



Air quality and health-related impacts of traditional and alternate jet fuels from airport aircraft operations in the U.S.

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ABSTRACT

Aviation emissions from landing and takeoff operations (LTO) can degrade local and regional air quality leading to adverse health outcomes in populations near airports and downwind. In this study we aim to quantify the air quality and health-related impacts from commercial LTO emissions in the continental U.S. for two recent years' inventories, 2011 and 2016. We quantify the LTO-attributable PM_{2.5}, O₃, and NO₂ concentrations and health outcomes for mortality and multiple morbidity health endpoints. We also quantify the impacts from two scenarios representing a nation-wide implementation of 5% or 50% blends of sustainable alternative jet fuels. We estimate 80 (68–93) and 88 (75–100) PM_{2.5}-attributable and 610 (310–920) and 1,100 (570–1,700) NO₂-attributable premature mortalities in 2011 and 2016, respectively. We estimate a net decrease of 28 (14–56) and 54 (27–110) in O₃-attributable premature mortalities across the U.S. in 2011 and 2016, respectively due to the large O₃ titration effects near the airports. We also find that the asthma exacerbations due to NO₂ exposures from LTO emissions increase from 100,000 (2,500–200,000) in 2011 to 170,000 (4,400–340,000) in 2016. Implementing a 5% or 50% blend of sustainable alternative jet fuel in 2016 results in a 1% or 18% reduction, respectively in PM_{2.5}-attributable premature mortalities. Monetizing the value of avoided total premature mortalities, we find that a 50%-blended sustainable alternative jet fuel results in a 19% decrease in PM_{2.5} damages per ton of fuel burned and a 2% decrease in total damages per ton of fuel burned as compared to damages from traditional jet fuel. We also quantify health impacts by state and find California to be the most impacted by LTO emissions. We find that LTO-attributable PM_{2.5} and NO₂ premature mortalities increase by 10% and 80%, respectively from 2011 to 2016 and that NO₂-attributable premature mortalities are responsible for 91% of total LTO-attributable premature mortalities in both 2011 and 2016. And since we find LTO-attributable NO₂ to be unaffected by the implementation of sustainable alternative jet fuels, additional approaches focused on NO_x reductions in the combustor are needed to mitigate the air quality-related health impacts from LTO emissions.

1. Introduction

Commercial aviation emissions' impact on air quality has been estimated to be responsible for approximately 16,000 premature mortalities each year globally (Yim et al., 2015; Grobler et al., 2019). Both international and domestic forecasts for aviation traffic call for continued growth, 2.1% and 3.5% growth in revenue passenger miles over the next 20 years, respectively (FAA, 2014), with total fuel burn from commercial aircraft globally increasing by 71% between 1992 and 2006 (Olsen et al., 2013) and increasing by 28% between 2006 and 2015

(Grobler et al., 2019). Modeling studies have aimed to quantify air quality impacts and climate effects from all stages of an aircraft's flight trajectory, often looking at impacts at the landing and takeoff (LTO) stages (Arunachalam et al., 2011; Unal et al., 2005; Woody et al., 2011; Levy et al., 2012b; Lee et al., 2013; Woody and Arunachalam, 2013; Wolfe et al., 2014; Vennam et al., 2015; Woody et al., 2015; Woody et al., 2016; Penn et al., 2017) (below 3,000 ft) and full-flight (cruise) stages (Barrett et al., 2010; Barrett et al., 2012; Koo et al., 2013; Caiazzo et al., 2017; Cameron et al., 2017; Vennam et al., 2017; Grobler et al., 2019; Quadros et al., 2020; Lee et al., 2021).

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A recent paper by Dedoussi et al. 2021 (Dedoussi, 2021) has highlighted the important research questions surrounding aviation-attributable air pollution such as global impacts versus local impacts, aviation's impact in a changing atmosphere, and emission reduction strategies focused on technological changes such as alternative jet fuels. Recent modeling studies have aimed to quantify global impacts from full-flight cruise emissions; with over 75% of health impacts due to aviation-attributable air pollution estimated to be from full-flight emissions (Barrett et al., 2010; Yim et al., 2015; Dedoussi, 2021). However, global studies may not be capturing the local impacts due to limited chemistry and coarser model grid cell resolutions (Barrett et al., 2010; Levy et al., 2012b; Dedoussi et al., 2020). Local to regional scale modeling efforts to quantify health impacts from aviation-attributable pollution in the U.S. have utilized emission inventories from the early 2000s (Ratliff et al., 2009; Levy et al., 2012b; Lee et al., 2013; Koo et al., 2013; Ashok et al., 2013; Wolfe et al., 2014; Yim et al., 2015; Penn et al., 2017) which do not represent the impacts of the economic recession in 2007–2009 and the impacts of a changing U.S. atmosphere (Pye et al., 2020; Dedoussi, 2021). In fact, the recession in 2007–2009 had slowed aviation growth such that U.S. commercial air carriers' total number of domestic departures had not increased above 2007 levels until 2016 (FAA, 2014). And as far as the authors are aware, no study that has looked at health impacts from aviation-attributable air pollution have quantified the impacts from aviation-attributable NO₂, a potentially dangerous local-scale pollutant from mobile sources (Khreis et al., 2017; Atkinson and Butland, 2018; Achakulwisut et al., 2019; Mohegh et al., 2020). And while one study has looked at the air-quality-related health benefits from desulfurizing jet fuel (Barrett et al., 2012), and two others have looked at the non-volatile particulate matter (nvPM) emissions reductions from alternative jet fuels (Speth et al., 2015; Lobo et al., 2011), no study has looked to quantify the air-quality-related health benefits from fleet-wide implementation of sustainable alternative jet fuel.

This study aims to address some of the gaps of the prior studies by quantifying LTO aviation-attributable fine particulate matter (PM_{2.5}), ozone (O₃), and nitrogen dioxide (NO₂) concentrations and health impacts in the continental U.S. (CONUS) for the two most recent U.S. Environmental Protection Agencies' (EPA) National Emissions Inventories' (NEI) years of 2011 and 2016. Two additional 2016 LTO emission inventories have been generated and modeled to quantify the air quality impacts of a 5% and 50% blend of sustainable alternative jet fuel implemented across the U.S. This will be the first study to quantify the impacts from aviation-attributable NO₂ in the U.S. and the implementation of sustainable alternative jet fuel blends nationwide. We hope these results can add to the growing list of literature surrounding aviation-attributable air pollution and associated health impacts.

2. Methods

We utilize a modeling platform consisting of the Weather Research and Forecasting (WRF) model (Skamarock et al., 2008) to process meteorology inputs, the Sparse Matrix Operator Kernel Emissions (Baek and Seppanen, 2018) (SMOKE) model to process background emissions, and the Community Multi-Scale Air Quality Model (CMAQ) to estimate LTO aviation-attributable PM_{2.5}, O₃, and NO₂ across the continental U.S. (CONUS) for the years 2011 and 2016. CMAQv5.2 (US EPA Office of Research and Development, 2017) with the Carbon Bond 6 revision 3 mechanism (CB6r3) (Luecken et al., 2019) is used to quantify pollutant concentrations across our modeling domain. Our modeling domain covers the continental U.S. with 12 km × 12 km horizontal grid cell resolution. Background emissions (all sectors excluding commercial aviation) are from the U.S. EPA's 2011 (2011 NEI v2) (US Environmental Protection Agency, 2011) and 2016 (2016 NEI v1) (US Environmental Protection Agency, 2016) National Emissions Inventories. Aircraft emissions in this study were constructed from the FAA's Aviation Emission Design Tool (AEDT) (FAA-AEDT, 2016) for National Air

Space (NAS)-wide LTO aircraft activity for the years 2011 and 2015. LTO segments from AEDT datasets (Wilkerson et al., 2010) were processed into gridded emission rate files using AEDTProc (Baek et al., 2012) such that LTO emissions of nitrogen oxides (NO_x) (speciated to nitrogen oxide (NO), NO₂, and nitrous acid (HONO)), carbon monoxide (CO), sulfur dioxide (SO₂), total organic gases (TOG) (speciated to CB6r3 model species according to EPA's speciation profile (US Environmental Protection Agency and Federal Aviation Administration, 2009)), and primary PM (sulfate, organic aerosols, and elemental carbon (Wayson et al., 2009; Wilkerson et al., 2010)) are allocated to CMAQ-ready emission input files. Two additional LTO emissions inventories were generated to represent the expected emissions if sustainable alternative jet fuels (AJFs) were implemented across the U.S. as a 5% blend of existing jet fuel and a 50% blend of existing jet fuel. These inventories were generated by scaling hourly emission rates of the 2015 LTO inventory by fractional impact factors (values in S8) derived from field studies as defined in a recent synthesis study from the Airports Cooperative Research Program (ACRP) that quantified emissions reductions due to use of sustainable alternate jet fuels (Hamilton, 2019). AJFs are produced from non-petroleum sources which reduces the life-cycle greenhouse gases (Pavlenko and Searle, 2021; Martinez-Valencia et al., 2021) and when burned in aircraft engines, AJFs emit less air pollutants. In this study we are concerned with the latter as it pertains to air quality impacts from airport aircraft operations. AJF use has been shown to reduce emissions of sulfur oxides (SO_x) and nvPM significantly (~90% and ~50% SO_x reductions for 100% and 50% blends of AJFs, respectively (Beyersdorf et al., 2014; Corporan et al., 2010) and ~65% nvPM reductions for 50% blends (Beyersdorf et al., 2014; Chan et al., 2015; Lobo et al., 2011)) and to a lesser extent, CO (~10% CO reductions for 50% blends of AJF (Shouse et al., 2012)). There are currently three potential routes for AJFs: synthetic liquid fuels manufactured from current fossil fuels or biomass, bio-jet fuels made from agricultural oil crops, and hydrogen. Current use of AJFs requires a certain percentage to be blended with conventional petroleum-based jet fuel. Hence, the 5% and 50% impact factors derived in the ACRP report are from field campaigns that quantify the emissions and effects of different AJFs at specified blends up to 50%. In addition to the two NEI background emission inventories and LTO emission inventories, we made use of two meteorology datasets for 2011 and 2016. Details regarding model settings and datasets used can be found in tables S1–S2 and details regarding model evaluation against observations can be found in figures S1–S7 and table S5.

The health impact assessment tool BenMAPR is used to quantify the expected mortalities and morbidities due to LTO-attributable pollutants. Similar to the U.S. EPA's Benefits Mapping and Analysis Program (BenMAP) (Sacks et al., 2018), BenMAPR is a geospatial air pollution health impact assessment modeling platform written in the statistical computing language R (R Core Team, 2017) that links air pollution exposures to data on exposed populations and their background health. It then calculates the health impacts of these exposures using concentration response functions (CRFs) from the epidemiological literature. BenMAPR has been used in other health impact analyses (Arter et al., 2021) and relies on the same calculations and datasets used in BenMAP. For quantifying PM_{2.5}-attributable premature mortalities, we make use of a CRF from a recently published meta-analysis (Vodonos et al., 2018) that found a 1.29% (95% CI 1.09–1.5) increase in all-cause mortality per 10 μg/m³ increase in PM_{2.5}. For O₃-attributable premature mortalities, we use a CRF associating all-cause mortality to long-term O₃ exposure with a hazards ratio of 1.02 (95% CI 1.01–1.04) per 10 ppb increase in O₃ (Turner et al., 2016). For NO₂-attributable premature mortalities, we use a CRF from a meta-analysis that found a pooled effect on mortality to be 1.04 (95% CI 1.02–1.06) with an increase in 10 μg/m³ in NO₂ (Faustini et al., 2014). In addition to mortality estimates, we expand the analysis to incorporate multiple morbidity estimates such as: respiratory and cardiovascular hospitalizations (Levy et al., 2012a; Zanobetti et al., 2009), non-fatal heart attacks (Mustafić et al., 2012), and asthma

hospitalizations, emergency department visits, and exacerbations in those ages 5 to 17 (Orellano et al., 2017). Information regarding the morbidity CRFs and the underlying datasets used can be found in table S4. We chose CRFs in studies that best represent present day health outcomes and North American populations. If no suitable CRF could be found, we then relied on the CRFs that are currently used in BenMAP.

3. Results

3.1. Emissions

Table 1 shows the number of arrivals and departures at CONUS airports and annual emissions of CONUS LTO CO, NO_x, primary elemental carbon (PEC), and SO₂ for each simulation year. While the total number of arrivals and departures slightly decrease from 2011 to 2016, total fuel burned increases by ~10% (from ~4.66 Tg in 2011 to ~5.13 Tg in 2016) resulting in emissions increasing from 2011 to 2016. Emissions of CO, PEC, and SO₂ decrease under the 5% and 50% 2016 AJF scenarios with the amount decreased given by the impact factors in the ACRP report (table S8). It is important to note that other emission species, such as NO_x, HAPs, and volatile organic compounds (VOC) were estimated to be negligibly impacted by currently approved blend percentages (up to 50%) of sustainable alternative jet fuels. The ACRP report found no statistically significant impact for NO_x and HAPs and no statistically meaningful results for VOCs from their fitting of the experimental data. Hence, the authors cautioned applying impact factors for those species and we left those emission species unchanged from 2016 in the 5% and 50% blend scenarios.

We are able to quantify airport level emissions by looking at the emissions in the airport-containing grid cell. Fig. 1 shows the CO, NO_x, primary elemental carbon (PEC), and SO₂ emissions at the airport-containing grid cells for the top 28 airports in terms of passenger boardings (Air Carrier Activity Information System (ACAIS), 2015). The ORD (Chicago O'Hare International)-containing grid cell has the most NO_x emissions at 2,184 tons in 2011 and the JFK (John F. Kennedy International)-containing grid cell has the most NO_x emissions at 2,880 tons in 2016. The ATL (Hartsfield-Jackson Atlanta International)-containing grid cell has the most CO emissions at 1,421 tons in 2011 and the LAX (Los Angeles International)-containing grid cell has the most CO emissions at 1,519 tons in 2016. The ATL-containing grid cell has the most PEC emissions at 3.7 tons in 2011 and at 3.7 tons in 2016. The ORD-containing grid cell has the most SO₂ emissions at 172 tons in 2011 and at 192 tons in 2016. Emissions at the grid cell level are largely linear with respect to the number of flights at these airports. Figures S8-S11 show the number of arrivals and departures at 66 airports across the U.S. versus emissions of NO_x, CO, SO₂, and PEC in the airport-containing grid cell.

3.2. Air quality

Table 2 shows the CONUS-averaged population-weighted

Table 1

LTO emissions in the U.S. for each scenario, ^a percent change from 2011, ^b percent change from 2016.

	2011	2016	2016AJF5	2016AJF50
Arrivals and Departures	24,151,176	24,141,257 (−0.04%) ^a	24,141,257	24,141,257
CO (Tons/year)	44,637	49,382 (11%) ^a	48,888 (−1%) ^b	44,049 (−11%) ^b
NO _x (Tons/year)	64,056	73,926 (15%) ^a	73,926 (0%) ^b	73,926 (0%) ^b
PEC (Tons/year)	161	164 (2%) ^a	149 (−9%) ^b	58 (−65%) ^b
SO ₂ (Tons/year)	5,481	6,038 (10%) ^a	5,814 (−4%) ^b	3,773 (−38%) ^b

concentrations for LTO-attributable PM_{2.5}, O₃, and NO₂. PM_{2.5} and NO₂ concentrations represent the annual average while O₃ concentrations represent the annual average of the daily 8 h maximum. PM_{2.5} population-weighted concentrations increase by 7.7% from 2011 to 2016. PM_{2.5} population-weighted concentrations are expected to decrease in 2016 by 2.4% if a 5% blend of sustainable alternative jet fuel is implemented and by 18.4% if a 50% blend is implemented. O₃ population-weighted concentrations are negative in both 2011 and 2016, meaning that LTO emissions near the airport are removing O₃. This is due to localized NO_x/VOC ratios impacting whether O₃ formation is limited by NO_x or VOC concentrations and as it is often the case for densely populated urban areas; NO_x emissions can lead to a reduction in O₃ in VOC-limited areas. From Table 1 we can see that NO_x emissions from LTO across the U.S. are larger in 2016 than 2011 which corresponds to an 82% decrease in O₃. As both NO_x and VOC emissions are unaffected by the implementation of sustainable alternative jet fuels, O₃ concentrations are not impacted. NO₂ population-weighted concentrations increase by 77% from 2011 to 2016 and are also unaffected by the implementation of sustainable alternative jet fuels.

Figs. 2–4 show the LTO-attributable PM_{2.5}, O₃, and NO₂ concentrations, respectively across the continental U.S. aggregated to the county scale for both 2011 and 2016. Impacts are mostly seen in counties either containing an airport or surrounding an airport. Large negative O₃ concentrations (in blue in Fig. 3) outweigh positive O₃ concentrations that are occurring downwind of the airports in NO_x-limited areas. In 2016, two counties in Northwest Washington, Pierce and Island Counties, have negative LTO-attributable PM_{2.5} due to the negative LTO-attributable nitrate (NO₃[−]), ammonium (NH₄⁺), and organic matter (OM) components of PM_{2.5}. These components are reduced with LTO emissions due to LTO emissions of NO_x reacting with available O₃ and gas phase NO₃[−] near the airport to form nitrogen pentoxide (N₂O₅); resulting in a reduction of both gas phase NO₃[−] and oxidants available to form SOA (Woody and Arunachalam, 2013). In the summer, warmer temperatures can limit the particle-phase partitioning of ammonium nitrate (NH₄NO₃) resulting in negative LTO-attributable NH₄⁺. The higher temperatures limiting the partitioning of NH₄NO₃ into particle phase can also make the impacts on aerosol sulfate (SO₄^{2−}) formation more pronounced. Aerosol SO₄^{2−} formation can be reduced by the additional LTO NO_x increasing aqueous-phase acidity and subsequently reducing the dissolution of SO₂ in the aqueous-phase; as well as LTO NO_x limiting the oxidants available to oxidize SO₂ to sulfuric acid (H₂SO₄). And while these impacts are seen across the U.S. for other grid cells, these two Washington counties that are north and south of the Seattle-Tacoma airport are impacted enough by these effects to cause each county to have on average negative LTO-attributable PM_{2.5}.

Figs. 5–7 show the monthly LTO-attributable PM_{2.5}, O₃, and NO₂ concentrations, respectively across the continental U.S. for 2011 and 2016 as boxplots for all grid cells within the continental U.S. (top of each figure) and the monthly mean LTO-attributable concentrations (bottom of each figure). Figures S17 and S18 show the monthly mean LTO-attributable PM_{2.5} component concentrations for 2011 and 2016, respectively. In 2011, LTO-attributable NO₃[−] and NH₄⁺ are largest in the month of March due to the most LTO NO_x being emitted in March. The LTO NO_x can positively impact NO₃[−] and NH₄⁺ formation far enough downwind of airports in the month of March where the competition for available oxidants between LTO NO_x and SO₂ to form NO₃[−] and SO₄^{2−}, respectively, and the competition of the subsequent nitric acid (HNO₃) and H₂SO₄ for available NH₃ to determine the phase state of HNO₃ and subsequent aerosol phase NH₄NO₃ near the airport is no longer occurring. In cooler months, limited NH₃ near the airports will keep HNO₃ in the gas phase and H₂SO₄ neutralized. But downwind of the airports, LTO NO_x can positively impact NH₄NO₃ concentrations where this competition is no longer occurring. And while NO_x emissions are also large in the summer months, we see NO₃[−] being a much smaller portion of the LTO-attributable PM_{2.5} due to effects that occur near the airport and the

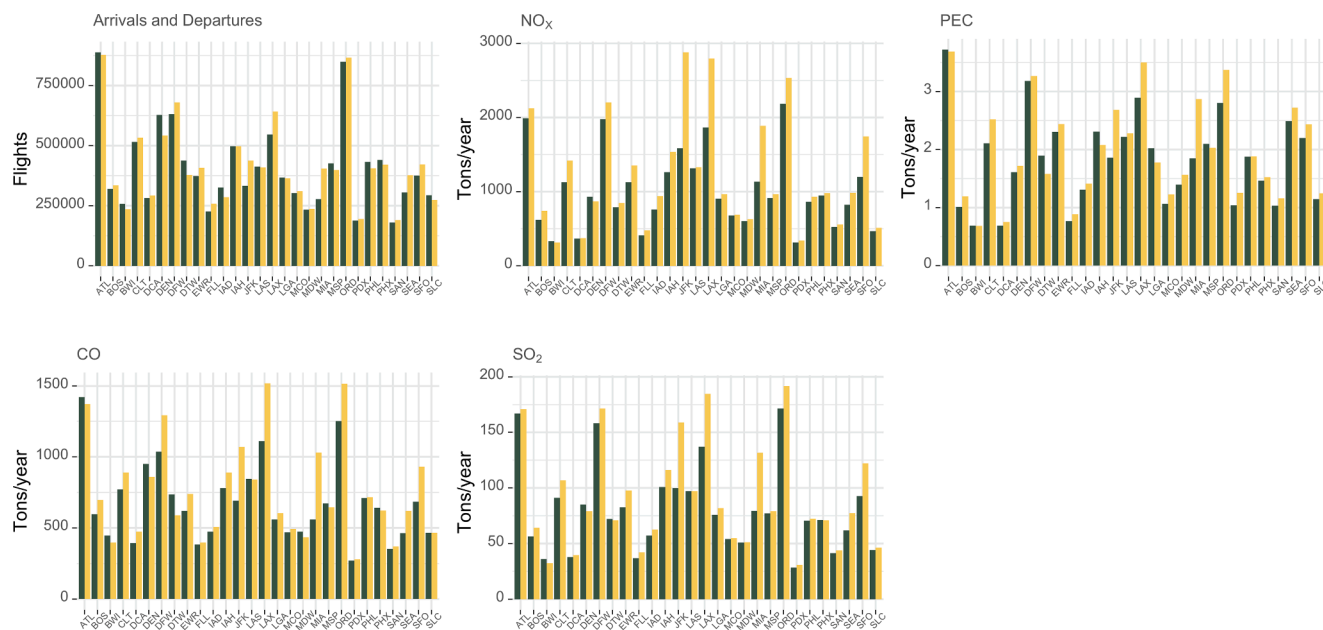


Fig. 1. Number of arrivals and departures and the LTO emissions in the airport-containing grid cells, table S6 describes the full names associated with each of the airport ID codes.

Table 2
Annual average population weighted concentrations due to LTO emissions in the U.S.

	2011	2016	2016AJF5	2016AJF50
PM _{2.5} (μg/m ³)	2.72E-03	2.93E-03	2.86E-03	2.39E-03
O ₃ (ppb)	-8.24E-03	-1.50E-02	-1.50E-02	-1.50E-02
NO ₂ (ppb)	3.92E-02	6.93E-02	6.94E-02	6.94E-02

warmer temperatures impacting the particle-phase partitioning of NH₄NO₃ and reduced formation of SO₄²⁻ as described in the previous paragraph. The distribution of grid cell concentrations shows the localized impacts of LTO emissions with outlier points' absolute values being order of magnitudes larger than the mean values and indicative of airport-containing grid cells (Figures S13-S15 show the monthly LTO-attributable PM_{2.5}, O₃, and NO₂ concentrations for airport-containing grid cells). These outliers drive the population-weighted exposures and as we will see, the health outcomes. Mean values of O₃ and NO₂

show opposite monthly trends with mean LTO-attributable O₃ concentrations peaking in the summer and diminishing in the winter and LTO-attributable NO₂ exhibiting the opposite. Both LTO-attributable O₃ and NO₂ impacts are driven by LTO NO_x emissions.

Similar to individual airport's emissions, we can look at the concentrations of PM_{2.5}, O₃, and NO₂ at the airport-containing grid cell level. It is important to note that the airport-grid cell concentrations are not indicative of the total impacts felt by LTO emissions near an airport, and this analysis is simply a comparison of concentrations in the grid cells containing the top 28 airports in terms of passenger boardings. Fig. 8 shows the pollutant concentrations at the airport-containing grid cell level for the same group of airports in Fig. 1. The SFO (San Francisco International)-containing grid cell has the largest increase in PM_{2.5} concentration of 5.4E-02 μg/m³ in 2011 and the LAX-containing grid cell has the largest increase in PM_{2.5} concentration of 4.9E-02 μg/m³ in 2016. O₃ titration impacts are seen at the airport-containing grid cell level where O₃ concentrations are decreased by the addition of LTO emissions. The SFO-containing grid cell has the largest decrease in O₃ concentration at -1.13 ppb in 2011 and -1.86 ppb in 2016. The LAX-

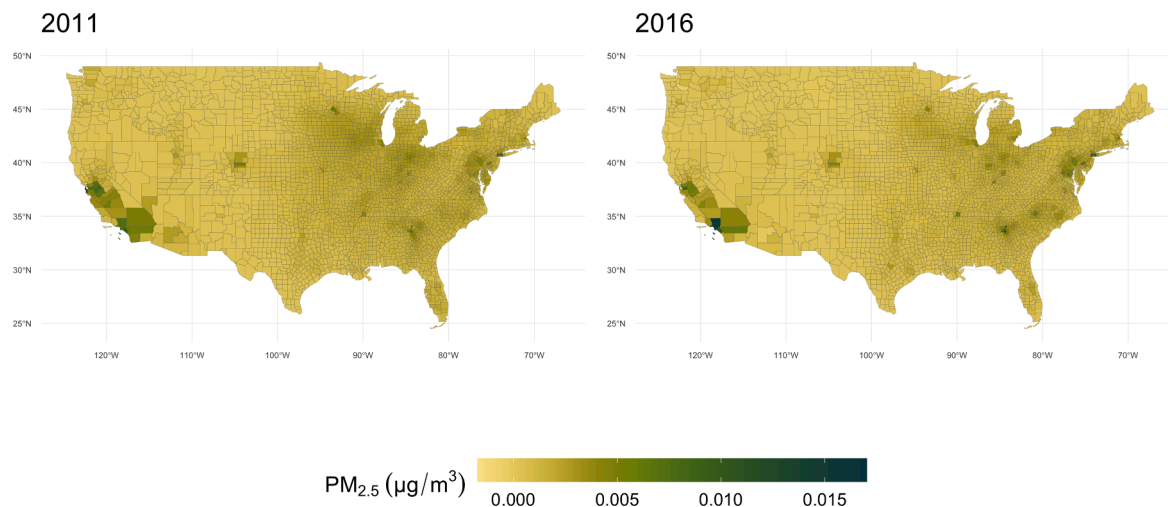


Fig. 2. LTO-attributable PM_{2.5} concentrations in 2011 (left) and 2016 (right).

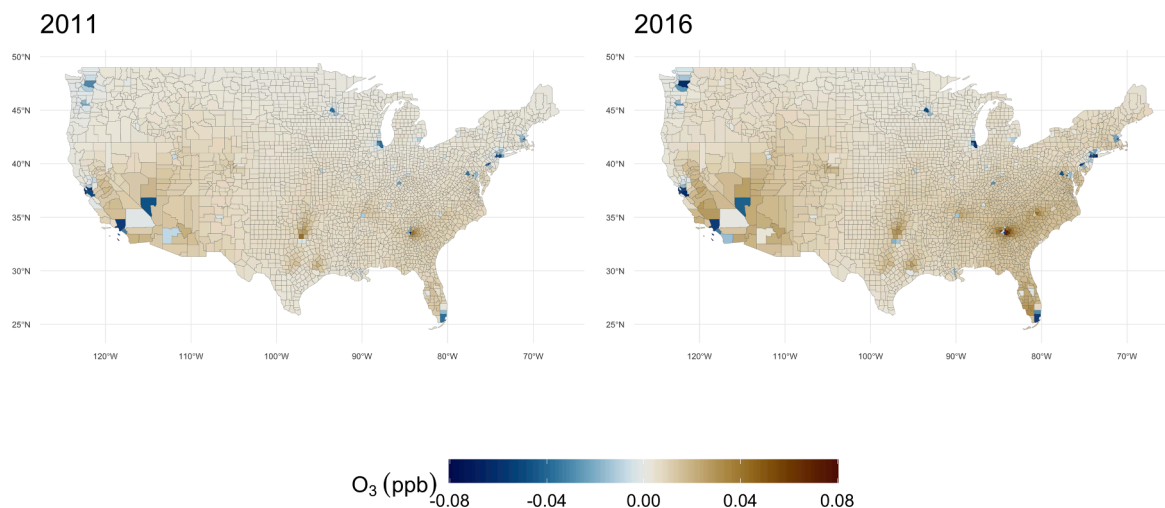


Fig. 3. LTO-attributable O₃ concentrations in 2011 (left) and 2016 (right).

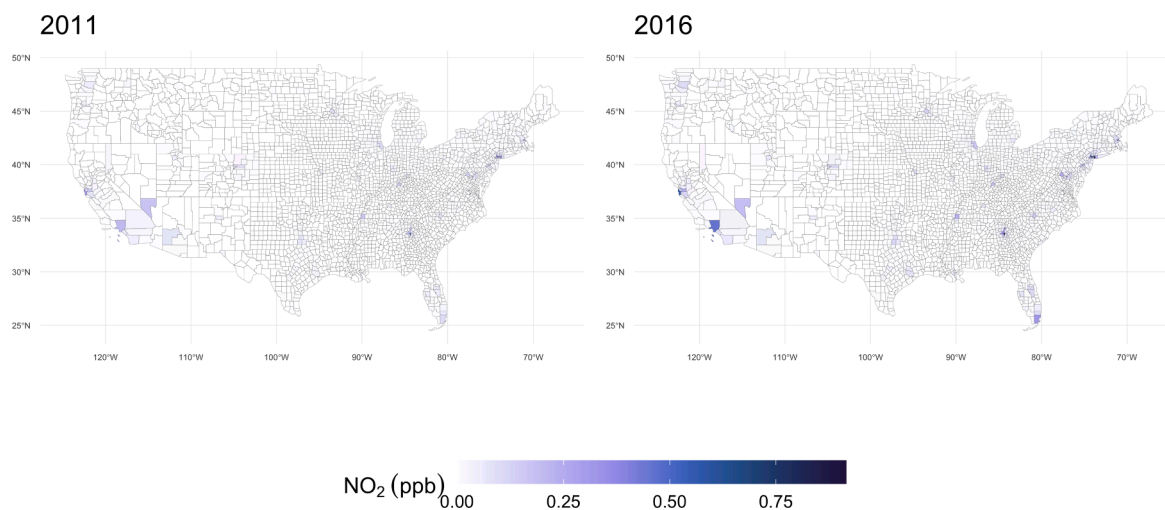


Fig. 4. LTO-attributable NO₂ concentrations in 2011 (left) and 2016 (right).

containing grid cell has the largest increase in NO₂ concentration at 1.76 ppb in 2011 and 4.44 ppb in 2016. Impacts in the SFO-containing grid cell may not be entirely due to LTO activity from SFO due to the close proximity to OAK (Oakland International) (see figure S12). Figure S16 breaks down the PM_{2.5} at the airport-containing grid cell level into concentrations of NH₄⁺, NO₃⁻, SO₄²⁻, elemental carbon (EC), and OM chemical constituents. LTO emissions are positively contributing to concentrations of NH₄⁺, SO₄²⁻, EC, and OM while concentrations of NO₃⁻ are negatively impacted. Decreases in NO₃⁻ and O₃ concentrations at the grid cell level can be explained by LTO emissions of NO being converted to NO₂. This process is possible due to the reaction of NO with O₃, which results in O₃ concentrations being reduced and NO₂ concentrations increasing. NO₂ is able to react with NO₃⁻ to form N₂O₅. Hence, the reduction of both NO₃⁻ and O₃ can also impact secondary organic aerosol (SOA) formation as both NO₃⁻ and O₃ are precursors needed for SOA formation (Woody and Arunachalam, 2013); which we are able to see by looking at the few cases in which OM is negatively impacted in 2016 in the ATL, IAH (George Bush Intercontinental/Houston), MCO (Orlando International), and SFO-containing grid cells. LTO emissions of NO_x at the airport-containing grid cell level are fairly linear with NO₂ concentrations (Figure S19) and O₃ concentrations (Figure S20) with opposite correlations.

3.3. Health impacts

Table 3 shows the total adverse health outcomes due to air quality degradation from LTO emissions in the U.S. in 2011 and 2016 and under two sustainable alternative jet fuel implementation scenarios in 2016. PM_{2.5}-attributable premature mortalities increase by 10% from 2011 to 2016. We estimate a net decrease in O₃-attributable premature mortalities across the U.S. in both 2011 and 2016 due to the large O₃ titration effects near the airports. Hence, the titration of O₃ in heavily populated regions around airports outweigh the formation of O₃ downwind. That is not to say that there are no adverse O₃ health outcomes in communities downwind of airports as we will see for health outcomes broken down by state in the next section. NO₂-attributable premature mortalities increase by 80% from 2011 to 2016 and are responsible for 91% of total premature mortalities in both 2011 and 2016. Implementing a 5% or 50% blend of sustainable alternative jet fuel in 2016 results in a 1% or 18% reduction, respectively in PM_{2.5}-attributable premature mortalities in 2016.

Figs. 9–11 show the number of PM_{2.5}, O₃, and NO₂-attributable mortalities, respectively in each state. California saw the most PM_{2.5}-attributable mortalities in 2011 and 2016 with 17.7 (14.9–20.6, 95% CI) and 23.7 (20.0–27.5) mortalities, respectively. Texas saw the most O₃-attributable mortalities in 2011 with 2.0 (1.0–4.1) mortalities while Georgia saw the most O₃-attributable mortalities in 2016 with 3.3

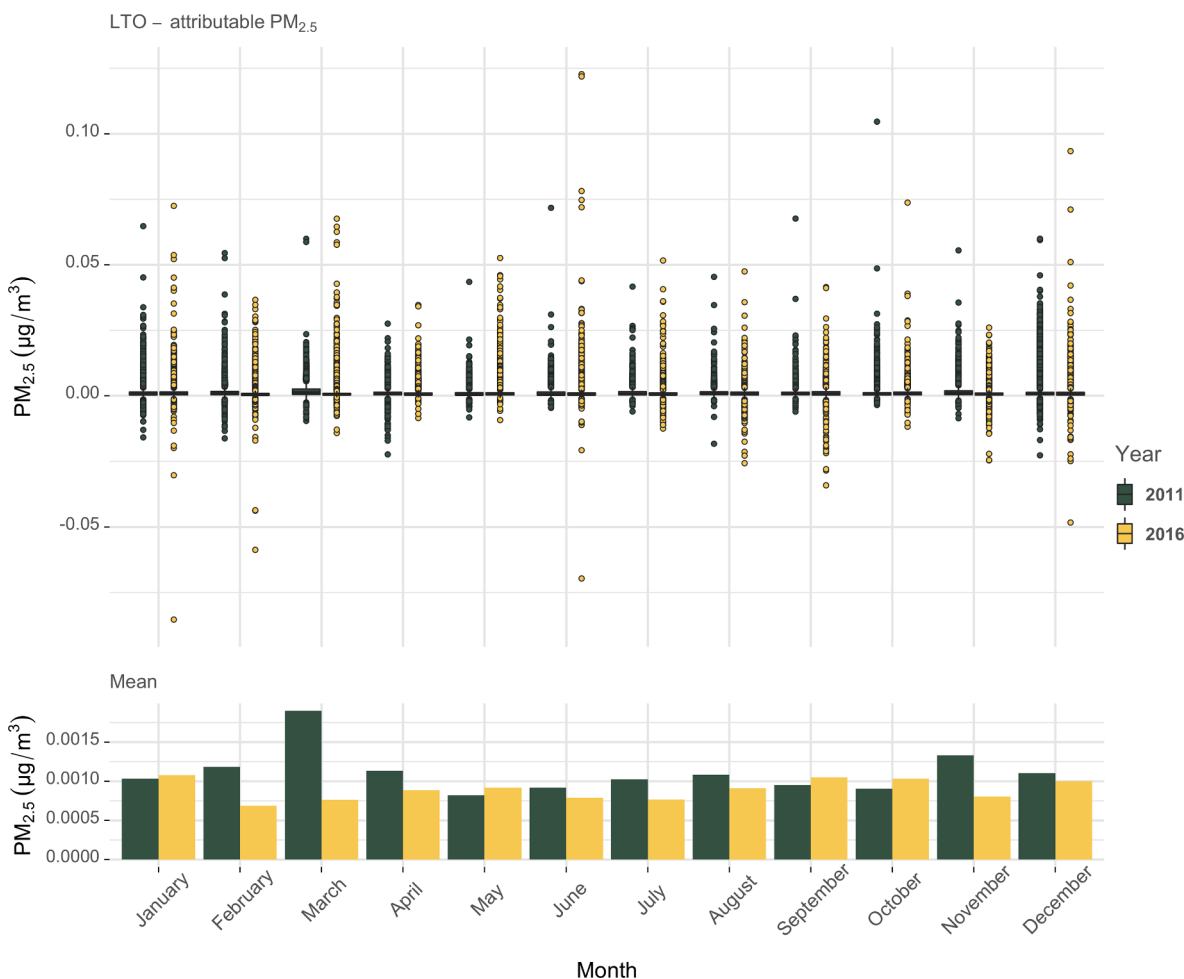


Fig. 5. Monthly LTO-attributable $PM_{2.5}$ concentrations for the grid cells that comprise the continental U.S.

(1.6–6.6) mortalities. California saw the most NO_2 -attributable mortalities in 2011 and 2016 with 155 (78–233) and 323 (161–484) mortalities, respectively.

We can monetize the value of avoided mortalities by applying a value of statistical life (VSL) approach as recommended by the EPA (US Environmental Protection Agency, 2010) and divide by the estimated fuel burned to approximate the economic damages per ton of fuel burned. We multiply the number of $PM_{2.5}$ and total (the sum of $PM_{2.5}$, O_3 , and NO_2) attributable premature mortalities in 2016 by a 2016 USD income-adjusted value of \$10.3 million and divide by the total fuel burn estimated through the total SO_2 emitted. We do this for our three scenarios in 2016 representing the economic damages per ton of traditional jet fuel burned, 5%-blended sustainable alternative jet fuel, and 50%-blended sustainable alternative jet fuel. We find LTO activity in 2016 to be responsible for 177 (90–292) USD $PM_{2.5}$ damages per ton of fuel burned. Implementing a 5% or 50% blend of sustainable alternative jet fuels results in 175 (88–292) and 144 (74–234) USD $PM_{2.5}$ damages per ton of fuel burned, respectively. Total damages per ton of fuel burned in 2016 are 2,211 (738–4,850) USD and drop to 2,209 (736–4,850) and 2,178 (723–4,791) USD for a 5% and 50% blend, respectively. The values in parentheses indicate an uncertainty in the willingness to pay for a reduction in mortality risk as given in table S3. Hence, a 50%-blended sustainable alternative jet fuel results in a 19% decrease in $PM_{2.5}$ damages per ton fuel burned from traditional jet fuel and a 2% decrease in total damages.

4. Discussion

We can compare health outcome estimates from prior studies that aimed to quantify the impacts from LTO-attributable $PM_{2.5}$ and O_3 . Levy et al. 2012 (Levy et al., 2012b) utilized a $36\text{ km} \times 36\text{ km}$ CMAQ platform for 99 airports in the U.S. that account for 94% of passenger enplanements in 2005 and quantified the LTO-attributable $PM_{2.5}$ premature mortalities to be 75. Ashok et al. 2013 (Ashok et al., 2013) and Ratliff et al. 2009 (Ratliff et al., 2009) also utilized a $36\text{ km} \times 36\text{ km}$ CMAQ platform for a 2005 LTO inventory and quantified 195 and 160 $PM_{2.5}$ premature mortalities, respectively. Each of these studies utilized different CMAQ versions and emissions inventories which owe to the variation in results. Barrett et al. 2010 (Barrett et al., 2010) utilized GEOS-Chem at a $1^\circ \times 1^\circ$ resolution for a 2006 LTO inventory and quantified the LTO-attributable $PM_{2.5}$ premature mortalities to be ~ 92 (20% of 458 premature mortalities due to full flight emissions). Koo et al. 2013 (Koo et al., 2013) utilized an adjoint of GEOS-Chem to quantify 20 $PM_{2.5}$ premature mortalities from LTO SO_2 emissions and 150 from LTO NO_x emissions in 2006. These studies both relied on global simulations in GEOS-Chem with limited chemistry and coarser model resolution. We estimate 80 and 88 LTO-attributable $PM_{2.5}$ premature mortalities in 2011 and 2016, respectively which are in line with the findings of these prior studies.

Fewer studies have aimed to quantify LTO-attributable O_3 mortalities. Yim et al. 2015 (Yim et al., 2015) utilized a $36\text{ km} \times 36\text{ km}$ CMAQ platform for a LTO inventory in 2006 and quantified LTO-attributable premature mortalities from both $PM_{2.5}$ and O_3 to be 650 in North America. Dedoussi et al. 2020 (Dedoussi et al., 2020) utilized GEOS-

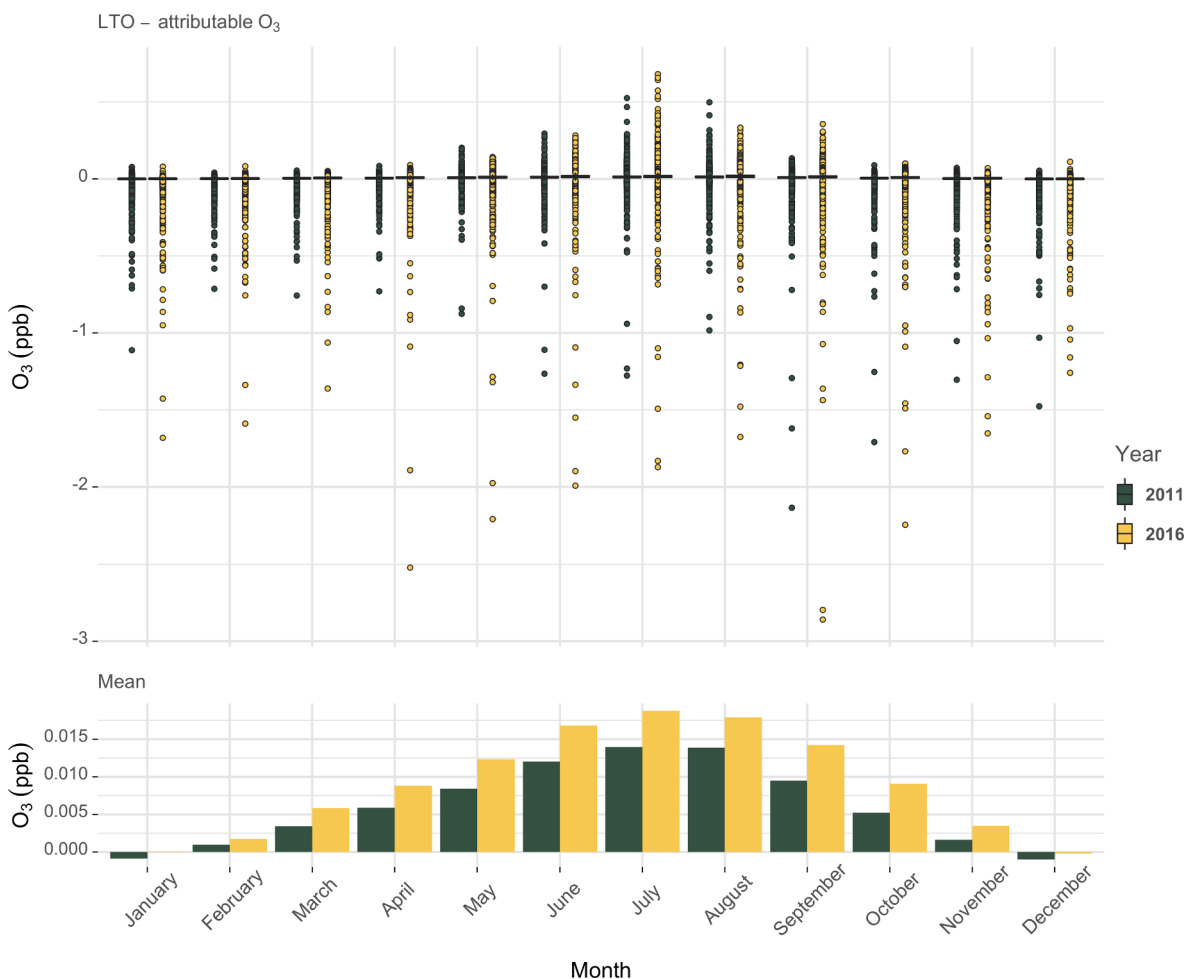


Fig. 6. Monthly LTO-attributable O₃ concentrations for the grid cells that comprise the continental U.S.

Chem at $0.5^\circ \times 0.67^\circ$ resolution for a LTO inventory in 2011 and quantified LTO-attributable premature mortalities from both PM_{2.5} and O₃ to be 209 in the U.S. Both of these studies have quantified total O₃ mortalities from LTO to be positive. However, these studies also highlight the impact of coarser model resolutions not being able to capture LTO NO_x emissions decreasing O₃ formation in VOC-limited areas. A recent global aviation sensitivity study by Qaudros et al. 2020 (Qaudros et al., 2020) stated that LTO emissions cause close to zero net increase in surface level ozone over the year due to airports often being located in areas with high NO_x surface level concentrations and the additional NO_x from aircraft is able to decrease O₃, especially in the winter. Two CMAQ-DDM studies aimed at quantifying sensitivities to individual airports (Penn et al., 2017; Arter and Arunachalam, 2021) also found O₃ depletion at airports from LTO NO_x emissions, with Penn et al. 2017 claiming that the 36 km × 36 km model resolution used may still be reducing the likelihood of NO_x-saturated/VOC-limited conditions (Penn et al., 2017). Missing from all of these studies is the quantification of LTO-attributable NO₂ adverse health outcomes which is closely tied to the localized O₃ impacts. In VOC-limited areas, NO₂ and O₃ will often be inversely correlated as a reduction in NO_x emissions can lead to an increase in O₃. Hence, additional NO_x from LTO activity in VOC-limited areas will impact both NO₂ and O₃ concentrations. A recent study in Hong Kong (Hossain et al., 2021) has aimed to quantify the added health risk (AR) associated with NO₂ and O₃ in a VOC-limited, heavily populated city and found the decrease of AR from ambient NO₂ from decreased NO_x emissions measured at roadside stations was considerably higher than the increase in AR associated with increased ambient O₃ in those urban areas. Another recent study (Moheg et al., 2020)

aimed to quantify the sensitivity of estimated NO₂-attributable health outcomes to grid resolution and found performing their analyses across U.S. cities at 10 km and 100 km resulted in 6% and 32% fewer adverse health outcomes, respectively than the analysis being performed at 100 m up to 1 km. Hence, by not taking into account LTO-attributable NO₂, especially at the resolutions needed to capture NO_x-saturated/VOC-limited conditions, we may be underestimating the air-quality related health impacts from LTO emissions.

Yim et al. 2015 (Yim et al., 2015) and Grobler et al. 2019 (Grobler et al., 2019) monetized the damages from LTO-attributable PM_{2.5} and O₃ premature mortalities. Yim et al. 2015 estimated 439 USD per ton of fuel burned in North America and Grobler et al. 2019 estimated 320 USD per ton of fuel burned in the U.S. for damages from PM_{2.5} and O₃. We estimate 177 USD per ton of fuel burned for damages from PM_{2.5} and 2,211 USD for damages from PM_{2.5}, O₃, and NO₂. We did not apply a cessation lag as was done in Yim et al. 2015 (Yim et al., 2015) and Grobler et al. 2019 (Grobler et al., 2019) when monetizing the damages. As we are comparing LTO-attributable PM_{2.5}, O₃, and NO₂ premature mortalities across LTO inventories, a PM cessation lag used in the results would not vary between the comparisons, and not match the CRF used here. In addition, the authors are unaware of research pertaining to cessation lags for use in NO₂-attributable premature mortalities. Future work can make use of proposed cessation lags for PM_{2.5}-attributable premature mortalities (Walton, 2010). While no other studies have quantified the impacts due to implementation of sustainable alternative jet fuel blends, Barrett et al. 2012 quantified the air quality related health impacts from implementation of ultra low sulfur fuel by reducing aircraft SO₂ emissions by 97.5% and found ~120–230 avoided PM_{2.5} mortalities in the U.

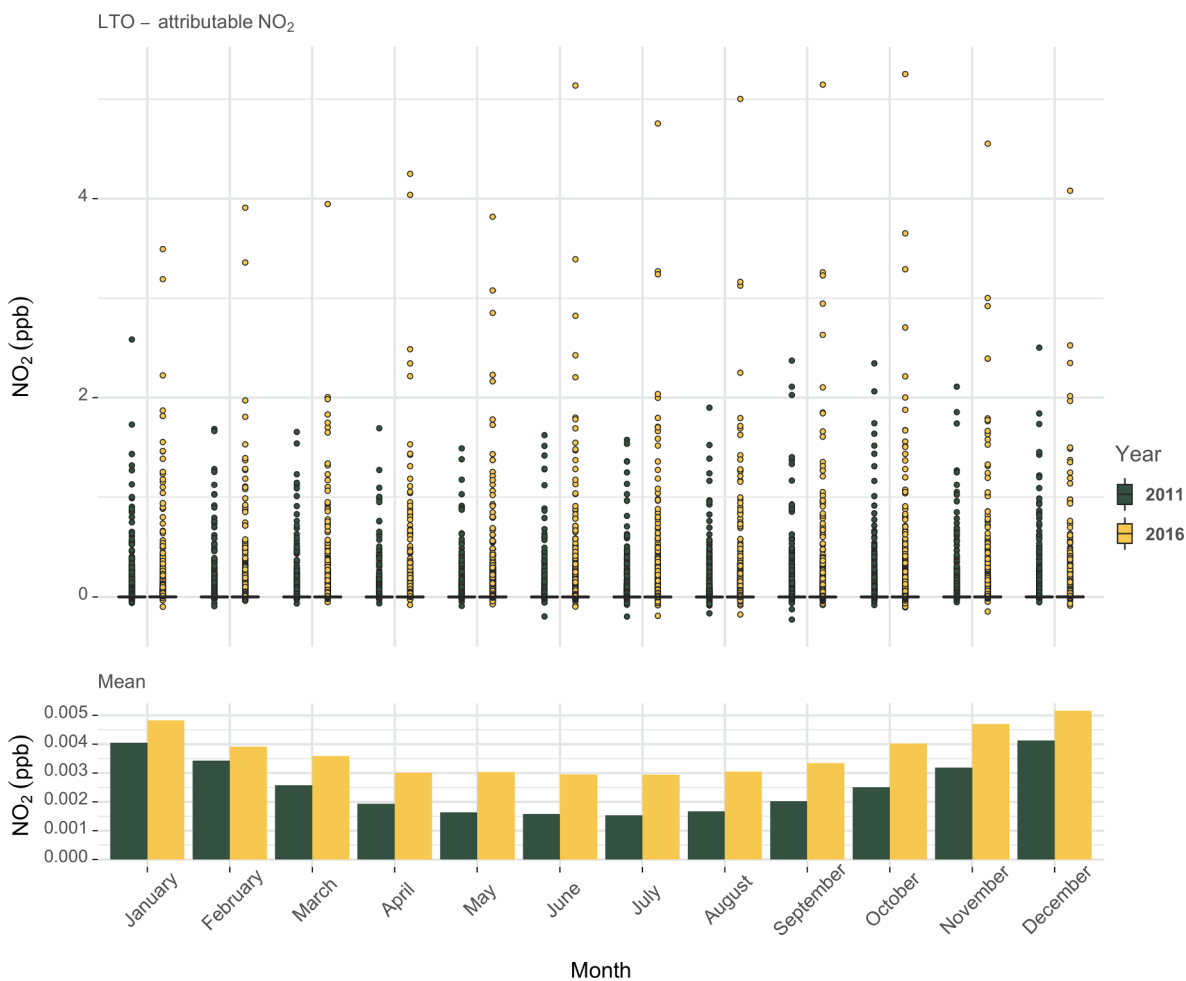


Fig. 7. Monthly LTO-attributable NO₂ concentrations for the grid cells that comprise the continental U.S.



Fig. 8. LTO-attributable PM_{2.5}, O₃, and NO₂ concentrations in the airport-containing grid cells.

Table 3

Total U.S. LTO-attributable air quality-related mortality and morbidity outcomes for each scenario (Values in parentheses reflect the 95% confidence intervals from the uncertainties in the CRFs, and estimates are rounded to two significant figures).

		2011	2016	2016AJF5	2016AJF50
Premature Mortalities	PM _{2.5}	80 (68–93)	88 (75–100)	87 (73–100)	72 (61–84)
	O ₃	–28 (–14 to –56)	–54 (–27 to –110)	–54 (–27 to –110)	–54 (–27 to –110)
	NO ₂	610 (310–920)	1,100 (570–1,700)	1,100 (570–1,700)	1,100 (570–1,700)
	Total	670 (360–960)	1,200 (610–1,700)	1,200 (610–1,700)	1,100 (600–1,700)
Cardiovascular Hospitalizations	PM _{2.5}	3.7 (2.6–4.9)	4.6 (3.1–6)	4.5 (3.1–5.9)	3.7 (2.6–4.9)
Respiratory Hospitalizations	PM _{2.5}	3.7 (1.9–5.4)	4.5 (2.3–6.6)	4.4 (2.3–6.5)	3.7 (1.9–5.4)
	O ₃	–12 (–4.5 to –20)	–25 (–8.9 to –40)	–25 (–8.9 to –40)	–25 (–8.9 to –41)
	Total	–8.7 (–2.6 to –15)	–20 (–6.6 to –34)	–20 (–6.6 to –34)	–21 (–7 to –35)
Non-fatal Heart Attacks	PM _{2.5}	3.7 (2.2–5.3)	4.1 (2.5–6)	4.1 (2.4–5.8)	3.4 (2–4.9)
	NO ₂	41 (22–59)	75 (41–110)	75 (41–110)	75 (41–110)
	Total	44 (24–65)	79 (43–110)	79 (43–110)	78 (43–110)
Asthma Hospitalizations Ages 5 to 17	PM _{2.5}	0.27 (0–0.54)	0.28 (0–0.57)	0.28 (0–0.56)	0.23 (0–0.46)
	NO ₂	14 (0.35–27)	23 (0.6–47)	23 (0.6–47)	23 (0.6–47)
	Total	14 (0.35–28)	24 (0.6–47)	24 (0.6–47)	24 (0.6–47)
Asthma Emergency Department Visits Ages 5 to 17	PM _{2.5}	2.6 (0–5.3)	2.7 (0–5.5)	2.7 (0–5.4)	2.2 (0–4.5)
	NO ₂	130 (3.2–250)	220 (5.6–430)	220 (5.6–430)	220 (5.6–430)
	Total	130 (3.2–260)	220 (5.6–440)	220 (5.6–440)	220 (5.6–440)
Asthma Exacerbations Ages 5 to 17	PM _{2.5}	2,100 (0–4,300)	2,300 (0–4,600)	2,200 (0–4,500)	1,900 (0–3,800)
	NO ₂	100,000 (2,500–200,000)	170,000 (4,400–340,000)	170,000 (4,400–340,000)	170,000 (4,400–340,000)
	Total	100,000 (2,500–200,000)	170,000 (4,400–340,000)	170,000 (4,400–340,000)	170,000 (4,400–340,000)

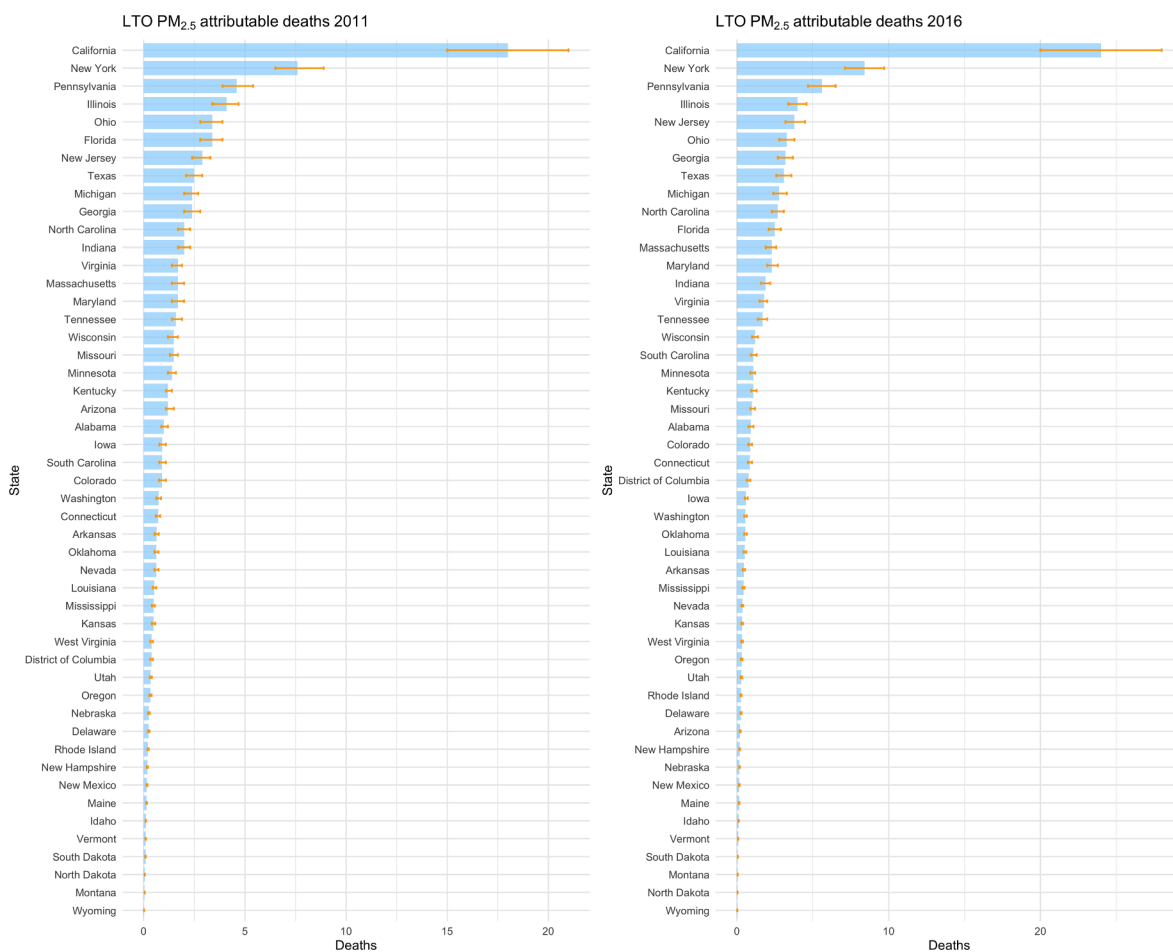


Fig. 9. PM_{2.5}-attributable mortalities in each state in 2011 (left) and 2016 (right), the orange bars reflect the 95% confidence intervals from the uncertainties in the CRFs.

S. depending on model choice and CRF. Barrett et al. 2012 (Barrett et al., 2012) estimated PM_{2.5} exposure to be reduced by 6% globally by using ultra low sulfur fuel. We estimate a reduction in PM_{2.5} population-weighted concentrations of 2.4% and 1 fewer PM_{2.5} mortalities in the U.S. if a 5% blend of sustainable alternative jet fuel is implemented and

an 18.4% reduction in PM_{2.5} population-weighted concentrations and 16 fewer PM_{2.5} mortalities if a 50% blend is implemented.

To account for uncertainty in the health outcome estimates, we have included results from the 95% confidence intervals in the CRFs. The confidence intervals for each mortality and morbidity outcome reflect

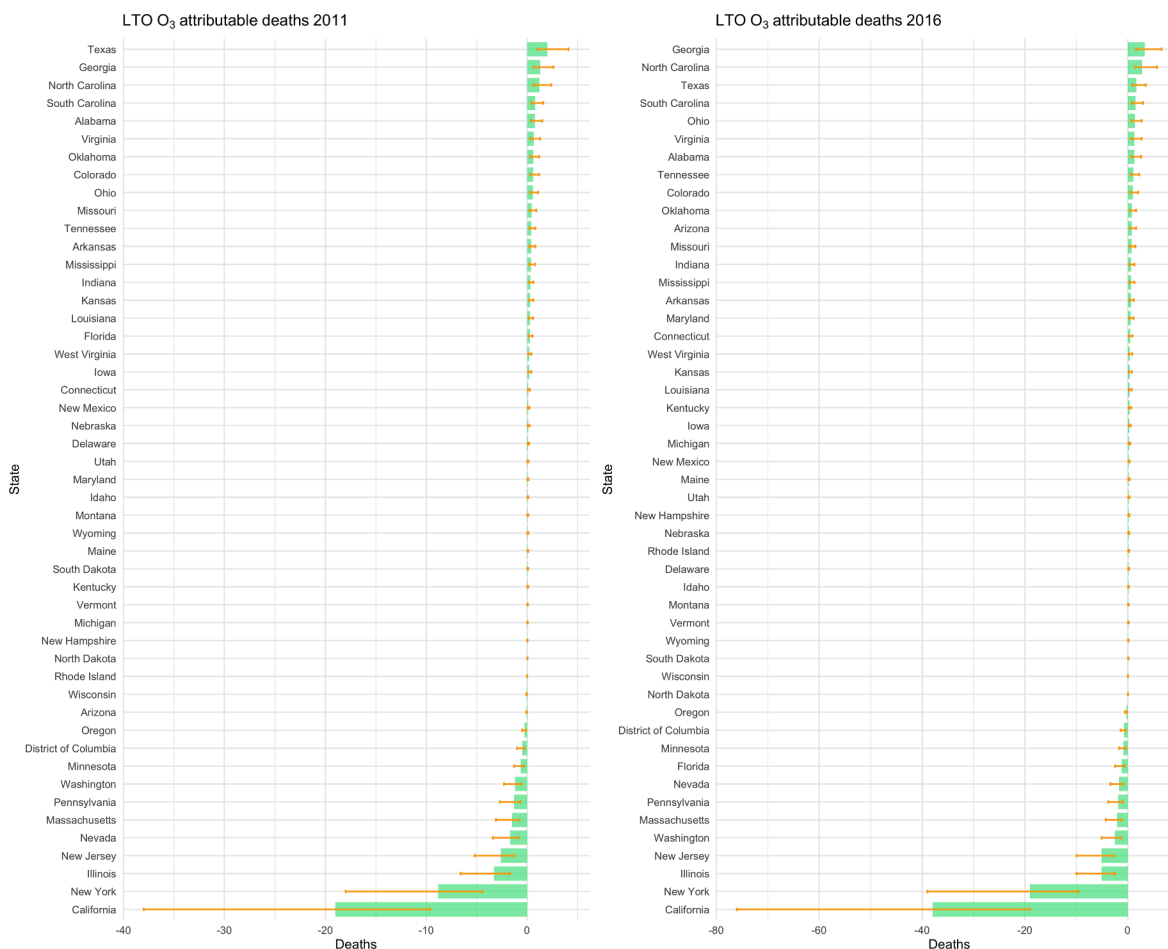


Fig. 10. O₃-attributable mortalities in each state in 2011 (left) and 2016 (right), the orange bars reflect the 95% confidence intervals from the uncertainties in the CRFs.

the variability in the results from the epidemiological studies used to construct the CRFs. We accounted for uncertainty in the damages per ton of fuel burn by including lower and upper bounds for the 2016 USD (\$) income-adjusted value of \$10.3 million as given in table S3. Additional uncertainties exist in each stage of the emission-to-health outcome modeling chain. Uncertainties in the emission inventories and the air quality model outputs are not accounted for in this study as the focus is a comparison of emission inventories. Additional work is needed to assess the uncertainty surrounding the emission inventories and air quality model outputs. Follow up studies should also look to distinguish the impacts from various types of sustainable alternative jet fuels. The impact factors used in this work are derived from multiple types and blends of sustainable alternative jet fuels. Next steps in this work should aim to quantify the impacts by fuel, engine, and aircraft type to get a better estimate of local air quality impacts from an airport's fleet makeup and choice of sustainable alternative jet fuel.

In this study we compare impacts of the LTO emission sector between two years, 2011 and 2016, that have varying background and LTO emission inventories, meteorological datasets, and population estimates. We did not perform sensitivity analyses that determine the impacts due to each of those varying datasets. Hence, when we make comparisons between the two years we are assuming that we are capturing the most accurate representations of each modeling year. Additional work can quantify the impacts due to each variable similar to the work done in Levy et al. 2012 (Levy et al., 2012b) which quantified LTO impacts by varying background emissions, LTO emissions, and populations in 2005 and projected 2025 values. However, that study also did not quantify impacts due to changing meteorology which can be

significant especially for timeframes indicative of changing climates (Fann et al., 2021).

While we are the first to report large adverse health outcomes from LTO-attributable NO₂, other studies have started to include NO₂ in traffic-related pollution health burden studies (Atkinson and Butland, 2018; Mohegh et al., 2020; Hossain et al., 2021; Southerland et al., 2021). As NO₂ is localized to the emission source, coarsely resolved modeling efforts may not capture the spatial variability of NO₂ (Anenberg et al., 2017) and its health impacts (Mohegh et al., 2020). While 12 km × 12 km resolution is not as fine as 4 km × 4 km or 1 km × 1 km, the results from this study are more finely resolved than results from prior aviation-related health impact modeling studies. That being said, 12 km × 12 km resolution may still be missing the fine resolution needed to capture NO₂ variability leading to an underestimation of impacts. And while other studies have quantified the impact of LTO emissions on PM_{2.5} concentrations at varying grid cell resolutions (Arunachalam et al., 2011; Woody et al., 2016; Rissman et al., 2013) and found PM_{2.5} population exposure to be mostly unaffected by grid cell resolution, NO₂ exposure is dependent on grid cell resolution. In addition, 12 km × 12 km resolution may still be too coarse to accurately describe the NO_x-saturated/VOC-limited conditions that govern O₃ response near airports. Future modeling studies should continue to move to finer spatial resolutions when estimating NO₂ and O₃ concentrations. In addition to finer spatial resolutions, incorporating plume level processes may improve LTO-attributable NO₂ and O₃ estimates. Two studies (Cameron et al., 2013; Fritz et al., 2020) have looked at the impacts of including plume level treatment for aircraft sources as compared to instant mixing within a photochemical model's grid cell. Both of these studies found an

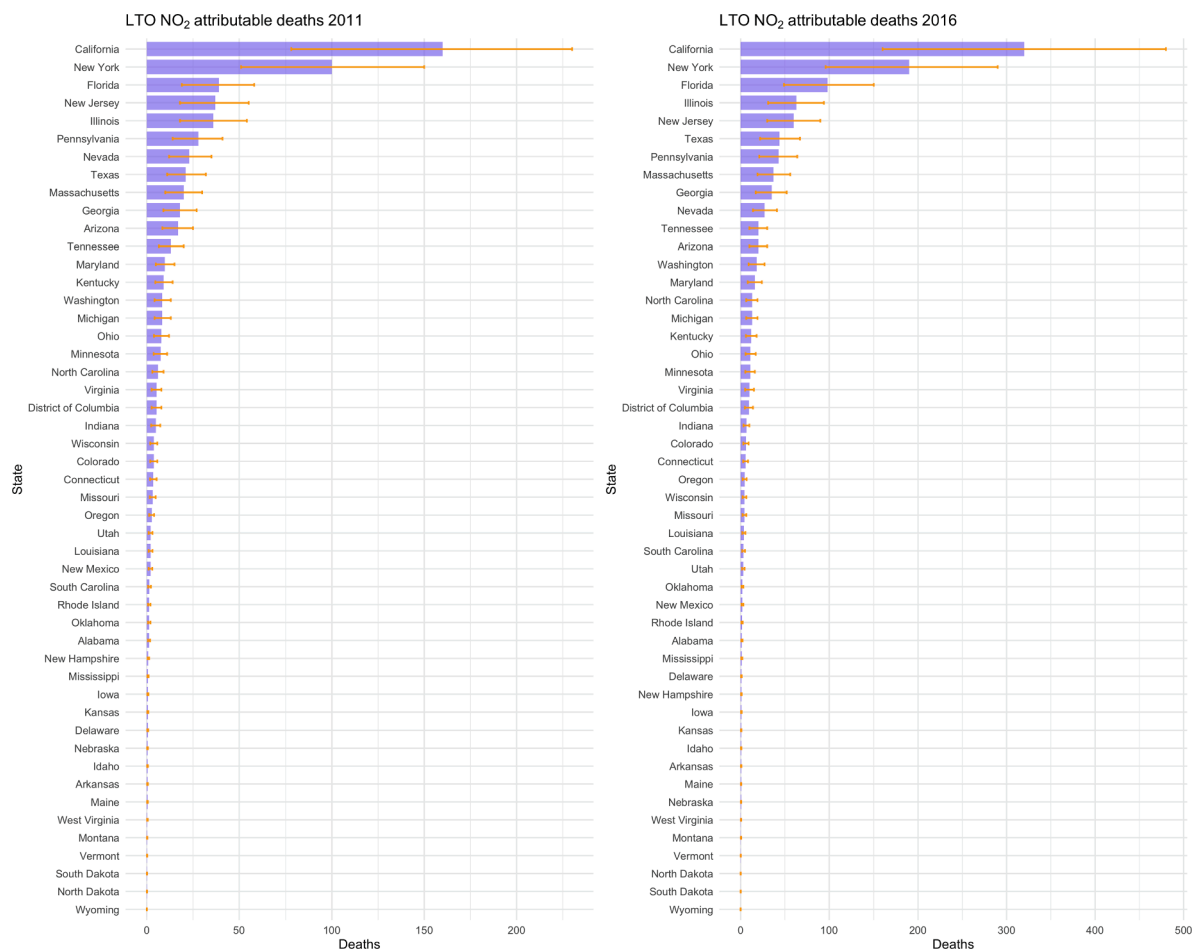


Fig. 11. NO_2 -attributable mortalities in each state in 2011 (left) and 2016 (right), the orange bars reflect the 95% confidence intervals from the uncertainties in the CRFs.

overestimation of O_3 production without plume level treatment and an underestimation of the remaining NO_x . In addition these studies also quantified the conversion of NO_x into reservoir species with and without plume treatment, with one study (Cameron et al., 2013) finding the plume treatment results in more NO_x remaining in the plume than compared to the grid treatment, and the other study (Fritz et al., 2020) finding that plume treatment results in NO_x to HNO_3 conversion being reduced by 16% and NO_x to N_2O_5 by 37%. Hence, we can surmise that by not including plume level treatment, we are underestimating NO_2 impacts.

From this study we have two rather impactful findings: 1. LTO-attributable NO_2 is responsible for 91% of total LTO-attributable premature mortalities in the U.S. and 2. Implementing a 5% or 50% blend of sustainable alternative jet fuels leaves LTO-attributable NO_2 concentrations untouched as sustainable alternative jet fuels do not reduce NO_x emissions. Table 10 in the ACRP report (Hamilton, 2019) shows the impact factors for NO_x for each experimental study considered. Of the 34 distinct measurements made with varying engine type, conventional jet fuel, sustainable alternative jet fuel, and blend percentage cited in Table 10 of the ACRP report; only 5 measurements showed non-zero impact factors, and after the ACRP authors attempted to construct a fit of the data, the uncertainty was greater than the absolute value of the impact factor. Hence, the authors of the ACRP report concluded that there is no statistically significant impact associated with sustainable alternative jet fuels for NO_x . As LTO NO_x emissions are almost entirely responsible for LTO-attributable NO_2 , we find that sustainable alternative jet fuel implementation will do little to mitigate the NO_2 air quality-related health impacts from LTO emissions in the U.S. NO_x emissions are

primarily due to combustion in the air flow, and compared to nitrogen content in the fuel; atmospheric nitrogen is the dominant source of NO_x (National Academies of Sciences, Engineering, and Medicine, 2018). Hence, to reduce LTO NO_x emissions and the associated adverse air quality-related health outcomes, technologies aimed at combustion modifications such as Lean Premixed Prevaporized (LPP) combustion (Sattelmayer et al., 1998; Anacleto et al., 2003), Lean Direct Injection (LDI) combustion (Liu et al., 2017), and water/steam injection (Block Novelo et al., 2019; Golzari et al., 2021) need to be explored.

As recent studies surrounding aviation-related environmental impacts have shifted toward the impacts from full flight emissions at global scales, the results from this study hope to highlight the importance of a continued effort in reducing the air quality-related impacts from LTO emissions on vulnerable populations near airports. By including LTO-attributable NO_2 health burden estimates, we introduce an additional factor to consider when assessing the public health impact of the aviation sector. This study also shows the benefits of implementing sustainable alternative jet fuel blends at a national scale which can help to inform aviation sustainability efforts while also protecting public health.

CRedit authorship contribution statement

Calvin A. Arter: Methodology, Investigation, Data curation, Software, Formal analysis, Writing – original draft. **Jonathan J. Buonocore:** Software, Data curation, Writing – review & editing. **Chowdhury Moniruzzaman:** Software, Investigation, Data curation. **Dongmei Yang:** Software, Investigation, Data curation. **Jiaoyan Huang:** Conceptualization, Methodology. **Saravanan Arunachalam:**

Conceptualization, Methodology, Validation, Supervision, Project administration, Resources, Funding acquisition, Writing – review & editing.

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 material

Supplementary data associated with this article can be found, in the online version, at <https://doi.org/10.1016/j.envint.2021.106958>.

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