

# Impact of Meteorology and Aerosol Sources on PM<sub>2.5</sub> and Oxidative Potential Variability and Levels in China

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**Abstract:** China has long-term high PM<sub>2.5</sub> levels, and its Oxidative Potential (OP) is worth studying as it may unravel the impacts of aerosol pollution on public health better than PM<sub>2.5</sub> alone. OP refers to the ability of PM<sub>2.5</sub> to induce oxidative stress (OS). OP and PM<sub>2.5</sub> are influenced by meteorological factors, anthropogenic emissions sources and atmospheric aging. Although their impact on PM<sub>2.5</sub> have been

20 studied, OP measurements only recently became available and on a limited scale, as they require considerable technical expertise and resources. For this, the joint relationship between PM<sub>2.5</sub> and OP for a wide range of meteorological conditions and emissions profiles remain elusive. Towards this, we estimated PM<sub>2.5</sub> and OP over China using the Danish Eulerian Hemispheric Model (DEHM) system with meteorological input from WRF weather forecast model. It was found that higher values of PM<sub>2.5</sub> and

25 OP were primarily concentrated in urban agglomerations in the central and eastern regions of China, while lower values were found in the western and northeastern regions. Furthermore, the probability density function revealed that about 40% of areas in China had an annual average PM<sub>2.5</sub> concentrations exceeding the Chinese concentrations limit. For OP, 36% of the regions have OP below 1  $nmol\ min^{-1}\ m^{-3}$ , 41% have OP between 1 and 2  $nmol\ min^{-1}\ m^{-3}$ , and 23% have OP above

30 2  $nmol\ min^{-1}\ m^{-3}$ , which are in line with previous measurement studies. Analysis of the simulations indicate that meteorological conditions contributed 46% and 65% to PM<sub>2.5</sub> concentrations and OP variability, respectively, while anthropogenic emissions contributed 54% and 35% to PM<sub>2.5</sub> concentrations and OP variability, respectively. The emission sensitivity analysis also highlighted PM<sub>2.5</sub> and OP levels are mostly determined by secondary aerosol formation and biomass burning.

35 **Keywords:** PM<sub>2.5</sub>; Oxidative potential; Meteorology; Anthropogenic emission sources; Probability density function

## 1. Introduction

Fine particulate matter, with an aerodynamic diameter of less than 2.5  $\mu\text{m}$  ( $\text{PM}_{2.5}$ ), is the primary atmospheric pollutant in China (Chen et al., 2021; Chen et al., 2021; Liu et al., 2023) with respect to human health.  $\text{PM}_{2.5}$  exposure in China for 2017 resulted in an estimated 1.8 (95% CI: 1.6, 2.0) million premature deaths (Liu et al., 2021). Many recent studies have suggested that the oxidative potential (OP) of  $\text{PM}_{2.5}$  may better explain the negative impact of  $\text{PM}_{2.5}$  exposure on human health than the well-established metric of mass concentrations (Yu et al., 2019; Gao et al., 2020). OP refers to the ability of  $\text{PM}_{2.5}$  to induce oxidative stress (OS) (Yang et al., 2021). Liu et al. (2020) summarized OP measurements conducted in nine regions of China around 2014. The results showed that the average OP content in northern Beijing was highest during the winter of 2016 ( $\sim 14.0 \text{ nmol min}^{-1} \text{ m}^{-3}$ ), while the average OP level in Shanghai during the spring of 2016 was lowest ( $\sim 0.15 \text{ nmol min}^{-1} \text{ m}^{-3}$ ). However, there is currently no exact threshold division of OP values. Exposure to high levels of OP (from compounds such as quinones and soluble transitional metals) induces an excess production of reactive oxygen species (ROS) in cells and lead to OS effects and ultimately trigger inflammation and disease. Therefore, reducing  $\text{PM}_{2.5}$  pollution and its associated OP (the volume-normalized dithiothreitol activity) are critical to addressing China's environmental and environmental health issues.

Anthropogenic emissions, as the main source of  $\text{PM}_{2.5}$  pollution and environmental health risks, have been studied extensively (Chen et al., 2019; Liu et al., 2022). Zhu et al. (2018) and Pui et al. (2014) summarized the studies on the PM sources in China and reported that secondary inorganic aerosols (SIA), industry, residential combustion, biomass burning, industry, and transportation are the main source categories in China in the historical and future business-as-usual scenarios. Due to the significant influence of various sectors on  $\text{PM}_{2.5}$  emissions, and research (Liu et al., 2018; Liu et al., 2020) indicating a close association between  $\text{PM}_{2.5}$  and OP, the connection between OP, serving as a toxicity indicator for  $\text{PM}_{2.5}$ , and its sources (Liu et al., 2020) is becoming increasingly crucial and the topic of numerous studies. For instance, Yu et al. (2019) used the dithiothreitol (DTT) assay to measure the  $\text{PM}_{2.5}$  samples in Beijing throughout the year and identified vehicle emissions as the main contributing source based on the source analysis of OP. However, studies conducted in three coastal cities of the Bohai Sea region (Liu et al., 2018) and in Nanjing (Zhang et al., 2023) using the same DTT assay indicated that coal combustion was the most active source of OP. Together, these studies suggest that obtaining the

spatial distribution characteristics of PM<sub>2.5</sub> and OP and their links to emission sources, is of paramount importance for implementing region-specific control measures.

Apart from anthropogenic emissions, meteorological conditions (i.e., temperature, humidity, wind speed, precipitation) also play a crucial role in the formation, accumulation, transformation, and dispersion of PM<sub>2.5</sub> (Liu et al., 2022; Liu et al., 2022). Utilizing a multiple linear regression model, Gong et al. (2022) conducted an analysis of the trends of meteorological elements and PM<sub>2.5</sub> levels across various regions in China from 2013 to 2020. Furthermore, they separated and quantified the impacts of meteorological factors and emissions on these trends. The findings indicate that meteorology alone can account for approximately 20~33% of the variability in PM<sub>2.5</sub> levels. Xing et al. (2023) conducted a study in the Shenzhen region using DTT, ascorbic acid (AA), and glutathione (GSH) OP assays. They analyzed meteorological conditions and PM<sub>2.5</sub> chemical composition to understand how the prevalence of monsoons in winter (northern and northeastern winds) and summer (southern and southeastern winds) affected the sources and contributed to the seasonal variation in PM<sub>2.5</sub> composition and OP (mass-normalized). Similarly, Molina et al. (2023) and Wang et al. (2019) revealed that meteorological conditions indirectly influence OP (volume-normalized and mass-normalized) through their impact on the chemical properties of the components. Ainur et al. (2023) employing a DTT assay, investigated outdoor health risks associated with atmospheric particulate matter in Xi'an, find a positive correlation between winter OP (volume-normalized) and relative humidity. Although several studies have identified linkages between meteorological conditions and PM<sub>2.5</sub>/OP, quantitative assessment of meteorological conditions to both PM<sub>2.5</sub> and OP variability is lacking.

As of the present, research on the influence of both meteorological conditions and anthropogenic emissions on OP primarily relies on measurement methods (Yu et al., 2019; Gao et al., 2020; Campbell et al., 2021), such as DTT, AA, and GSH, which are difficult and costly to test and hard to provide the spatial distribution of OP comprehensively. Although mechanistic models of OP do exist (Shahpoury et al., 2024), their links to experimental metrics of OP are qualitative. For this, we propose a hybrid approach combining existing observations of OP with a chemistry transport model (CTM). So, using OP from assays and their observed links to sources and chemical constituents can then be parameterized and implemented in CTM for a comprehensive assessment of OP exposure over large areas and time periods. This study quantifies the contribution of meteorological and anthropogenic emission factors (i.e., coal combustion, biomass burning, secondary aerosol formation that originate from a series of atmospheric

reactions, industry, and transportation source) to OP and PM<sub>2.5</sub> levels throughout China with the Danish Eulerian Hemispheric Model (DEHM) model. The study hence provides a method for calculating OP across China and using OP as an indicator to assess the impacts of anthropogenic emission sources on human health in China.

## 2. Materials and methods

### 2.1 Methodological flow

The research strategy of this study consists of three main parts: model setup, spatial distribution characteristic analysis, and quantification of meteorological conditions and anthropogenic emissions' contributions (Figure 1). In the first part, DEHM was employed to obtain hourly pollutant concentrations, followed by model evaluation, where the numerical weather prediction model WRF v4.1 (Skamarock et al., 2008) driven by either ERA5 or global climate data from CESM was used as meteorological input to DEHM and with exactly same spatial setup for China as in DEHM. Sensitivity experiments were designed for meteorological conditions, emission inventories, and anthropogenic emission sources. From these simulations, the spatial-scale estimation of OP was estimated by incorporating simulated values of primary and secondary PM<sub>2.5</sub> concentrations from various anthropogenic sources into Equation (1) (see Sect.2.2 for detail).

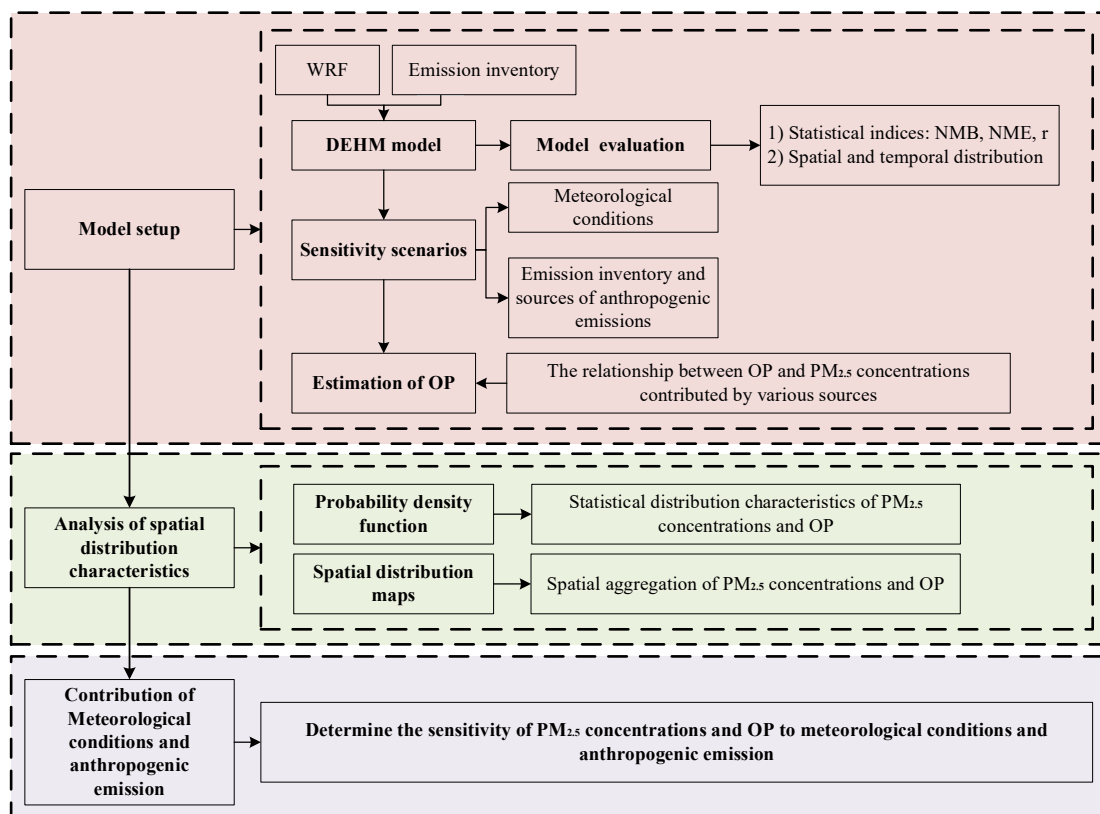


Figure 1. Schematic diagram of the study strategy; NME, NMB,  $r$ , and OP are normalized mean error, normalized mean bias, correlation coefficient, and oxidative potential, respectively.

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In the second part, the spatial distribution characteristics of  $PM_{2.5}$  and OP were determined using probability density functions (PDF) and spatial distribution maps. In the third part, quantitative analysis was conducted based on the simulation results from the sensitivity experiments to determine the extent of influence of meteorology and emissions on  $PM_{2.5}$  and OP, as well as the primary sources of  $PM_{2.5}$  and OP.

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## 2.2 Estimation of OP

Most of current data on OP of  $PM_{2.5}$  in China are obtained by means of measurement, and the research objects are basically limited to specific cities, which to some extent hinders the conduct of research on OP in a large-scale region. Considering that Liu et al. (2018) collected samples across four seasons from multiple representative locations in China, their developed OP prediction model (Equation (1)) can support us in estimating OP (with a unit of  $nmol\ min^{-1}\ m^{-3}$ ) in China, thereby exploring the spatial distribution characteristics of OP and the contributions of different anthropogenic sources to OP. In the present study, we have used this relationship, in combination with the sensitivity simulations (Sect. 2.4), to calculate the OP.

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$$OP = 0.088 \times re + 0.076 \times bi + 0.041 \times se + 0.034 \times in + 0.017 \times tr \quad (1)$$

130 where, *re*, *bi*, *in*, and *tr* represent the primary PM<sub>2.5</sub> concentrations (with a unit of  $\mu g m^{-3}$ ) for coal combustion, biomass burning, industry source, and transportation source, respectively. *se* (secondary aerosol formation) refers to the concentrations of secondary organic and inorganic (SOA and SIA, respectively) components (with a unit of  $\mu g m^{-3}$ ). In this study, the coal combustion refers to coal heating from the residential sector. Biomass burning includes open burning of agricultural biomass, 135 domestic biomass burning for cooking and heating, and biomass burning from biomass power plants and coal-fired power plants. Industry source is mainly derived from specific industrial processes in the iron and steel industrial base, metallurgical production plants for non-ferrous metals (e.g., titanium and molybdenum), and so on. Transportation source primarily comes from tailpipe emissions. It's worth mentioning that secondary aerosol formation originates from a series of atmospheric reactions. Some 140 identified sources (i.e., coal combustion, biomass burning, industrial processes, and transportation) may generate secondary inorganic and organic aerosols through the emission of their precursor components. The coefficient (with a unit of  $nmol min^{-1} m^{-3} source^{-1}$ ) reflect the intrinsic OP of each source.

### 2.3 Model setup

The DEHM can well capture many features of PM and its precursors' changes in large-scale space 145 (Christensen, 1997; Brandt et al., 2012; Im et al., 2019). To date, the DEHM model has been widely used in air pollution and health risk assessment research in Europe and Asia (Brandt et al., 2013a; b; Zare et al., 2014; Geels et al., 2015; Im et al., 2018; Im et al., 2019; Lehtomäki et al., 2020; Cramer et al., 2020; Liu et al., 2021; Geels et al., 2021; Thomas et al., 2022; Im et al., 2023), but this will be the first time that DEHM is applied to estimate OP. Thus, The DEHM model system was used to simulate 150 the pollutant concentrations in 2014 by using a two-way nested domain in this study (Kumar et al., 2016). A mother domain with a resolution of 150 km × 150 km was employed on a polar stereographic projection, true at 60°N to cover the entire Northern Hemisphere. The nested domain covered the whole of China consisting of 150 × 150 grid cells with a resolution of 50 km × 50 km, which was used for the analysis. The mother domain provided initial and boundary conditions for the nested domain. Vertically, there 155 were 29 unevenly distributed layers, with the highest level reaching 100 hPa, and the lowest layer was approximately 20 m in height. The meteorological fields were simulated using the WRF model (Skamarock et al., 2008) with the same domain and resolution driven by global meteorological data

obtained from the ERA5 dataset and the Community Earth System Model (CESM) global model, respectively. The simulations utilized the Revised MM5 surface layer scheme, the Yonsei University (YSU) boundary layer parameterization scheme (Hong et al., 2006), the Multi-scale Kain-Fritsch cumulus parameterization scheme (Zheng et al., 2016), the CAM longwave radiation scheme, the CAM shortwave radiation scheme (Skamarock et al., 2021), etc. The gas-phase chemistry module included 66 species, 9 primary particles (including natural particles such as sea salt), and 138 chemical reactions and was based on the scheme by Strand et al. (1994) (Brandt et al., 2012). The gas-phase species considered in this study included SO<sub>2</sub>, NO<sub>2</sub>, CH<sub>4</sub>, C<sub>2</sub>H<sub>6</sub>, etc. PM<sub>2.5</sub> was formed by BC, OC, sea salt, ammonium (NH<sub>4</sub><sup>+</sup>), nitrate (NO<sub>3</sub><sup>-</sup>), sulfate (SO<sub>4</sub><sup>2-</sup>), and secondary organic aerosols (SOA), among others (Frohn et al., 2022). Biogenic volatile organic compounds (BVOCs), such as isoprene, contributed to the formation of SOA (Zare et al., 2012). Further details on the configuration of the chemical scheme and the list of chemical reactions can refer to the literature (Zare et al., 2012; Brandt et al., 2012; Collin, 2020; Frohn et al., 2022). The SOA were calculated using the volatility basis set (see details in Im et al. (2019)). In addition to the anthropogenic emissions, DEHM also includes emissions from biogenic emissions, such as vegetation, sea salt, lightning, soil, etc. [The current version of the DEHM model does not include wind-blown, resuspended dust emissions or road dust nor aerosol-radiation or radiation-cloud interactions.](#) The time resolution of the DEHM model output is one hour.

In the current study, the DEHM model used anthropogenic emissions from the Emissions Database for Global Atmospheric Research – Hemispheric Transport of Air Pollution (EDGAR-HTAP) database and Eclipse V6. The biomass burning emissions are obtained from the Global Fire Assimilation System (GFAS) from ECMWF (Kaiser et al., 2012), which has a horizontal resolution of a 0.1° × 0.1° on a daily time basis. Natural emissions for DEHM are based on the Global Emissions Initiative (GEIA, Zare et al., 2012; Frost et al., 2013) with monthly inventories for emissions of nitrogen oxides from soil and lightning, and annual inventories for emissions of ammonia from natural sources. The production of sea salt (Soares et al., 2016) and biogenic volatile organic compounds (Zare et al., 2014) are calculated online in the model as a function of meteorological parameters like wind speed and temperature (Frohn et al., 2022).



## 185 2.4 Sensitivity scenarios

### 2.4.1 Relative contributions from meteorological conditions and emissions

Table 1 summarizes the scenarios for assessing the relative contributions of meteorological conditions and emissions to PM<sub>2.5</sub> and OP variability in 2014. ERA5 (Hersbach et al., 2020; ERA, 2023) is a global reanalysis dataset that is based on the assimilation of historical observations and model data. Studies  
190 (Thomas et al., 2021; Xu et al., 2022) have demonstrated that ERA5 performs well relative to MERRA, NCEP, and ERA-Interim, with higher temporal and spatial resolutions. Therefore, Scenarios C<sub>1</sub> and C<sub>2</sub> used ERA5 as input to WRF. Considering the robust representation of aerosol effective radiative forcing and good predictive capabilities for key surface variables in CESM (2023) (García-Martínez et al., 2020; Richter et al., 2022), Scenario C<sub>3</sub> utilized meteorological data based on CESM version 2.1.1  
195 (Danabasoglu et al., 2020) climate model as input for WRF. Scenarios C<sub>2</sub> and C<sub>3</sub> employed the Eclipse V6 emissions inventory, while Scenario C<sub>1</sub> used the EDGAR-HTAP inventory.

The ECLIPSE project by the International Institute for Applied Systems Analysis (IIASA) aims to generate a global gridded anthropogenic emission inventory for various emission scenarios. The Greenhouse Gas - Air Pollution Interactions and Synergies (GAINS) model has been employed to  
200 estimate emissions of air pollutants and GHGs (such as SO<sub>2</sub>, NO<sub>x</sub>, NH<sub>3</sub>, NMVOC, BC, OC, OM, PM<sub>2.5</sub>, PM<sub>10</sub>, CO, CH<sub>4</sub>) using source characteristics and emission factors at a resolution of 0.5° × 0.5° latitude-longitude (Upadhyay et al., 2020; Eclipse, 2020). The following sector-layers are available: energy, industry, solvent use, transport, domestic combustion, agriculture, open burning of agricultural waste, waste treatment. a number of scenarios are provided for which the key economic assumptions and energy  
205 use originate from IEA World Energy Outlook (IEA, 2011), the POLES model, or Energy Technology Perspectives (IEA, 2012) for the period 2010-2050, while statistical data for the period 1990-2010 came from IEA. For agriculture the FAO databases and long-term global projections were used (Alexandratos et al., 2012). It is noteworthy that this inventory considers China's 13th 5-year plan. The EDGAR-HTAP (Joint et al., 2011; Crippa et al., 2023) emission inventory endeavors to employ official or  
210 scientific inventories within a national or regional scale, with a spatial resolution of 0.1° × 0.1°. EDGAR-HTAP comprehensively accounts for all major emission sectors, including residential, transportation, industrial, energy, and agricultural sectors. EDGAR offers independent emission estimates for various pollutants, including CO, CH<sub>4</sub>, SO<sub>2</sub>, NO<sub>x</sub>, NMVOC, NH<sub>3</sub>, PM<sub>2.5</sub>, PM<sub>10</sub>, BC, and OC. These estimates

follow a standardized methodology provided by the Intergovernmental Panel on Climate Change (IPCC).

215 The data from EDGAR allows for comparisons with emission reports published by European Member States or Parties under the United Nations Framework Convention on Climate Change (UNFCCC) (Kumar et al., 2023).

Equations (2~5) were used to quantitatively evaluate the contributions of meteorological conditions and emission inventories.

$$\text{Con(Met)} = \frac{C_2 - C_3}{C_3} \quad (2)$$

$$\text{Con(Emi)} = \frac{C_1 - C_2}{C_2} \quad (3)$$

$$\text{NCon(Met)} = \frac{\text{abs(Con(Met))}}{\text{abs(Con(Met))} + \text{abs(Con(Emi))}} \quad (4)$$

$$\text{NCon(Emi)} = \frac{\text{abs(Con(Emi))}}{\text{abs(Con(Met))} + \text{abs(Con(Emi))}} \quad (5)$$

220 where,  $C_1$ ,  $C_2$  and  $C_3$  represent the  $\text{PM}_{2.5}$  concentrations and OP from scenarios  $C_1$ ,  $C_2$  and  $C_3$ , respectively.  $\text{Con(Met)}$  represents the impact of changing meteorological datasets on changes in  $\text{PM}_{2.5}$  and OP.  $\text{Con(Emi)}$  represents the impact of changing emission inventory on changes in  $\text{PM}_{2.5}$  and OP.  $\text{NCon(Met)}$  and  $\text{NCon(Emi)}$  represent the normalized contributions of meteorology and emission. In the equations, the  $\text{abs}$  function represents the absolute value of the quantity in parentheses.

225 **Table 1. Emission inventory and meteorological datasets in three simulation scenarios.**

Scenarios	Emission inventory	Meteorological datasets
$C_1$	EDGAR-HTAP	ERA5
$C_2$	Eclipse V6	ERA5
$C_3$	Eclipse V6	CESM

#### 2.4.2 Relative contributions from individual emissions

As mentioned above, OP's main source contributions include five parts, i.e., coal combustion, biomass burning, secondary aerosol formation, industrial sources, and transportation sources (Equation (1)), we conducted perturbation experiments targeting these five sources to quantitatively assess their contributions to  $\text{PM}_{2.5}$  concentrations and OP (Figure 2). These experiments were carried out within the three scenarios described in Section 2.4.1, and we performed a total of 15 runs. Under the non-perturbation condition (referred to as the NPC case), all aforementioned emission sources were considered. Under the perturbation condition (referred to as the PC case), reduction designs were

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implemented for emissions from coal combustion, biomass burning, industrial sources, and transportation sources. The emissions of both primary aerosols and tracegases from each individual source are reduced by 30%. The choice of 30 % was motivated by the consideration that the perturbation would be large enough to produce a sizeable impact (i.e., more than numerical noise) even at long distances, while small enough to be in the near-linear atmospheric chemistry regime (Galmarini et al., 2017; Im et al., 2019). Notably, the transportation sector in the DEHM model only considers tailpipe emissions, excluding non-exhaust emissions from vehicles like road dust, brake dust, and tire wear. The contribution of tailpipe emissions in the transportation sector to OP is estimated by incorporating simulated values of primary PM<sub>2.5</sub> concentrations from tailpipe emissions into Equation (1). In this study, the transportation sector refers to road transport, excluding other transportation sources like ships and airplanes. Additionally, to estimate the PM<sub>2.5</sub> concentrations and OP from coal and biomass burning, it is necessary to obtain the percentage contributions of PM<sub>2.5</sub> emissions from coal combustion for residential heating, domestic biomass burning for cooking and heating to PM<sub>2.5</sub> emissions of the residential sector, respectively, as well as the percentage contributions of PM<sub>2.5</sub> emissions from biomass burning in power plants to the total PM<sub>2.5</sub> emissions from the power sector. The percentage contributions of each anthropogenic source can be estimated using Equations (6–8).

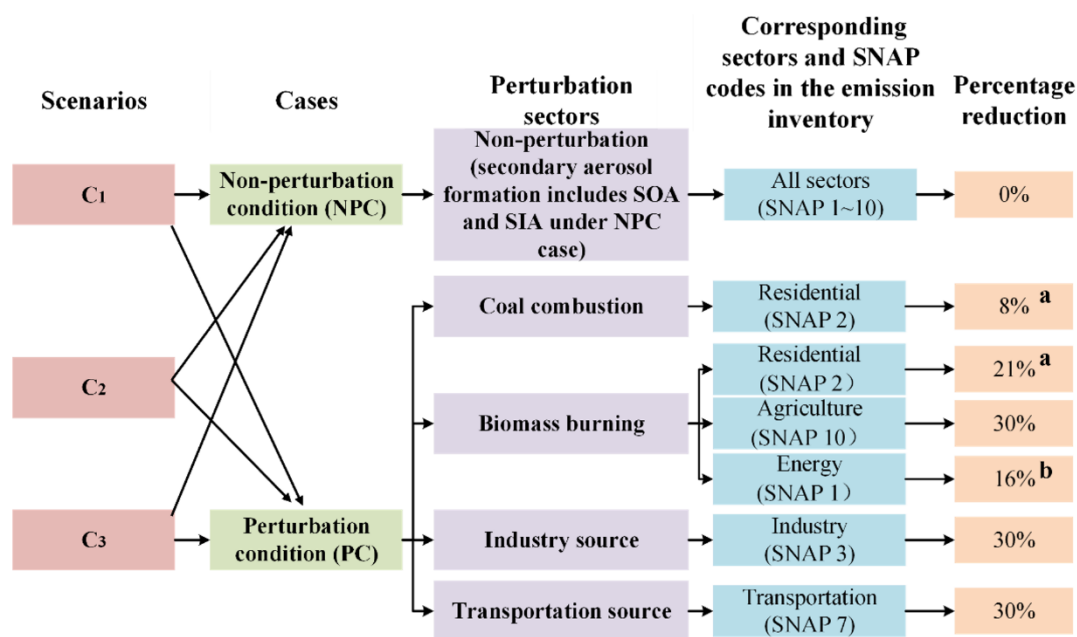
$$PC_{re-j} = \frac{E_{re-j}}{E_{re}} \quad (6)$$

$$E_{pp-bi} = EF \times FQ \quad (7)$$

$$PC_{pp-bi,cf} = \frac{E_{pp-bi} + E_{pp,cf}}{E_{pp}} \quad (8)$$

where,  $PC_{re-j}$  denotes the percentage contribution of PM<sub>2.5</sub> emissions from the residential subsector j (including coal cooking, coal heating, biomass cooking, biomass heating, clean energy, and nonresidential) to the total PM<sub>2.5</sub> emissions from the residential sector.  $E_{re-j}$  represents the PM<sub>2.5</sub> emissions from the residential subsector j, while  $E_{re}$  represents the total PM<sub>2.5</sub> emissions from the residential sector. The values of  $E_{re-j}$  and  $E_{re}$  are obtained from the literature (Yun et al., 2020). After calculation, the values for  $PC_{re\text{coal cooking}}$ ,  $PC_{re\text{coal heating}}$ ,  $PC_{re\text{biomass cooking}}$ , and  $PC_{re\text{biomass heating}}$  are determined to be 21%, 27%, 33%, and 19%, respectively.  $E_{pp-bi}$  refers to the PM<sub>2.5</sub> emissions from biomass power plants,  $EF$  refers to the PM<sub>2.5</sub> emission factor of biomass power plants, and  $FQ$  refers to the fuel quantity.  $PC_{pp-bi,cf}$  refers to the percentage contribution of PM<sub>2.5</sub> emissions from biomass power plants and coal-fired power plants to the PM<sub>2.5</sub> emissions of the power plants. To accelerate carbon

260 reduction in coal-fired power generation, the Chinese government has issued a series of policies supporting and encouraging the coupling of coal and biomass for power generation (Mao, 2017). This undoubtedly adds complexity to distinguishing between  $PM_{2.5}$  emissions from coal combustion and biomass fuel. Furthermore, in the energy strategy where biomass serves as a clean alternative to fossil fuels, the scale of biomass utilization and the biomass power generation industry in China continue to expand (Lin et al., 2021). Considering the aforementioned reasons, we include  $PM_{2.5}$  emissions from coal-fired power plants in our analysis.  $E_{pp\_cf}$  refers to the  $PM_{2.5}$  emissions from coal-fired power plants, and  $E_{pp}$  refers to the  $PM_{2.5}$  emissions from power plants.  $EF$ ,  $FQ$ ,  $E_{pp\_cf}$ , and  $E_{pp}$  are obtained from the literature (Zheng et al., 2018; Tong et al., 2018; Yun et al., 2020; MEE, 2020; Wang et al., 2020; Tang et al., 2020; Lin et al., 2021; Chen et al., 2022). After calculation,  $PC_{pp\_bi\_cf}$  is determined to be 270 54%. Previous studies (Hodan et al., 2004; Chen et al., 2018; Zhang et al., 2022) showed that in China, the proportion of secondary and primary  $PM_{2.5}$  mass to the total  $PM_{2.5}$  mass is close, so we assume that they account for 50% respectively. Figure 2 showed the emission reduction design for perturbed emissions.



275 **Figure 2. Emission reduction design for perturbed emissions; <sup>a</sup> was obtained from the literature (Yun et al., 2020), <sup>b</sup> were obtained from the literature (Zheng et al., 2018; Tong et al., 2018; Yun et al., 2020; MEE, 2020; Wang et al., 2020; Tang et al., 2020; Lin et al., 2021; Chen et al., 2022).**

Furthermore, the  $PM_{2.5}$  concentrations for each sector is calculated using Equations (9~11).

$$C_{P,i} = \frac{C_{NPC,i} - C_{PC,i}}{30\%} \quad (9)$$

$$C_{P,primary\ PM_{2.5}} = C_{P,total\ PM_{2.5}} - C_{P,SIA} - C_{P,SOA} \quad (10)$$

$$C_{secondary} = C_{NPC,SOA} + C_{NPC,SIA} \quad (11)$$

where,  $i$  refers to the type of pollutants, i.e., total  $PM_{2.5}$ , SOA, SIA, and primary  $PM_{2.5}$ .  $C_{NPC,i}$  represents

280 concentrations of the pollutant  $i$  in the NPC case.  $C_{PC,i}$  represents concentrations of the pollutant  $i$  in the

PC case.  $C_{P,i}$  represents concentrations of the pollutant  $i$  by the specific emission sector P which is perturbed (perturbation sectors P include coal combustion, biomass burning, industry, traffic source).

$C_{P,primary\ PM_{2.5}}$  represents concentrations of primary  $PM_{2.5}$  by the perturbation sectors P.  $C_{secondary}$  represents  $PM_{2.5}$  concentrations by the secondary aerosol formation.

## 285 2.5 Probability density function

Taking into account the substantial spatial heterogeneity of  $PM_{2.5}$  concentrations and OP, we employ PDF to characterize the statistical distribution characteristics of  $PM_{2.5}$  concentrations and OP across China. This offers a more generalized and robust probability for criteria limits. In this study, all three functional types (Lognormal, exponential, and Gamma) were tested for annual average of  $PM_{2.5}$  concentrations and OP at the monitoring stations. To determine the representative distributions for the datasets, we further performed goodness-of-fit tests such as the Sum of Squared Error (SSE) and the Kolmogorov-Smirnov (K-S) test (de Melo et al., 2000), using the fitter package in Python.

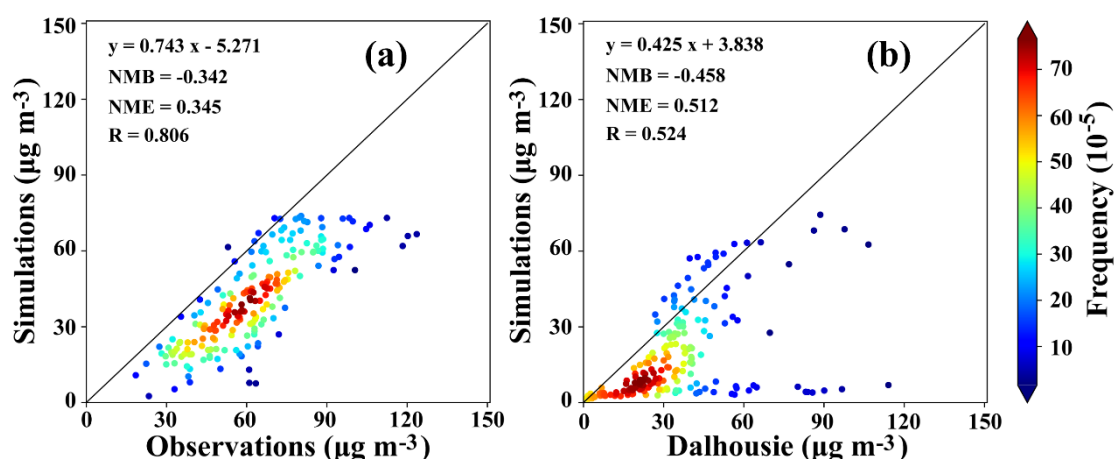
## 3. Results and discussion

### 3.1 Model evaluation

295 The hourly observation data was obtained from the Ministry of Ecology and Environment of China (MEE, 2014). The MEE website first released  $PM_{2.5}$  measurement data in January 2013. In accordance with the Chinese environmental protection standards, the hourly  $PM_{2.5}$  concentrations are measured using the micro-oscillation balance method and beta absorption method, with an uncertainty of less than  $5\ \mu g\ m^{-3}$  (Zeng et al., 2021). The  $PM_{2.5}$  monitoring stations are primarily distributed in urban areas, particularly in major metropolitan areas of China (Zeng et al., 2021). In 2014, the observation stations are mainly concentrated in eastern China, while stations in western China are limited. Therefore, in the

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present study, we also evaluated with the gridded annual-mean global reanalysis Dalhousie surface  $PM_{2.5}$  dataset (van Donkelaar et al., 2021), which combines satellite retrievals of aerosol optical depth, chemical transport modeling, and ground-based measurements. The Dalhousie dataset compensated for the non-uniform distribution spatially of observation stations to comprehensively evaluate the performance of the DEHM model. The density scatter plot of model performance and evaluation for China in scenario  $C_1$  based on annual mean  $PM_{2.5}$  observations from MEE and  $PM_{2.5}$  derived from the Dalhousie dataset are shown in Figure 3. Overall, the model performance in terms of correlation coefficient ( $R$ ) and normalized mean error (NME) calculated based on annual mean observations met the performance criteria suggested by Emery et al. (2017) ( $NME < 0.5$ ,  $R > 0.4$ ), and the normalized mean bias (NMB) was also close to the performance criteria suggested by Emery et al. (2017) ( $NMB \leq \pm 0.3$ ). Compared to the observations, the model performance in terms of  $R$ , NME, and NMB calculated based on the Dalhousie dataset was slightly poorer but still close to the performance criteria suggested by Emery et al. (2017). Additionally, this study evaluated the model performance in scenarios  $C_2$  and  $C_3$  (Figure S2). It was found that under scenarios  $C_2$  and  $C_3$ , the model performance in terms of  $R$  and NME, calculated based on both annual mean observations and Dalhousie dataset met the performance criteria suggested by Emery et al. (2017). The NMB under scenarios  $C_2$  and  $C_3$  calculated based on both annual mean observations and Dalhousie dataset were also close to the performance criteria suggested by Emery et al. (2017). Therefore, the simulated annual mean  $PM_{2.5}$  concentrations in scenarios  $C_1$ ,  $C_2$  and  $C_3$  are considered reliable.

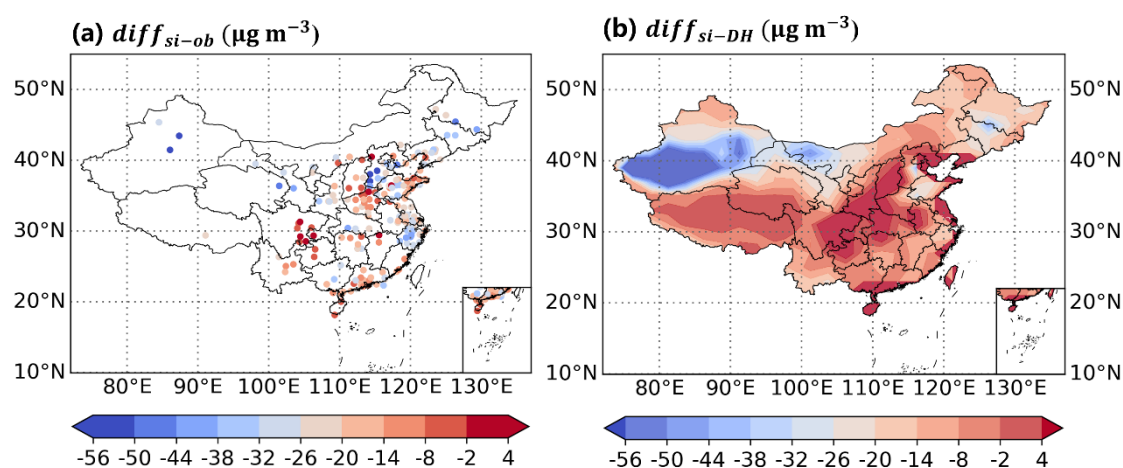


**Figure 3.** Density scatterplots of model performance and validation for China in scenario  $C_1$  based on (a) annual mean  $PM_{2.5}$  observations from MEE and (b) annual mean  $PM_{2.5}$  derived from the Dalhousie dataset in 2014.

325 To verify the spatial accuracy, a comparison of the spatial distribution of simulated and observed  $\text{PM}_{2.5}$ , both from MEE and Dalhousie, was conducted. Figure 4 showed the spatial distribution of the annual mean simulated minus annual mean observed values (denoted as  $\text{diff}_{si-ob}$ ) (Figure 4a), as well as the spatial distribution of the annual mean simulated values minus the Dalhousie dataset (denoted as  $\text{diff}_{si-DH}$ ) (Figure 4b). Both Figure 4a and Figure 4b indicated that the majority of regions (central and eastern China) exhibited differences ranging from  $-18 \mu\text{g m}^{-3}$  to  $0 \mu\text{g m}^{-3}$ , which is an underestimation of 37% compared to the average annual observations. The simulated  $\text{PM}_{2.5}$  concentrations in eastern, central, northeastern, and western China were 37%, 21%, -49%, and 41% lower than the observations, respectively; the simulated values were 28%, 3%, 54%, and 48% higher than the Dalhousie dataset, respectively. The disparities in model performance across regions may be attributed to uncertainties in

335 the simulation of meteorological fields, coupled with insufficient consideration of species in the reaction processes within the model. Considering the existing literature (Huang et al., 2021; Jia et al., 2021), it is known that bias within approximately 50% is acceptable. For example, the  $\text{PM}_{2.5}$  concentrations in East China in 2014 simulated by Jia et al. (2021) was overestimated by 48%. Shi et al. (2021) also reported  $\text{PM}_{2.5}$  concentrations being overestimated or underestimated by 40% compared to observed

340 values. Hence, the simulated bias in this study falls within an acceptable range, meeting the research requirements.



345 **Figure 4. Spatial distribution of the annual mean simulated minus annual mean observed values (a), as well as the spatial distribution of the annual mean simulated values minus the Dalhousie dataset (b) for China in 2014 under scenario C1.**

Similarly, the model performance over time scales was also investigated. Scatter density plots and distribution characteristics of monthly average observations and simulations for all monitoring sites in

2014 were depicted in Figure S1 and Figure 5, respectively. From Figure 5, it can be observed that the simulated values closely align with the observed values from April to September. However, in other months, there was a slightly poorer alignment between simulated and observed values. Nonetheless, considering the overall performance throughout the year, as analyzed in conjunction with Figure S1, it can be deduced that both the correlation R and NME met the performance criteria suggested by Emery et al. (2017) for all months except December. Furthermore, the results in Figure 4 indicated that the bias across various regions in DEHM is acceptable. Consequently, on an aggregate level for China, the model demonstrates acceptable performance in simulating monthly average  $PM_{2.5}$  concentrations.

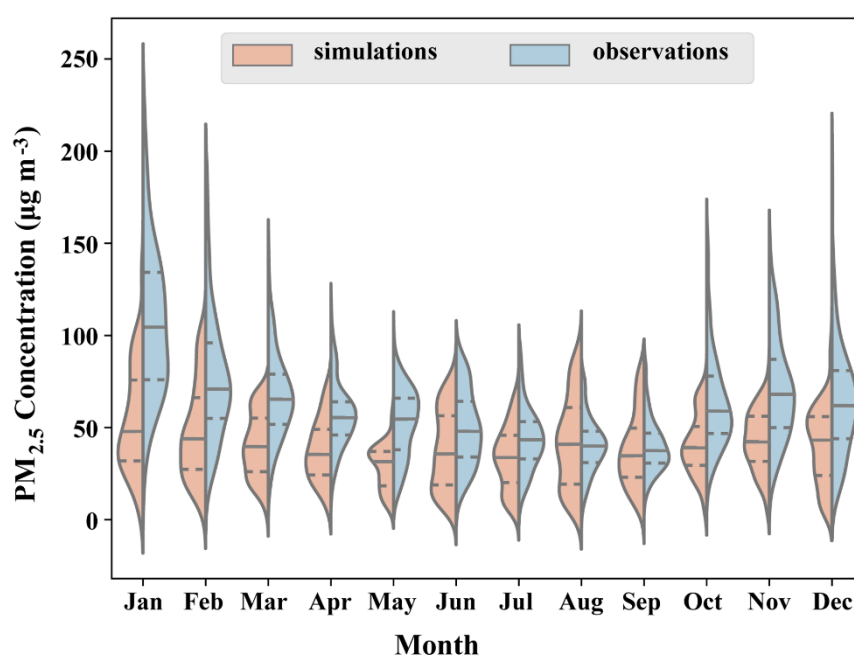


Figure 5. Violin plots of monthly average from MEE observations and simulations averaged over various observation stations for China in 2014 under scenario C<sub>1</sub>; The red and blue colors represent the statistical distribution of simulated and observations, respectively; The width of the violin represents the sample size; The solid black line inside the violin indicates the median. The upper and lower dashed black lines within the violin indicate the upper quartile (the 75th percentile) and lower quartile (the 25th percentile), respectively.

### 3.2 Spatial distribution characteristics of $PM_{2.5}$ and OP

To learn about the spatial distributions of  $PM_{2.5}$  concentrations and OP, we plot maps of surface  $PM_{2.5}$  and OP for scenario C<sub>1</sub> (Figure 6a and 6b) and quantified the average annual  $PM_{2.5}$  concentrations and OP across different regions of China (Figure 6c). Figure 6d depicted the geographical location of the study area. High  $PM_{2.5}$  concentrations and High OP are mainly located in central and eastern urban clusters. Low  $PM_{2.5}$  concentrations and Low OP are mainly distributed in northeastern and western China.



The results in Figure 6c indicated that the annual average  $PM_{2.5}$  concentrations/OP in eastern, central, northeastern, and western China are  $33 \mu g m^{-3} / 1.4 nmol min^{-1} m^{-3}$ ,  $46 \mu g m^{-3} / 2.0 nmol min^{-1} m^{-3}$ ,  $19 \mu g m^{-3} / 0.8 nmol min^{-1} m^{-3}$ , and  $12 \mu g m^{-3} / 0.5 nmol min^{-1} m^{-3}$ , respectively.

Due to differences in city types, pollutant emission intensities, and pollutant chemical components in different regions, there are significant spatial heterogeneity in  $PM_{2.5}$  concentrations and therefore in OP (see Sect. 3.4 and Figure 11 for details). Due to high population density, socio-economic activities and winter heating needs, large amounts of anthropogenic emissions, especially from industry, transportation, coal burning and biomass burning, exacerbate  $PM_{2.5}$  and redox active component pollution.

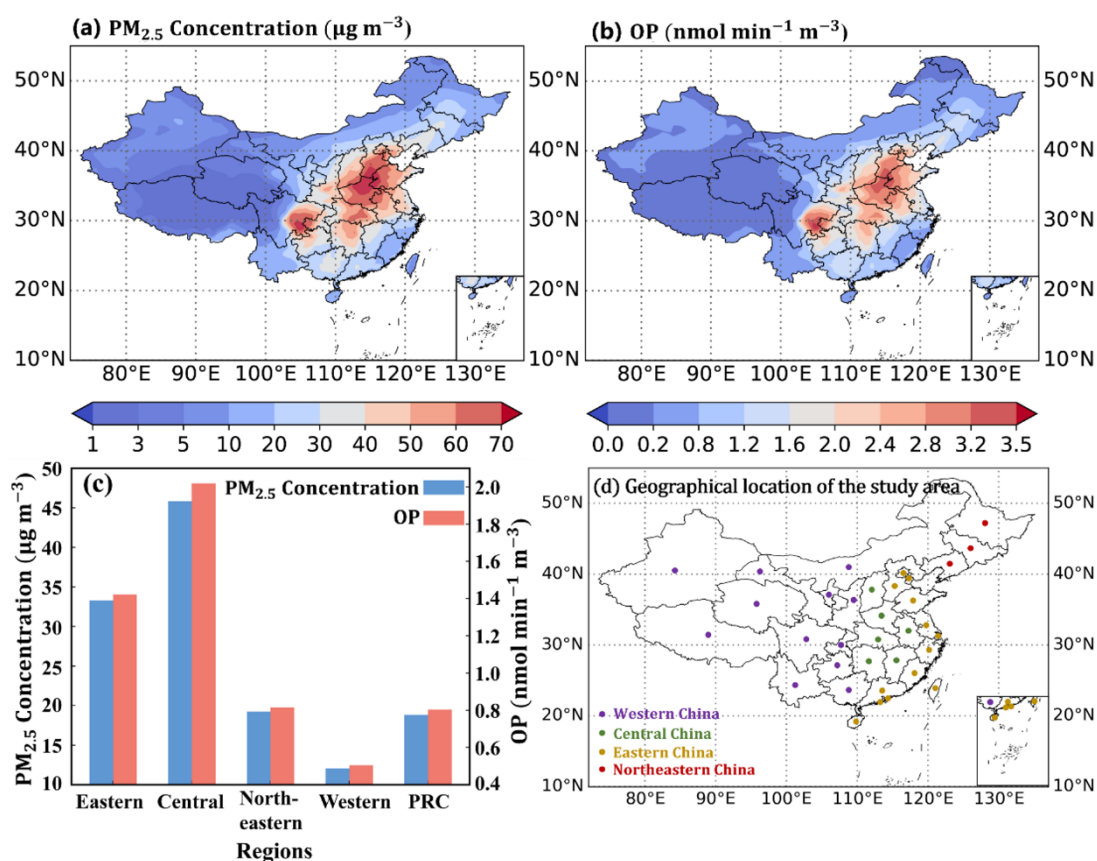


Figure 6. Spatial clustering of annual mean  $PM_{2.5}$  concentrations (a) and annual mean OP (b) in China, annual mean  $PM_{2.5}$  concentrations and annual mean OP (c) in different regions of China in 2014 under scenario C<sub>1</sub>, and geographical location of the study area (d); the central region of China comprises Shanxi, Anhui, Jiangxi, Henan, Hubei, and Hunan provinces; the eastern region of China comprises Beijing, Tianjin, Hebei, Shanghai, Jiangsu, Zhejiang, Fujian, Shandong, Guangdong, Hainan, Hong Kong, Macao, and Taiwan; It should be noted that the eastern region in this study includes Hong Kong, Macao and Taiwan; the western region of China consists of twelve provinces (autonomous regions and municipalities): Inner Mongolia, Guangxi, Chongqing, Sichuan, Guizhou, Yunnan, Tibet, Shaanxi, Gansu, Qinghai, Ningxia and Xinjiang; the northeastern region of China comprises Liaoning, Jilin and Heilongjiang provinces.

To quantitatively analyze the regional distribution characteristics of  $PM_{2.5}$  concentrations and OP in China, we determined the distribution function that is suitable for a specific dataset (Table 3), investigated the frequency histogram (FH) of  $PM_{2.5}$  concentrations and OP, fitted the PDF, and then obtained the cumulative distribution function (CDF) by integrating PDF, as shown in Figure 7. It was found that the gamma distribution performed the best in fitting  $PM_{2.5}$  concentrations and OP from Table 2. Considering the test results, the gamma distribution was used to explore the spatial distribution characteristics of  $PM_{2.5}$  concentrations and OP. Figure 7a depicted the probability distribution of  $PM_{2.5}$  concentrations, while Figure 7b depicted the probability distribution of OP. The wide distribution interval indicated that both  $PM_{2.5}$  concentrations and OP have a similar and large spatial heterogeneity. According to the FH, the highest frequency density of  $PM_{2.5}$  concentrations ranges from 10.5 to 12.9  $\mu g m^{-3}$ ; The maximum frequency density of OP ranges from 0.26 to 0.34  $nmol min^{-1} m^{-3}$ . This reflects the overall pollution levels of  $PM_{2.5}$  and OP in the Chinese region. Taking into account the annual average  $PM_{2.5}$  concentrations limits set out in China's ambient air quality standard (AAQS, 2012), we focused on primary (15  $\mu g m^{-3}$ ) and secondary concentrations (35  $\mu g m^{-3}$ ) limits. The PDF and CDF results showed that 85% of the total area was above the primary concentrations limit and 40% was above the secondary concentrations limit. In addition, 36% of regions in China have an OP below 1.00  $nmol min^{-1} m^{-3}$ , 41% have an OP between 1.00 and 2.00  $nmol min^{-1} m^{-3}$ , and 23% have an OP above 2.00  $nmol min^{-1} m^{-3}$ .

**Table 2. Goodness-of-fit test results for China in 2014 under scenario C<sub>1</sub>.**

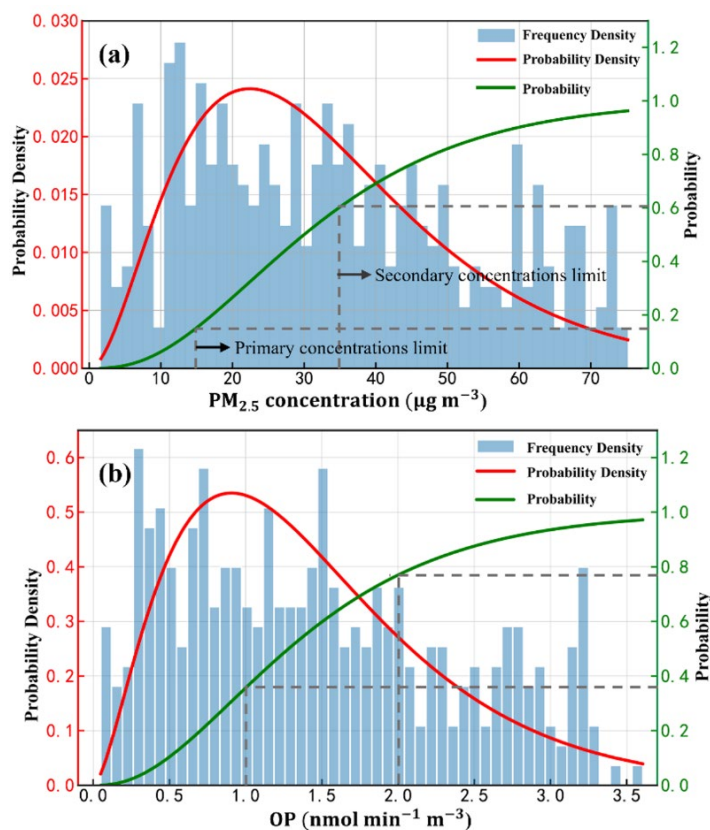
Item	goodness-of-fit test	Gamma	Lognormal	Exponential
$PM_{2.5}$ concentrations	SSE	0.002	0.023	0.003
	KS_pvalue	0.329	0.000	0.000
OP	Sumsquare_error	0.654	0.746	1.209
	KS_pvalue	0.231	0.271	0.000

Bold values indicate the best results.

\*Note:

SSE is Sum of Squared Error

KS\_pvalue is the P-value of the Kolmogorov-Smirnov test



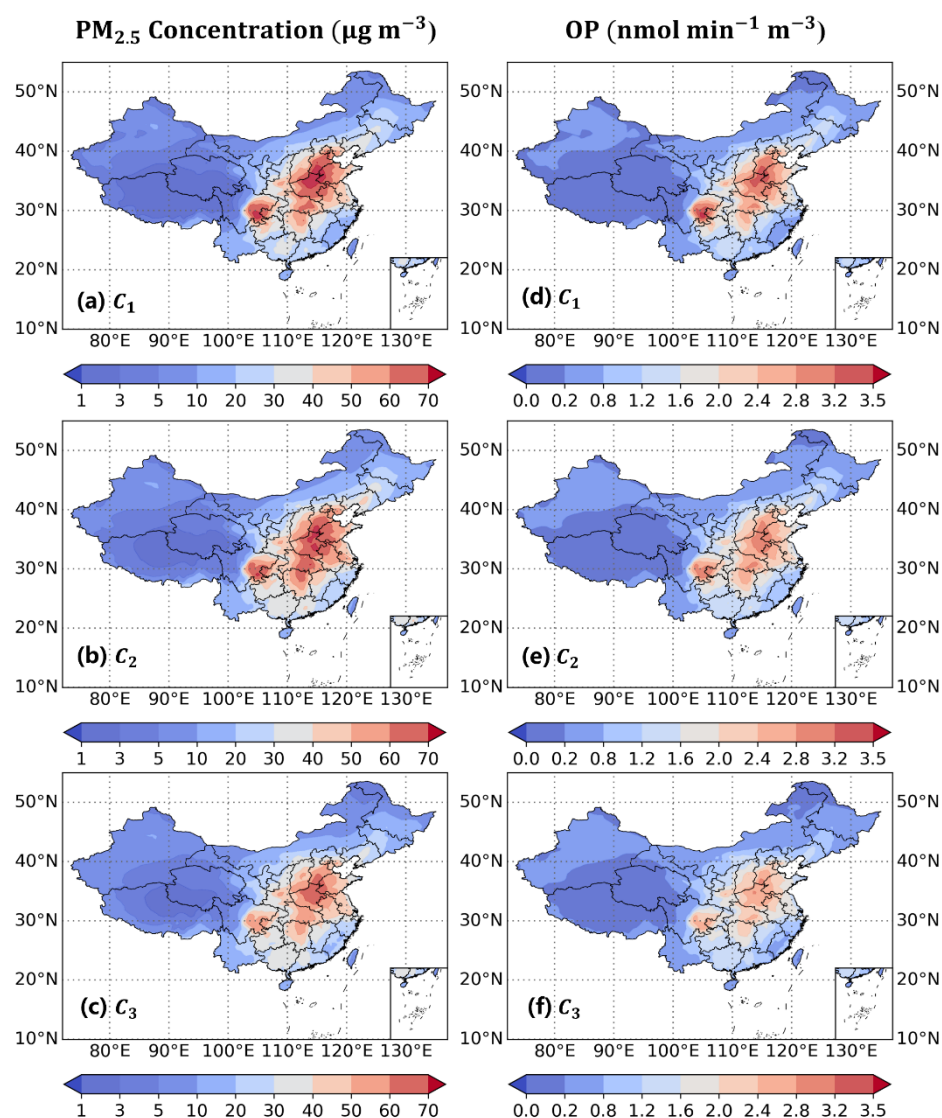
410 **Figure 7. Probability distribution of (a) annual mean  $PM_{2.5}$  concentrations and (b) annual mean OP for China in 2014 under scenario  $C_1$ .**

### 3.3 Contributions of meteorological conditions and emission inventories to the variations in $PM_{2.5}$ and OP

To determine the sensitivity of  $PM_{2.5}$  pollution and oxidation potential (OP) to meteorological conditions  
 415 (emission inventories), this study compared scenarios  $C_2$  and  $C_3$  ( $C_1$ ) and investigated the impacts and  
 contributions due to ERA5 and CESM (HTAP and Eclipse V6 emission inventories) on  $PM_{2.5}$  and OP.  
 Figure 8 illustrates the spatial distribution maps of  $PM_{2.5}$  concentrations and OP under scenarios  $C_1$ ,  $C_2$ ,  
 and  $C_3$ . Figure 9a presents the annual average  $PM_{2.5}$  concentrations and OP under different scenarios,  
 and Figure 9b shows the relative contributions of meteorological conditions and emission inventories.  
 420 From Figures 8 and 9, it can be observed that, compared to scenario  $C_2$ ,  $PM_{2.5}$  concentrations and OP are  
 lower in the western region and slightly higher in some eastern areas under scenario  $C_1$ , primarily due to  
 changes in emission inventories attributed to the inclusion or exclusion of specific local sources during  
 the compilation process. Compared to scenario  $C_3$ ,  $PM_{2.5}$  concentrations and OP are lower in the western  
 region and higher in some eastern areas under scenario  $C_2$ , primarily attributed to meteorological  
 425 contributions. For the entire China region, the transition in emission inventories from Eclipse V6 to

HTAP resulted in an overall decrease in  $PM_{2.5}$  concentrations of  $1.55 \mu g m^{-3}$ , approximately 7.61%, and a decrease in OP of  $0.0339 nmol min^{-1} m^{-3}$ , approximately 4.05%. The shift in meteorological data from CESM to ERA5 led to an increase in  $PM_{2.5}$  concentrations of  $1.22 \mu g m^{-3}$ , approximately 6.4%, and an increase in OP of  $0.0585 nmol min^{-1} m^{-3}$ , approximately 7.5%. According to the normalization process using Equations (4)–(5), meteorological conditions contributed approximately 45.6% to the variations in  $PM_{2.5}$  and approximately 65.0% to the variations in OP. Meanwhile, emission inventories contributed approximately 54.4% to the variations in  $PM_{2.5}$  and approximately 35.0% to the variations in OP. Our findings highlight the significance of the quality of model input data, including emission inventories and meteorological data, for model performance.

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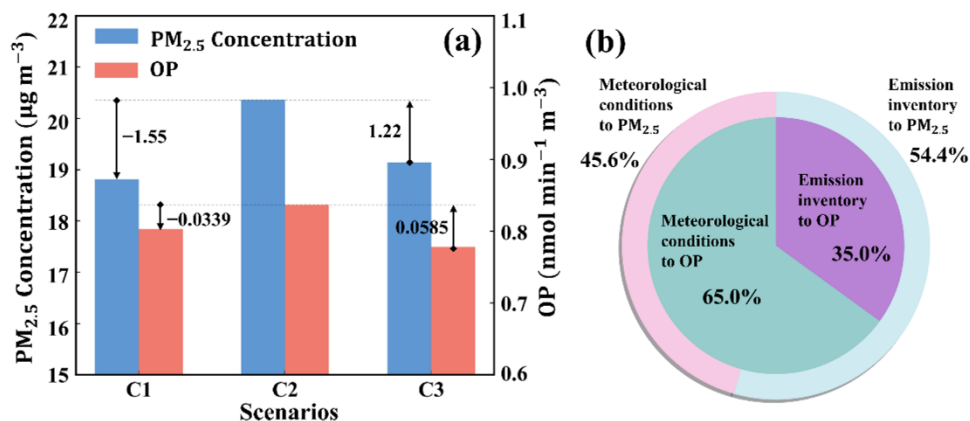


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**Figure 8.** Spatial distribution of annual mean  $PM_{2.5}$  concentrations and annual mean OP for China in 2014 under different scenarios; (a)–(c) are  $PM_{2.5}$  concentrations in scenarios  $C_1$ ,  $C_2$  and  $C_3$ , respectively; (d)–(f) is the OP in scenarios  $C_1$ ,  $C_2$ , and  $C_3$ , respectively; The meteorological datasets (emission inventories) employed

for scenarios C<sub>1</sub>, C<sub>2</sub>, and C<sub>3</sub> are ERA5 (EDGAR-HTAP), ERA5 (Eclipse V6), and CESM (Eclipse V6), respectively.

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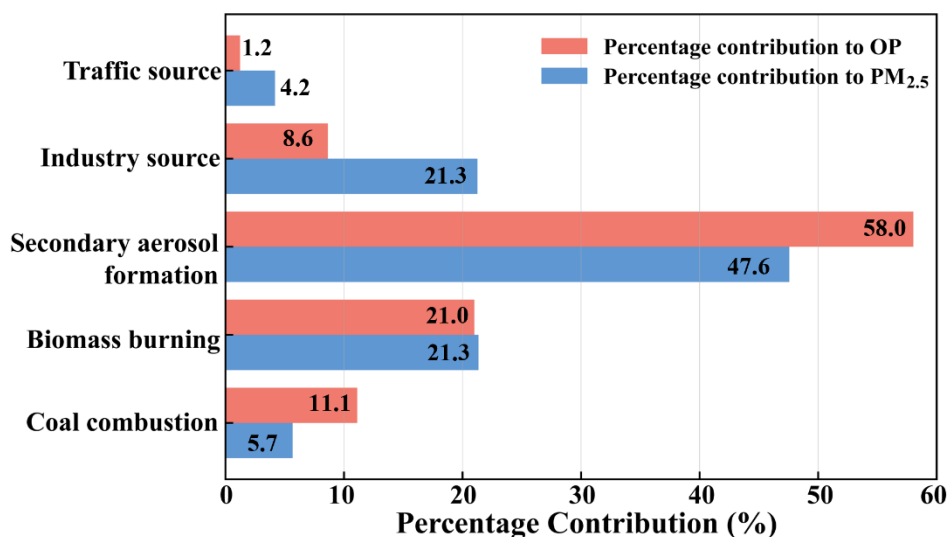
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Figure 9. (a) Average annual PM<sub>2.5</sub> concentrations and Average annual OP for China in 2014 under different scenarios; (b) The relative contribution of meteorological conditions and emission inventories to Average annual PM<sub>2.5</sub> and Average annual OP for China in 2014, with the outer circle representing PM<sub>2.5</sub> and the inner circle representing OP; The meteorological datasets (emission inventories) employed for scenarios C<sub>1</sub>, C<sub>2</sub>, and C<sub>3</sub> are ERA5 (EDGAR-HTAP), ERA5 (Eclipse V6), and CESM (Eclipse V6), respectively.

### 3.4 Contribution of anthropogenic emission sources to PM<sub>2.5</sub> and OP

To determine the impact of anthropogenic emissions on PM<sub>2.5</sub> and OP, we quantified their percent contribution (Figure 10). Secondary aerosol formation, biomass burning, industrial, coal combustion for residential heating, and transportation sources contributed 48%, 21%, 21%, 6% and 4% to PM<sub>2.5</sub>, respectively. Secondary aerosol formation, biomass burning, coal combustion for residential heating, industrial sources, and transportation sources contributed 58%, 21%, 11%, 9% and 1% to OP, respectively. This means that secondary aerosol formation and biomass burning are the main sources of PM<sub>2.5</sub> and OP.

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**Figure 10. Percentage contribution of different anthropogenic sources (coal combustion for residential heating, biomass burning, secondary aerosol formation, industry, and traffic) to total PM<sub>2.5</sub> concentrations and OP for China in 2014 under scenario C<sub>1</sub>.**

Thus, we explored the spatial distribution characteristics of PM<sub>2.5</sub> and OP from different anthropogenic  
 460 sources to reveal the spatial contributions of PM<sub>2.5</sub> concentrations and OP, as shown in Figure 11. It was  
 observed that the spatial distribution features of PM<sub>2.5</sub> concentrations and OP from each emission source  
 are similar to those in Figure 6, and they all adhere to the principle that the eastern region is higher than  
 the western.

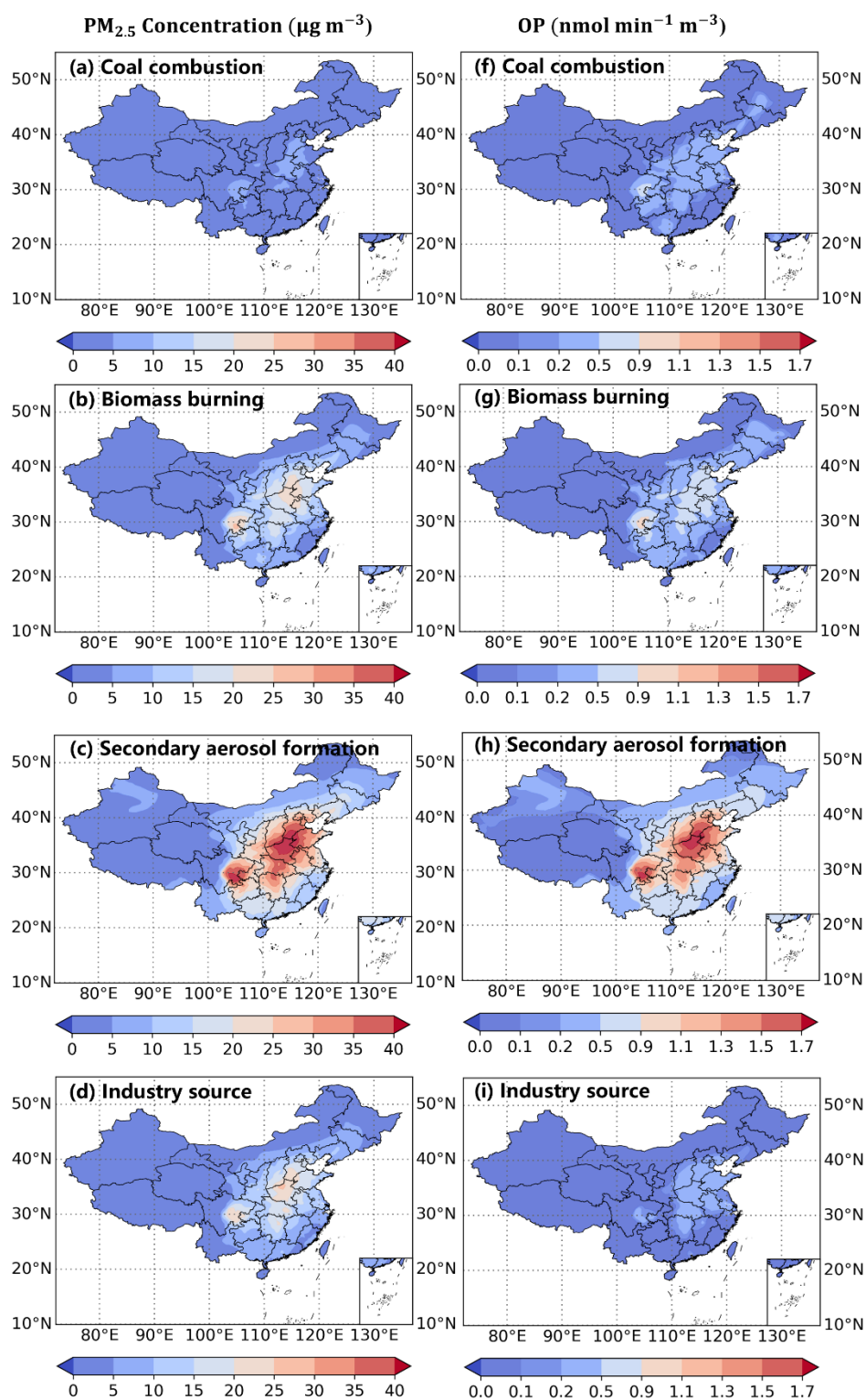
It can be seen from Figure 11 that the main reason that secondary aerosol formation is the main  
 465 anthropogenic source of both PM<sub>2.5</sub> concentrations and OP in China is due to the higher pollution levels,  
 more contributions to mass, and toxicity in the central and eastern regions. Relevant study (Molina et  
 al., 2023) has highlighted the significant contribution of secondary aerosol formation to particle mass  
 and intrinsic OP.

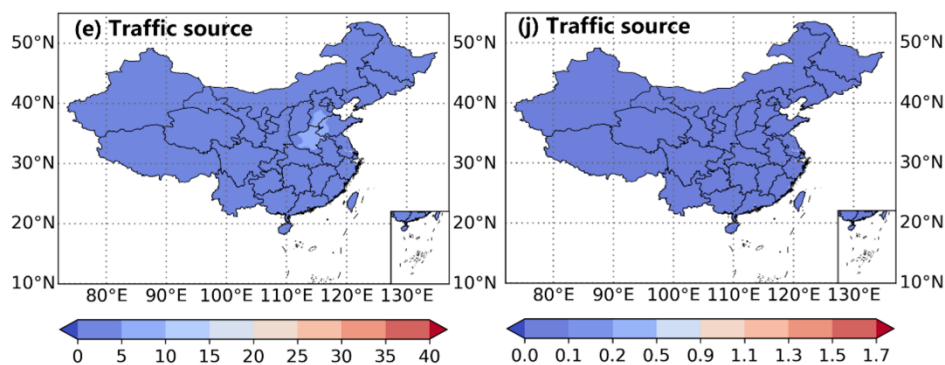
Due to Chinese crops, especially corn straw, and power plants are mainly concentrated in central and  
 470 eastern regions, northeast and part of the western region, as well as the bigger intrinsic OP (Equation (1)),  
 this results in biomass burning becoming the second contribution.

In this study, coal combustion refers to coal heating from the residential sector. Coal burning increases  
 secondary inorganic and organic aerosols in the air (Liu et al., 2018) which leads to stronger oxidative  
 toxicity. Thus, due to greater heating demand in locations with high population density and chilly winters  
 475 (e.g., the northern part of the central and eastern China), PM<sub>2.5</sub> concentrations and OP linked to coal  
 burning are higher.

Industrial emissions are mainly derived from specific industrial processes in the iron and steel industrial  
 base, metallurgical production plants for non-ferrous metals (e.g., titanium and molybdenum), and so on.  
 This is one main source for metals. Due to the correlation between these transition metals and OP (Fang  
 480 et al., 2017; Liu et al., 2018), China's four industrial zones (Liaozhong-South Heavy Industry Base,  
 Beijing-Tianjin-Tangshan Industrial Base, Shanghai-Nanjing-Hangzhou Industrial Base, and Pearl River  
 Delta Light Industry Base) are important contributors to PM<sub>2.5</sub> and OP emissions from industrial sources.  
 The transportation sector in the DEHM model only considers tailpipe emissions, excluding non-exhaust  
 emissions from vehicles like road dust, brake dust, and tire wear, which is a main reason that the traffic  
 485 sources exhibit the lowest contribution to PM<sub>2.5</sub> concentrations/OP. Moreover, it can be observed from

Figure 11 that the sector's emissions are mainly concentrated in a small number of regions, such as Henan, Hebei, and Shandong. This is valid for the top three provinces in terms of vehicle particulate matter and nitrogen oxide emissions in 2014 according to China Annual Vehicle Pollution Prevention and Control Report (MEE, 2015).





495 **Figure 11. Spatial distribution of annual mean  $PM_{2.5}$  concentrations and annual mean OP from different anthropogenic sources for China in 2014 under scenario C1; (a)~(e) are  $PM_{2.5}$  concentrations derived from coal combustion for residential heating, biomass burning, secondary aerosol formation, industry, and traffic respectively; (f)~(j) are the OP derived from coal combustion for residential heating, biomass burning, secondary aerosol formation, industry, and traffic respectively.**

### 3.5 Uncertainty of OP estimates

500 OP is considered an important indicator of  $PM_{2.5}$  toxicity and is associated with adverse health effects. Linking the predicted health effects of aerosols to OP may be more relevant than considering  $PM_{2.5}$  mass alone (Alwadei et al., 2020). However, previous studies of OP in China have mainly focused on local areas, and OP and its source are very different in space and time (Wen et al., 2023), which makes the health research of OP challenging. At present, there are two kinds of methods for evaluating OP of  $PM_{2.5}$ :

505 cellular method and non-cellular method. The reproducibility of cellular methods is poor, and it is difficult to achieve a large sample size analysis. And the choice of cell type or cell line can significantly affect the OP results (Xing et al., 2023). The non-cellular method has the advantages of fast speed, simple operation, high reproducibility, and low cost. The most common non-cellular methods are DTT, AA, GSH, and 2', 7' -dichlorofluorescein (DCFH) assays (Pietrogrande et al., 2019). However,

510 standardized experimental methods for evaluating OP have not been established (Song et al., 2021), and it is difficult to provide more consistent data on OP across samples at different locations and times. Moreover, each non-cellular OP assay is specific for ROS, making none of the methods used as a standard method for assessing the toxicity of environmental particles. Several studies have used a multi-measure approach to compensate for the specificity of a single probe response to ROS (Calas et al., 2018;

515 Puthussery et al., 2020; Yu et al., 2021; Xu et al., 2021). Xu et al. (2021) used three measurement methods ( $OP^{DTT}$ ,  $OP^{AA}$ , and  $OP^{GSH}$ ) to estimate the Canadian annual mean OP and found that the sensitivity of the three methods to different components varied widely. Choosing a variety of methods



for OP measurement can lead to more comprehensive results, but it can also lead to a significant increase in workload.

520 For this, we propose a hybrid approach combining existing observations of OP with the CTM. So, using OP from assays and their observed links to sources and chemical constituents can then be parameterized and implemented in CTM for a comprehensive assessment of OP exposure over large areas and time periods. The method considers the seasonal characteristics of the chemical composition of  $PM_{2.5}$  and the DTT activity measurement of  $PM_{2.5}$ . A positive matrix factorization (PMF) model and multiple linear regression (MLR) model were used to quantify the contribution of  $PM_{2.5}$  emission sources to OP (the volume-normalized DTT activity,  $DTT_v$ ). The normalized regression equation in this study provides the sensitivity of OP to each identified source. The advantage is that directly applying predicted and readily available  $PM_{2.5}$  data makes it easy to estimate the OP and assess health risks over large regions and across time and space. This approach enables the exploration of spatial and seasonal variations in aerosol OP

530 across China, providing insight into the contribution of sources, atmospheric processes, and meteorological conditions. There are some main limitations to this study that may lead to uncertainty in predicting OP outcomes. Firstly, the OP prediction considered is incomplete and does not include all sources of OP. For example, the transportation sector refers only to road transport, excluding emissions from ships and other mobile sources. And the transportation sector only considers tailpipe emissions in traffic. These lead to some OP uncertainties. Additionally, this study only considers the intrinsic OP of

535 total SOA due to limited long-term measurements of SOA. Different types of SOA may exhibit varied OP responses due to differences in their sources, formation pathways, and chemical compositions. Aging and fresh SOA may also exhibit varying toxicities (Liu et al., 2023). In future research, efforts should be made to comprehensively collect  $PM_{2.5}$  samples from various sources and fully explore the potential

540 relationships between OP and  $PM_{2.5}$  components/sources to further improve OP prediction models and reduce prediction uncertainties. Secondly, the OP prediction model adopted in this paper is based on Liu et al. (2018). The data samples are from DTT experimental measurements conducted in various coastal cities and from different emission sources, with limited data samples. In this study, it is used to predict the OP of cities across the country, which inevitably leads to a slight error in the forecast results. However,

545 due to the spatiotemporal and emission source differences of the data samples were considered, the three cities selected are representative, which reduces the errors caused by the data samples to a certain extent. Thirdly, a significant body of literature also indicates that vertical resolution can reflect atmospheric

thermodynamic environments and the evolution processes of mesoscale systems, which are related to the diffusion and transport of  $PM_{2.5}$ . Insufficient vertical resolution can hinder the accurate prediction of  $PM_{2.5}$  surface concentrations (HARA, 2011; Li et al., 2022; Li et al., 2023). Therefore, the configuration of the DEHM model (e.g., vertical resolution) also introduces uncertainties to this study. In conclusion, the results calculated by the method proposed in this study are compared with the existing measurement data (Liu et al., 2014; Liu et al., 2018; Wang et al., 2019), and a good agreement is observed. For instance, through DTT measurements, Zhang et al. (2023) reported an average  $OP^{DTTV}$  of  $1.33 \text{ nmol min}^{-1} \text{ m}^{-3}$  from January 2020 to June 2021 in downtown Nanjing, located in the YRD region of China, with a range of  $0.82\sim 2.08 \text{ nmol min}^{-1} \text{ m}^{-3}$ . This is close to our estimated results (Liu et al., 2024) for the YRD region, where the annual mean OP during 2010-2014 was  $1.56 \text{ nmol min}^{-1} \text{ m}^{-3}$ , and the annual mean OP for 2020 under two emission reduction scenarios were  $1.36 \text{ nmol min}^{-1} \text{ m}^{-3}$  and  $1.25 \text{ nmol min}^{-1} \text{ m}^{-3}$ , respectively. The relative errors for the two scenarios were 2.3% and 6.0%. Another study (Liu et al., 2020) investigating the OP of  $PM_{2.5}$  in Wuhan, located in the CC region, reported a mean  $OP^{DTTV}$  of  $1.8 \text{ nmol min}^{-1} \text{ m}^{-3}$  for the summer of 2012 in downtown Wuhan. This aligns closely with our estimated results (Liu et al., 2024) for the CC region ( $1.73 \text{ nmol min}^{-1} \text{ m}^{-3}$ ), with a relative error of 3.9%. Therefore, the method proposed in this study is reliable. The proposed method provides a possibility to solve the difficulty and high cost of OP measurement.

#### 4. Conclusions

This study established a spatial modelling for  $PM_{2.5}$  concentrations and OP, provided a method for calculating OP across China, and quantitatively assessed the impacts of meteorological conditions and anthropogenic emissions on  $PM_{2.5}$  and OP variability and levels in China. The following conclusions can be obtained:

- $PM_{2.5}$  and OP exhibited spatial clustering characteristics, with higher values mainly located in the central and eastern urban areas. About 85% and 40% of the areas had  $PM_{2.5}$  annual average concentrations exceeding the first-grade concentrations limit ( $15 \mu\text{g m}^{-3}$ ) and second-grade concentrations limit ( $35 \mu\text{g m}^{-3}$ ), respectively. Additionally, about 36% of the areas had OP concentrations lower than  $1 \text{ nmol min}^{-1} \text{ m}^{-3}$ , while 23% of the areas had OP concentrations higher than  $2 \text{ nmol min}^{-1} \text{ m}^{-3}$ .

- Variability in both  $PM_{2.5}$  and OP are influenced by a combination of meteorological conditions and emission inventories. Meteorological conditions contributed about 46% of  $PM_{2.5}$  variation and 65% of OP variation. The emission inventory contributed about 54% of the change in  $PM_{2.5}$  and about 35% of the change in OP.

- 580
- The percentage contributions of secondary aerosol formation, biomass burning, industry, coal combustion for residential heating, and traffic to  $PM_{2.5}$  were about 48%, 21%, 21%, 6%, and 4%, respectively. The percentage contributions of secondary aerosol formation, biomass burning, coal combustion for residential heating, industry, and traffic to OP were approximately 58%, 21%, 11%, 9%, and 1%, respectively.

585 A main finding of this study is that meteorological variability is the prime driver of OP variability, and not emissions. Furthermore, secondary aerosol formation and biomass burning are the main sources of OP. Thus, air pollution strategies should focus more on biomass burning and the emissions of the precursors taking part in the secondary aerosol formation, and it would be efficient to introduce special emissions controls during stagnation or other periods where OP accumulates.

590

**Author contributions**

Jiemei Liu performed the simulation and its validation, data analysis, investigation, and writing – original draft. Jesper H. Christensen conducted investigation, worked on methodology, and performed validation.

595 Zhuyun Ye conducted investigation, data analysis, worked on methodology. Shikui Dong provided the suggestion of data analysis, and manuscript feedback. Camilla Geels conducted investigation and provided the suggestion of simulation. Jørgen Brandt conducted investigation. Athanasios Nenes conducted investigation and provided manuscript feedback. Yuan Yuan provided manuscript feedback, supervision, provided acquired funding. Ulas Im provided the resources, supervised this work, provided

600 manuscript feedback, and managed the project administration. All authors discussed the results and commented on the manuscript.

**Competing interests**

The contact author has declared that none of the authors has any competing interests.

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610 101095457).

**Declaration of conflicting interests:**

The authors declare no conflicts of interests.

**Supplement:**

Density scatterplots of model performance and validation based on monthly mean observations (Figure

615 S1) is provided in the Supplementary Materials file.

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