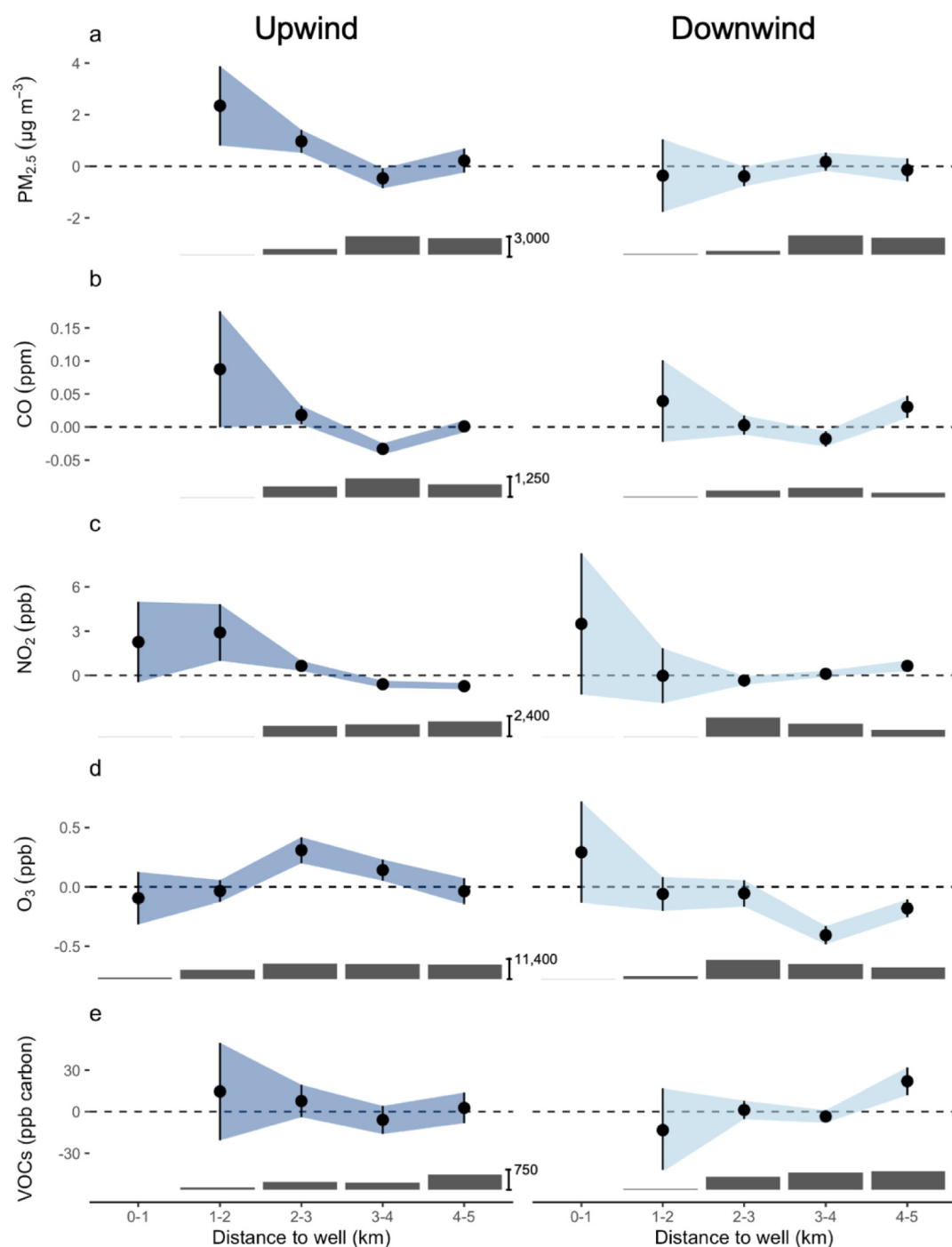


Fig. 3. Point estimates (95% CIs) for the marginal effect of one additional preproduction well upwind (left column) and downwind (right column) of the monitor. The bar plots show the number of monitor-days with exposure at least one preproduction well within each distance bin.

or shifts in wind direction during the day that were not adequately captured when we integrated wind direction over the course of a 24 h period. In addition to elevated PM_{2.5} levels, concentrations of O₃ increased when production activity increased between 1 and 4 km upwind of the monitor, but not for activity within 1 km of the monitor. This result may be attributable to secondary formation from primary pollutants emitted from during preproduction and production. Ground-level O₃ may be secondarily formed from photochemical reactions involving CO, NO_x, and VOCs, all of which we also observed were emitted from wells (Real et al., 2007; Rodriguez et al., 2009). We observed increased CO concentrations on days when preproduction wells were drilled within 3 km of the monitor.

Concentrations of NO₂ were higher on days when there was a preproduction well within 2 km or increased production volume within 1 km. For VOCs, we found higher concentrations when production volume increased within 1 km of the monitor. In the current study, VOCs comprised non-methane organic compounds including acetaldehyde, benzene, ethylene, and formaldehyde.

In models that considered both preproduction wells and production volume, we observed similar estimates to the models where we considered preproduction and production separately, as shown in Fig. S4. Preproduction activity near monitors was correlated with production volume, though this may not be apparent based on the correlation matrix in Table S3, which shows low correlation between preproduction

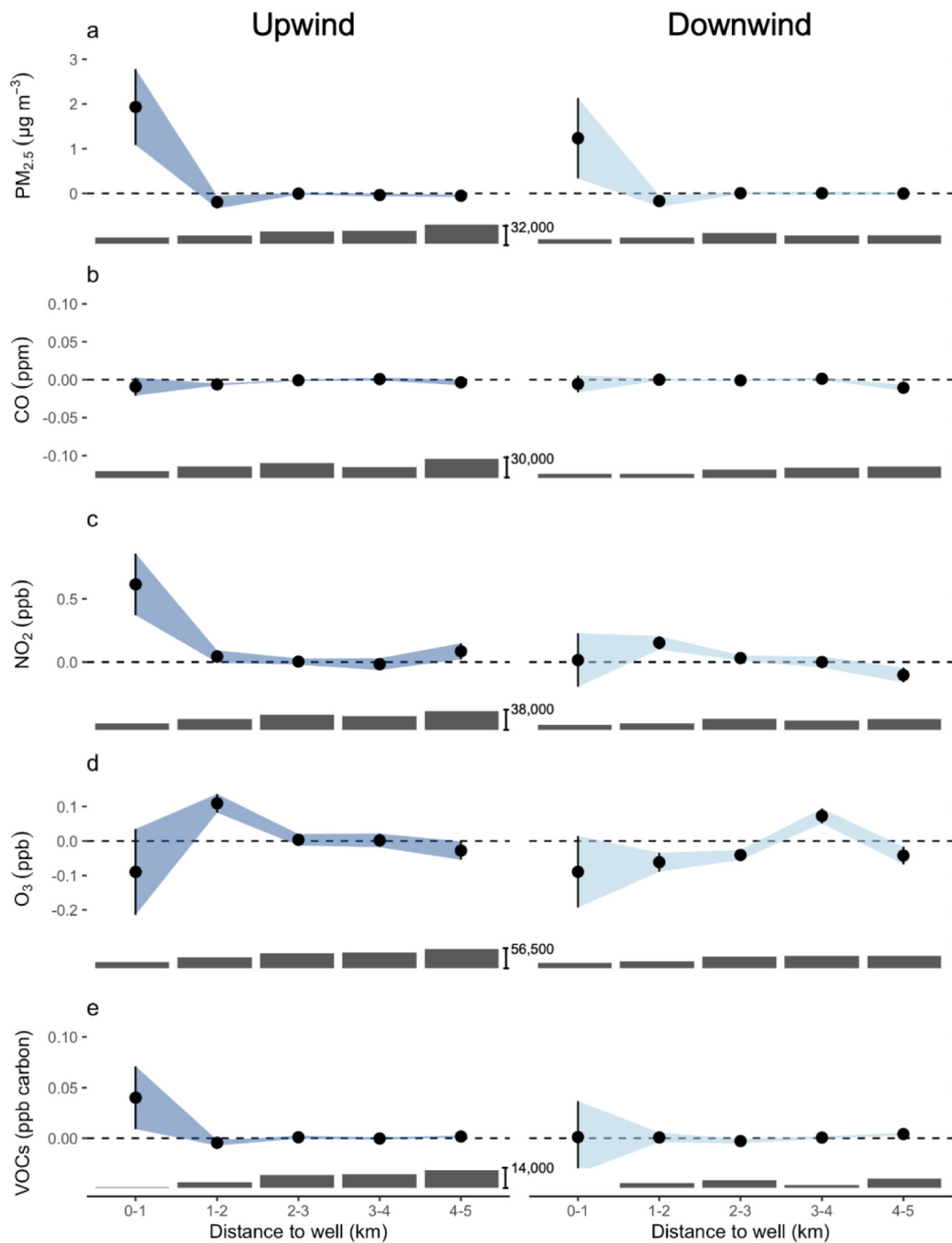


Fig. 4. Point estimates (95% CIs) for the marginal effect of 100 additional barrels of oil equivalent (BOE) of daily production volume, for wells upwind (left column) and downwind (right column) of the monitor. The bar plots show the number of monitor-days with exposure to at least 1 BOE of daily production volume within each distance bin. Note that more monitor-days had exposure to production volume than preproduction wells.

wells and production volume. However, among all monitor-days with a preproduction well within 1 km of the monitor, there was also >0 BOE of production volume.

In this study, we conducted a quasi-experimental analysis that relied on the existing network of air quality monitors. The siting of air quality monitors is delegated to local authorities and prior studies have found evidence of bias in where monitors are sited, which should be considered when interpreting the results from the current study (Grainger et al., 2017; Grainger and Schreiber, 2019). For example, in counties just marginally in attainment for National Ambient Air Quality Standards (NAAQS), regulators had an incentive to place new monitors far

from pollution sources, whereas in areas already in non-attainment, the regulators were incentivized to place monitors close to polluting sources (Grainger et al., 2017). This could lead to biased estimates of emissions from oil and gas wells, as monitors may be sited away from the most intensively producing oil fields. There is also evidence that monitors are less likely to be located in communities with racially and socioeconomically marginalized populations, which could lead to underestimation of oil and gas-related emissions if oil production in excluded areas was more intensive and polluting (Grainger and Schreiber, 2019). In the current study, the majority of oil and gas production was concentrated in Kern and Los Angeles Counties, both of which were in

non-attainment for PM_{2.5} throughout the study period (Environmental Protection Agency, 2021).

Findings from the current study indicate both primary emission and secondary formation of pollutants from upstream oil and gas production activities. However, identifying specific processes that resulted in observed pollutant emissions was outside the scope of the study.

4.1. Comparison to prior studies

Using proximity as a metric of exposure to upstream oil and gas production appears to adequately capture exposures to chemical contaminants. Proximity-based methods, such as inverse distance weighting or estimating production activity within 1 km of receptors, have been used in prior population health studies to estimate acute or chronic exposure to wells. The five pollutants we examined in this study represent a subset of potential hazards associated with exposure to oil and gas wells, which may include other air pollutants as well as water and noise pollution (Adgate et al., 2014; Jackson et al., 2014). Recent studies from California have reported fugitive methane from idle and unplugged wells, as well as urban oil and gas infrastructure, which may correlate with emissions of benzene, toluene, ethylene, xylene, and other air toxics (Lebel et al., 2020; Okorn et al., 2021). To differentiate risks conferred by air pollutants, population health researchers could utilize variations in wind direction.

Prior field studies have also found emissions of pollutants from upstream oil and gas facilities. A 2018 study in Texas found high concentrations of nitrous oxides and saturated hydrocarbons associated with oil and gas production in the Eagle Ford Shale (Schade and Roest, 2018). Another recent study in Colorado, which combined in situ monitoring and cancer risk assessment, found higher exposure to benzene and other non-methane hydrocarbons (toluene, ethylbenzene, and xylene) and elevated risk of cancer and other adverse health outcomes with close proximity to oil and gas facilities (McKenzie et al., 2018). Notably, the dataset in the current study did not include toluene, ethylbenzene, and xylene. Garcia-Gonzales et al. (2019a) found higher concentrations of VOCs downwind of a well site in Los Angeles. A study in Pennsylvania found that exposure metrics used in prior epidemiological studies were poorly correlated with observed pollutant concentrations (Wendt Hess et al., 2019). However, this study assessed exposure to wells at distances greater than 10 km, where we would not expect to detect increases in pollution, and the authors did not account for meteorological factors that may affect pollutant concentrations (Buonocore et al., 2020).

In prior studies, Tran et al. (2020) and Gonzalez et al. (2020) used differing proximity metrics to assess exposure to upstream oil and gas production and adverse birth outcomes in California. For their analysis of production volume and adverse birth outcomes, Tran et al. used a similar exposure assessment method to the one we employed in the current study, assessing “high” exposure to births with >100 BOE within 1 km of the residence. In the current analysis, we modeled exposure to production volume continuously rather than categorically. We found substantial increases in concentrations of PM_{2.5}, NO₂, and O₃ with exposure to an additional 100 BOE within 1 km, indicating that the metrics employed by Tran et al. likely were effective in capturing aspects of air pollution near active wells. Gonzalez et al. used inverse distance-squared weighting (IDW), a different approach that relies on the assumption that both density and proximity of wells confers risk of air pollution exposures. Notably, Gonzalez et al. (2020) conducted an exploratory analysis of the association between proximity to oil and gas wells, assessed using an IDW index, and concentrations of four pollutants (NO₂, O₃, PM₁₀, and PM_{2.5}). For that supplemental analysis, Gonzalez et al. also used data from EPA Air Quality System for mean monthly concentrations of air pollutants and fit fixed effects linear regression models estimating the effect of “high” exposure to wells (the highest tertile of the IDW index). These authors observed substantially higher concentrations of PM₁₀ and PM_{2.5}, lower concentrations of NO₂, and no substantial changes for O₃; for all

pollutants, effects. This indicates that the IDW method may be less effective as an exposure metric for the air pollutants investigated in this study than the methods employed in the current study. Additionally, the approaches in both Tran et al. (2020) and Gonzalez et al. (2020) may not adequately capture exposure to secondary pollutants such as O₃, which in the current study had higher concentrations several km downwind of wells.

4.2. Limitations and strengths

The current study had several limitations. We relied on daily changes in wind direction as a source of exogenous variation. On days with variable wind direction, estimating mean wind direction integrated over the course of the day could lead to exposure misclassification if, for example, wind blew from multiple directions during the course of a 24-h period. Data for many pollutants that may be emitted during upstream oil and gas production operations are not routinely monitored and reported in the EPA Air Quality System. Consequently, the results of the current study likely reflect only a subset of pollutants potentially emitted from upstream oil and gas production. Population health studies referring to our estimates of chemical contaminant exposure should consider the possibility of co-exposures to additional pollutants emitted during oil and gas production. We also did not have sufficient data to investigate specific VOC constituents, which may be associated with particular health endpoints of interest. Additionally, there were relatively few monitor-days with exposure to preproduction wells within 1 km. None of the monitors that measure concentrations of PM_{2.5} and VOCs were within 1 km of a preproduction well. We found evidence that drilling sites up within 1 to 3 km of air monitors increased PM_{2.5} concentrations, and concentrations of PM_{2.5} within 1 km of preproduction wells may be similar to or higher than our estimates for wells at 1–3 km. We did not expect to observe changes in VOC concentrations further than 1 km, as prior work has reported decay of VOCs within 100–200 m from well sites (Garcia-Gonzales et al., 2019a; Zielinska et al., 2014). Because of this, we were unable to make any inferences about the effect of preproduction activities on concentrations of VOCs.

In the primary analyses, we adjusted for exposure to wildfire smoke plumes to account for potential contributions of smoke to the pollutants of interest. Exposure was assessed as the number of overhead plumes for each monitor-day, but this method may not accurately indicate smoke conditions at ground level. A sensitivity analysis for PM_{2.5} omitting smoke days from the analysis yielded similar results to the smoke-adjusted models, suggesting that our statistical adjustment for smoke plumes was sufficient.

For the analyses of wells in the production stage, data on total oil and gas production volume were available at the monthly level. Because of this constraint, in the exposure assessment we assumed that production occurred evenly throughout the month. This could lead to exposure misclassification if production was concentrated in certain days of the month. Future researchers building on these findings should consider obtaining daily production volume data, if possible. Finally, we were not able to differentiate between drilling or production methods (i.e., conventional vs. unconventional methods, such as hydraulic fracturing), so we were not able to determine whether certain unconventional methods resulted in higher emissions.

Strengths of this study include the large panel dataset, comprising over 1 million daily observations from high quality air monitors with broad geographic and temporal variation. We were able to control for unobserved potential confounders through the study design, using wind as a plausibly exogenous source of variation uncorrelated to both upstream oil production and other sources of pollution. The monitor fixed effect accounts for average differences between monitoring locations, such as from pollution sources unrelated to oil and gas. Leveraging temporal variation from oil production activities and daily changes in wind direction accounts for other nearby pollution sources that are not both spatially collocated and temporally correlated with

oil and gas production. Based on this analytic approach, we think there is unlikely to be residual confounding. Additionally, we conducted several tests to validate the robustness of the results.

5. Conclusion

We conducted a quasi-experimental study to examine whether upstream oil and gas production results in emissions of ambient air pollutants. Adjusting for geographic, meteorological, seasonal, and time-trending factors, and leveraging daily changes in wind direction as an exogenous source of variation, we observed that proximity to oil and gas wells in both preproduction and production increased concentrations of PM_{2.5}, CO, NO₂, O₃, and VOCs at distances up to 4 km downwind of wells. These findings indicate that proximity to wells is an appropriate metric for air pollution-related exposures in population health studies. Notably, increases in PM_{2.5} concentrations near wells could be a mediating factor for previously reported increases in risk of adverse birth outcomes with proximity to wells in California (Bekkar et al., 2020; Gonzalez et al., 2020; Tran et al., 2020). Further research on hazards associated with upstream oil and gas production would improve understanding of potential health and environmental risks. Acute emissions of particular pollutants may be associated with specific steps of oil and gas preproduction or production, and more work is needed to determine if this is the case and, if so, which processes produce high emissions. Mitigating exposure to oil and gas wells would likely reduce exposure to ambient air pollutants.

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CRediT authorship contribution statement

David J.X. Gonzalez: Conceptualization, Data curation, Methodology, Formal analysis, Visualization, Writing – original draft, Writing – review & editing. **Christina K. Francis:** Data curation, Writing – original draft. **Gary M. Shaw:** Writing – review & editing. **Mark R. Cullen:** Methodology, Writing – review & editing. **Michael Baiocchi:** Methodology, Writing – review & editing. **Marshall Burke:** Conceptualization, Methodology, Supervision, Writing – review & editing.

Data availability

Data and code used in this analysis are available at <https://github.com/djxgonzalez/cal-drilling-air-quality>.

Declaration of competing interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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Appendix A. Supplementary data

Supplementary data to this article can be found online at <https://doi.org/10.1016/j.scitotenv.2021.150298>.

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