

# Deciphering Isoprene Variability Across Dozen of Chinese and Overseas Cities Using Deep Transfer Learning

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30 **Key Points:**

31     • An explainable deep transfer learning framework was developed to predict  
32         isoprene concentrations and their variations.

33     • Different drivers accounted for historical trends of isoprene concentrations in  
34         Hong Kong and London from 1990 to 2023.

35     • Reducing nitrogen oxides would alleviate ozone pollution driven by rising  
36         temperatures and isoprene levels in the warming climate.

37

38 **Abstract**

39 Isoprene, the globally most abundant volatile organic compound, significantly impacts  
40 air quality. Determining isoprene concentration variations and their drivers is a  
41 persistent challenge. Here, we developed a robust machine learning framework to  
42 simulate isoprene concentrations, without requiring localized emission inventories and  
43 explicit chemistry. Temperature, radiation, and surface pressure were the primary  
44 drivers of short-term isoprene variations across Chinese cities. On climatic timescales,  
45 urban greenspace expansion and climate warming drove isoprene increases by 341 pptv  
46 in Hong Kong during 1990–2023, but traffic emission reductions in London  
47 counteracted the isoprene rise that climate warming would have otherwise caused (-755  
48 pptv vs. +31 pptv). Driven by rising temperatures and isoprene levels, ozone would  
49 increase by up to 1.7-fold by 2100 under the high-emission scenario. However,  
50 ambitious reduction in nitrogen oxides would alleviate this growth to 1.2-fold. The  
51 study has the potential to inform air quality management in a warming climate.

52

53 **1 Introduction**

54 Isoprene is the most abundant non-methane volatile organic compound (VOC) globally,  
55 with the total emissions reaching approximately 500 TgC per year, exceeding those of  
56 the total anthropogenic VOCs (Guenther et al., 2012; Huang et al., 2017). The high  
57 atmospheric reactivity makes it a key precursor for tropospheric ozone ( $O_3$ ) and  
58 secondary organic aerosol, both of which significantly impact air quality and climate  
59 (Paulot et al., 2012; Lin et al., 2013; [Xi et al., 2025a](#)). In particular, the effect is  
60 pronounced in urban environments due to the interactions between isoprene and  
61 anthropogenic emissions (Xu et al., 2015).

62 Terrestrial vegetation is the primary source of atmospheric isoprene, and the emissions  
63 are influenced by plant species, geographical locations, and environmental conditions  
64 (Guenther et al., 1994; Guenther et al., 1993). Urban landscapes show remarkable  
65 diversity in isoprene production, exemplified by stark differences between tree species.  
66 While urban greenspace offers numerous benefits, it also emerges as a notable  
67 contributor to urban isoprene (Ma et al., 2022). The emissions are highly sensitive to  
68 meteorological conditions (Wang et al., 2024a). The combination of climate warming  
69 and urbanization lead to intensified urban heat, which in turn boosts isoprene emissions  
70 from greenspace (Li et al., 2024; Pfannerstill et al., 2024). In addition, studies have  
71 shown that a portion of urban isoprene may stem from motor vehicles, the contribution  
72 of which varies by location and season (Borbon et al., 2001). However, vehicle  
73 emissions of isoprene do not necessarily increase with growing vehicle population, due  
74 to stringent emission controls in many cities. This further complicates the task of  
75 accurately simulating the concentrations and trends of urban isoprene. While isoprene  
76 may also be emitted from other anthropogenic sources, such as petrochemical activities  
77 and coal combustion, the amounts are generally small compared to biogenic emissions,  
78 especially in warm seasons (Peron et al., 2024).

79 Modeling and measurement deficiencies remain a serious concern in isoprene research  
80 across multiple disciplines. For example, the Model of Emissions of Gases and  
81 Aerosols from Nature (MEGAN) estimates vegetation emissions based on theoretical

relationships with meteorology and vegetation dynamics. This model significantly underestimates isoprene emissions from urban greenspace when it is driven by coarse resolution (e.g., >30 m) satellite-derived vegetation data (Ma et al., 2019; Ma et al., 2022). It is also difficult for current chemical transport models to accurately simulate isoprene concentrations, mainly resulting from the grid resolution and uncertainties in isoprene emissions, vertical dispersion rates and oxidation parameterization schemes (Arneth et al., 2011; Guenther et al., 2012). Local vegetation surveys and emission factor measurements can be made to improve model performance. However, the work is challenging and the outcomes often point out additional uncertainties (Seco et al., 2022; Wang et al., 2024b). While isoprene measurements have demonstrated reliability in atmospheric chemistry research, the temporal and spatial coverage remains suboptimal. Given these constraints, there are insufficient robust isoprene data available over climatic timescales (e.g., several decades) to reveal the drivers of long-term trends.

To confront this dilemma, we developed a generalized physics-informed neural network based on a residual Multi-Layer Perceptron with a transfer training strategy to reproduce/predict ambient isoprene concentrations. The model was trained by a comprehensive set of isoprene data observed at ten sites in China (a total of ~65,000 hourly data) and validated by a total of ~8,500 hourly and daily data at six overseas sites (Table S1). The model was verified for its ability to predict isoprene with limited sizes of observational data and understand intricate relationships between isoprene and influencing factors. The model was then used to predict future trends of isoprene and the resulting O<sub>3</sub> variations in different climate scenarios. This study enhances our understanding of the responses of ambient isoprene concentrations to emissions and meteorology, and has the potential to inform urban planning and air quality policies in the warming climate.

107

108 **2 Data and Models**

109 **2.1 Isoprene Data and Deep Learning Model**

110 A total of over 72,000 hourly (and a small fraction of daily) data of isoprene  
111 concentrations in the daytime (06:00–20:00 local time) of warm seasons (May–October)  
112 were compiled from 16 sites worldwide. Around 88% the data was from different parts  
113 of China, and the remainder was from North America, Amazonia, India, and the UK  
114 (see Table S1). To ensure comparability, we included only online measurements,  
115 excluding offline sampling and analysis methods. While inter-instrument bias might  
116 exist, the isoprene variability within each site was expected to be much larger than any  
117 plausible inter-instrument bias. Moreover, this will not influence the analysis of  
118 isoprene variations at individual sites.

119 The residual multi-layer perceptron architecture (ResMLP) was employed to  
120 approximate the complicated responses of isoprene concentrations to input features,  
121 which was coupled with a physics-informed neural network, thereby PINN-ResMLP.  
122 This approach integrated domain knowledge by enforcing monotonicity constraints  
123 between isoprene and its major sources (e.g., vegetation and traffic emissions), thus  
124 ensuring physically consistent predictions. These constraints were implemented  
125 directly in the model’s loss function, which combined terms for data fitting,  
126 monotonicity regularization, and network structure penalties. As a fully data-driven  
127 model, ResMLP may learn patterns that are inconsistent with physical laws (Feng et al.,  
128 2025). Incorporating expert knowledge and physical constraints into the model can  
129 guide the learning processes (Zhu et al., 2024). In this study, we stipulated that isoprene  
130 concentrations were positively correlated with the biogenic and traffic emission sources.  
131 This relationship therefore can be expressed as:

$$\frac{\partial ISOP}{\partial VI} > 0 \quad (1)$$

$$\frac{\partial ISOP}{\partial BC_{traffic}} > 0 \quad (2)$$

134 where ISOP represents isoprene concentrations; VI is vegetation index derived from  
 135 Leaf Area Index (LAI) and Normalized Difference Vegetation Index (NDVI) (see Text  
 136 S1); and  $BC_{\text{traffic}}$  is traffic emissions of black carbon. To satisfy this priori knowledge,  
 137 we developed PINN-ResMLP to constrain the model. The optimization objective of  
 138 PINN-ResMLP included data item loss ( $\mathcal{L}_{\text{data}}$ ), physical inconsistency loss  
 139 ( $\mathcal{L}_{\text{monotonicity}}$ ), and additional structural loss ( $\mathcal{L}_{\text{structure}}$ ). Meanwhile, the L2 norm of  
 140 the network parameters, namely adding the sum of the squares of all network weights  
 141 (parameters) to the loss function, could effectively regularize and prevent overfitting in  
 142 PINN-ResMLP. Finally, the total function was formulated as:

$$143 \quad \mathcal{L} = \mathcal{L}_{\text{data}} + \alpha \cdot \mathcal{L}_{\text{monotonicity}} + \beta \cdot \mathcal{L}_{\text{structure}} \quad (3)$$

$$144 \quad \mathcal{L}_{\text{data}} = \frac{1}{2N} \sum_{i=1}^N \left[ (ISOP_{\text{obs},i} - ISOP_{\text{pred},i})^2 + |ISOP_{\text{obs},i} - ISOP_{\text{pred},i}| \right] \quad (4)$$

$$145 \quad \mathcal{L}_{\text{monotonicity}} = \frac{1}{N} \sum_{i=1}^N \left[ \textcolor{red}{1} - \frac{\text{sign}\left(\frac{\partial ISOP}{\partial VI}\right) + \text{sign}\left(\frac{\partial ISOP}{\partial BC_{\text{traffic}}}\right)}{2} \right] \quad (5)$$

$$146 \quad \text{sign}(\theta) = \begin{cases} -1, & \theta < 0 \\ \textcolor{red}{01}, & \theta \geq 0 \end{cases} \quad (6)$$

$$147 \quad \mathcal{L}_{\text{structure}} = \sum_{i=1}^M (W_i^2 + b_i^2) \quad (7)$$

148 where the  $\alpha$  and  $\beta$  are trade-off parameters;  $N$  is the number of training samples;  $i$   
 149 represents a certain sample;  $ISOP_{\text{obs}}$  and  $ISOP_{\text{pred}}$  are observed and predicted  
 150 isoprene concentrations, respectively;  $M$  is the number of layers in PINN-ResMLP.

151 Predictor variables were selected to capture key sources and sinks of isoprene, including  
 152 VI (see Text S1), meteorological parameters (e.g., temperature, solar radiation), and  
 153 black carbon emitted from traffic ( $BC_{\text{traffic}}$ ) as a proxy for anthropogenic emissions. **To**  
 154 **ensure consistency among all variables, datasets with different spatial resolutions (e.g.,**  
 155 **0.1° and 0.25°) were resampled to achieve a uniform resolution of 0.1° using bilinear**  
 156 **interpolation.** Full variable definitions and sources are provided in Table S2.

157 To address data scarcity at some sites, we implemented a supervised transfer learning  
 158 strategy. The PINN-ResMLP was pre-trained on data-rich sites and fine-tuned with

159 limited data from target sites (see Table S3). Three training strategies were adopted: (1)  
 160 transfer learning (T), where models were pre-trained on data from other sites and fine-  
 161 tuned on the target site; (2) no-transfer (NT), where models were trained solely on target  
 162 site data; and (3) mixed training (MIX) using combined data from all sites. Model  
 163 performance was evaluated using four-fold cross-validation (Table S3) and metrics  
 164 including normalized mean absolute error (NMAE) and coefficient of determination  
 165 ( $R^2$ ). Comparisons were made against standard machine learning algorithms, such as  
 166 Random Forest (RF), extreme gradient boosting (XGB), and support vector machine  
 167 (SVM). All algorithms were optimized using extensive grid search (see  
 168 hyperparameters in Table S4). NMAE and  $R^2$  are calculated as follows:

$$169 \quad NMAE = \frac{\frac{1}{N} \sum_{i=1}^N |ISOP_{obs,i} - ISOP_{pred,i}|}{\overline{ISOP}_{obs}} \quad (8)$$

$$170 \quad R^2 = 1 - \frac{\sum_{i=1}^N (ISOP_{obs,i} - ISOP_{pred,i})^2}{\sum_{i=1}^N (ISOP_{obs,i} - \overline{ISOP}_{obs})^2} \quad (9)$$

171 where  $\overline{ISOP}_{obs}$  represents the mean value of  $ISOP_{obs}$ .  
 172 Finally, the Shapley Additive Explanations (SHAP) approach (Lundberg et al., 2020)  
 173 was used to quantify the contributions of input variables to model predictions. SHAP  
 174 values allow us to assess the impact of each factor on isoprene concentrations. The  
 175 calculation is defined as follows:

$$176 \quad \varphi_i = \sum_{S \subseteq K \setminus \{i\}} \frac{|S|! (K - |S| - 1)!}{|K|!} [f(S \cup \{i\}) - f(S)] \quad (10)$$

177 where  $K$  represents the set of all features,  $S \subseteq K \setminus \{i\}$  denotes a feature subset that  
 178 excludes feature  $i$ ,  $|S|$  is the size of subset  $S$ ,  $f(S)$  is the model's prediction under  
 179 feature subset  $S$ , and  $f(S \cup \{i\}) - f(S)$  is the marginal contribution of feature  $i$ .

## 180 2.2 Attribution of Long-term Isoprene Trends and O<sub>3</sub> Projections

181 The PINN-ResMLP model was also used to quantify the contributions of different  
 182 factors to the long-term trends of isoprene concentrations at three sites in Hong Kong  
 183 and London using a scenario-based approach. Using the historical data of

184 meteorological parameters, VI and  $BC_{traffic}$  as the input of the PINN-ResMLPT model,  
185 we predicted the summertime (June to August) isoprene concentrations in Hong Kong  
186 and London for the period of 1990–2023 (base scenario). In order to reveal the impacts  
187 of the major drivers on the isoprene variations, we also predicted the isoprene  
188 concentrations by fixing the temperature, VI and  $BC_{traffic}$  as their averages over the  
189 above period one by one (controlled scenarios). The differences in the predicted  
190 isoprene ( $isoprene_{diff}$ ) between the base and controlled scenarios depicted the isoprene  
191 trends induced by the individual factors. The  $isoprene_{diff}$  was then compared between  
192 different time periods, e.g., the first and last 17 years and the first, middle and last  
193 decades. Besides, the coefficient of variation (CV) was calculated for the predicted  
194 isoprene concentrations in all the scenarios over the period of 1990–2023. The CV  
195 differences between the base and controlled scenarios indicate how the changes in  
196 temperature, VI and traffic emissions (represented by  $BC_{traffic}$ ) increased or decreased  
197 the variations in isoprene concentrations.

198 Additionally, future isoprene concentrations at the Hong Kong site were projected for  
199 2030–2100 based on temperature changes under different climate scenarios developed  
200 by the Coupled Model Intercomparison Project Phase 6 (CMIP6), while keeping other  
201 influencing factors constant. Briefly, we used the temperature data from four Shared  
202 Socioeconomic Pathways (SSPs), including SSP126 (low forcing), SSP245  
203 (intermediate forcing), SSP370 (medium-high forcing), and SSP585 (high forcing), and  
204 held the other influencing factors constant. The averages of outputs from four Coupled  
205 Model Intercomparison Project Phase 6 (CMIP6) models (ACCESS-CM2, CMCC-  
206 ESM2, MPI-ESM1-2-HR, and GFDL-ESM4) were adopted (Xu et al., 2024).

207 Using the future profiles of temperature and isoprene as constraints, we simulated  $O_3$   
208 concentrations as a function of isoprene and temperature using a zero-dimensional box  
209 model under different  $NO_x$  reduction scenarios. The Framework for 0-D Atmospheric  
210 Modeling (F0AM) incorporating Master Chemical Mechanism v3.3.1 was used to  
211 simulate  $O_3$  under different sets of temperatures and isoprene concentrations (Lyu et al.,  
212 2024). The model was constrained by the average diurnal profiles of air pollutants

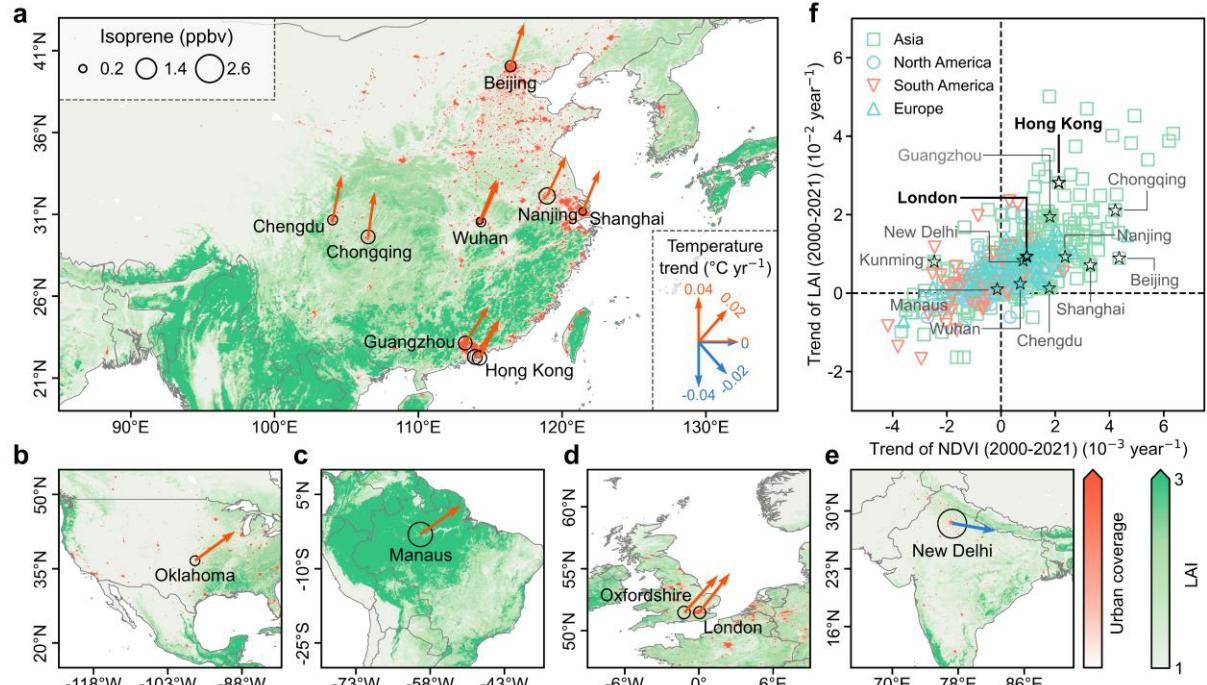
213 (excluding O<sub>3</sub>) and meteorological parameters observed in the summer of 2023 at the  
214 Hong Kong\_TC site, except that the daytime average temperature changed from 22 °C  
215 to 38 °C in 2 °C intervals and the daytime average isoprene concentrations varied in the  
216 range of 0.15–1.8 ppbv in intervals of 0.15 ppbv. The O<sub>3</sub> isopleths were depicted using  
217 the simulation results for 108 temperature-isoprene settings. Additionally, the above  
218 simulations were repeated in different scenarios of NO<sub>x</sub> reduction, i.e., 49.7% and 82.6%  
219 under the SSP370 and SSP126, respectively (Lou et al., 2023; Rogelj et al., 2018). It is  
220 worth noting that the diurnal profiles of other O<sub>3</sub> precursors, such as VOCs and carbon  
221 monoxide, were kept unchanged throughout all the simulations. Meanwhile, our future  
222 projections are designed to isolate the chemical response of O<sub>3</sub> to changes in  
223 temperature and isoprene and do not explicitly incorporate potential future changes in  
224 greenspace, urban morphology, or other anthropogenic emissions. Although these  
225 factors are expected to evolve under urban development and climate mitigation  
226 pathways, the present analysis focuses on quantifying the impacts of climate warming  
227 on isoprene emissions and the consequent O<sub>3</sub> responses.

228

229 **3 Results and Discussion**

230 **3.1 Simulating Isoprene Concentrations Using PINN-ResMLPT Model**

231 The isoprene concentrations averaged over the respective observation periods varied  
232 significantly from 0.15 ppbv in Wuhan to 2.79 ppbv in New Delhi (Figure 1 and Figure  
233 S1), due to the differences in sampling periods, climatic conditions, and vegetation type  
234 and density. We noticed that most cities have experienced an increase in greenspace in  
235 the last 20 years, and there existed significant differences in greenspace coverage and  
236 its recent trends between the cities (e.g., Hong Kong versus London). The high  
237 vegetation cover appeared to explain the elevated levels of isoprene in South China and  
238 Amazonia. Temperature also had a strong effect on isoprene concentrations, as  
239 indicated by the Pearson correlation coefficient ( $R$ ) between hourly isoprene and  
240 temperature at individual sites, i.e., 0.41–0.72. The temperature and temperature  
241 variation were spatially non-uniform, implying its inconsistent roles in affecting  
242 biogenic isoprene emissions. Additionally, anthropogenic emissions might have made  
243 significant contributions to isoprene in New Delhi, given the high nocturnal levels and  
244 the peak in evening rush hours (Figure S2). This was also mentioned in a previous study  
245 (Tripathi et al., 2022). The Weather Research and Forecasting model with Chemistry,  
246 which theoretically takes these factors into account, was used to simulate the isoprene  
247 concentrations. However, substantial divergences were noted between the simulated  
248 and observed values at hourly or even daily resolution (Figure S3), demonstrating the  
249 challenge for chemical transport models in accurately simulating isoprene (Morichetti  
250 et al., 2022; Wang et al., 2024b).

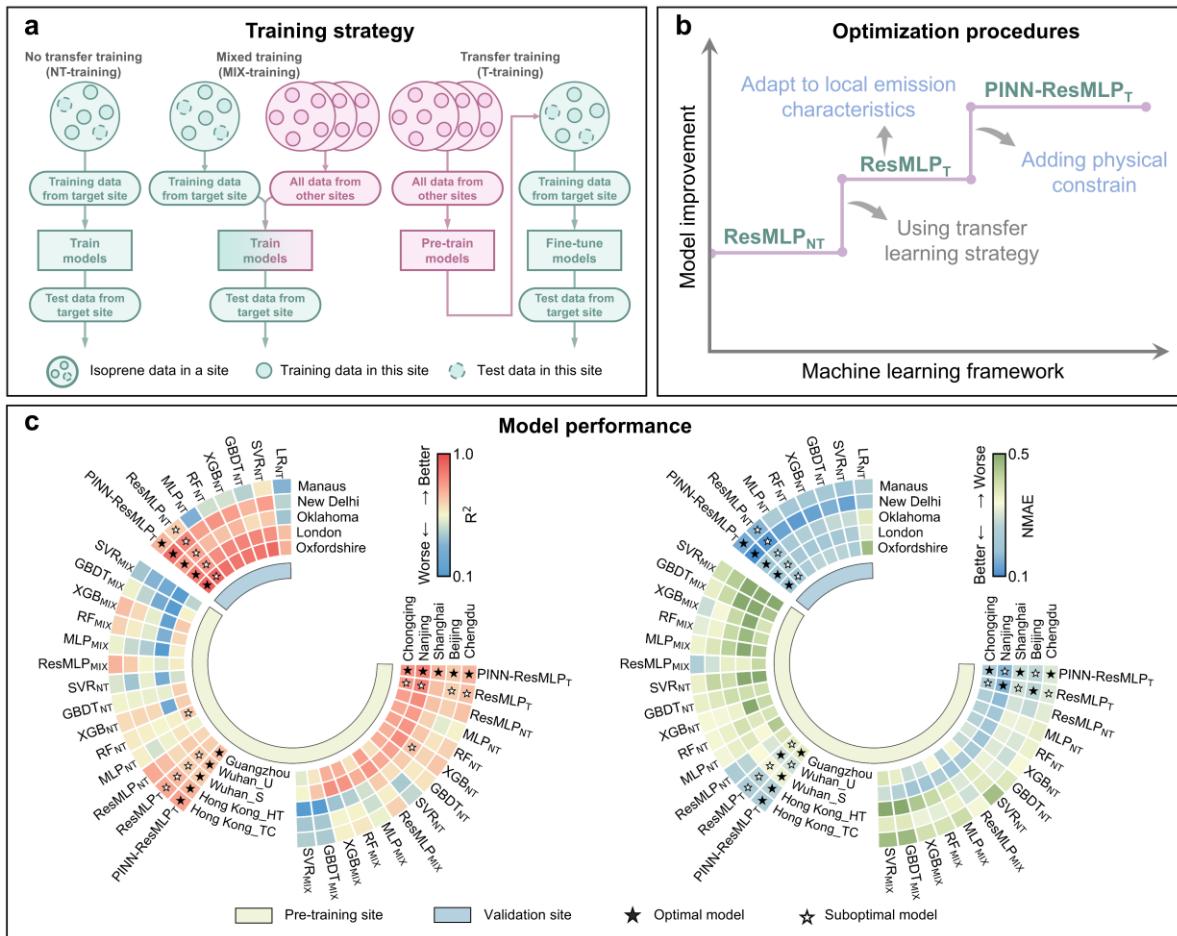


251 **Figure 1.** Geographical distribution of the isoprene sampling sites. (a-e) Locations of isoprene  
252 measurement sites in China (a), North America (b), Amazonia (c), United Kingdom (d), and  
253 India (e). The direction of arrows represents the trend of temperature from 1990 to 2023, and  
254 the size of the circle is proportional to isoprene concentration. (f) Trends of LAI (Leaf Area  
255 Index) and NDVI (Normalized Difference Vegetation Index) from 2001 to 2021 in major cities  
256 around the world.

258 We then examined the isoprene prediction ability of various machine learning  
259 algorithms with three training strategies: T, NT, and MIX (see Section 2.3). Overall,  
260 the model utilizing the NT training strategy exhibited higher fitting accuracy than the  
261 one employing the MIX training strategy (Figure 2). This suggests that training with  
262 data from different sites might introduce additional noises, due to the differences in  
263 isoprene emission dynamics. Particularly, isoprene emissions are highly sensitive to  
264 local vegetation profiles. While the ResMLP model with the NT training strategy  
265 (ResMLP<sub>NT</sub>) performed moderately among all the algorithms, the performance was  
266 improved by incorporating the T strategy. Specifically, the ResMLP<sub>T</sub> outperformed the  
267 other algorithms at 8 out of 10 sites, with the decrease in NMAE of 1%–5% and increase  
268 in  $R^2$  of 0.01–0.07. The results indicated that the ResMLP<sub>T</sub> model effectively exploited  
269 the implicit prior knowledge from the pre-training data to guide isoprene prediction at

270 the target sites. Importantly, the pre-trained parameters were fine-tuned using limited  
271 sizes of local data, which adapted the model to local isoprene emission dynamics  
272 without requiring region-specific vegetation profiles, such as vegetation types and  
273 corresponding emission factors. It is worth noting that the size of the retraining data at  
274 the validation sites was ~30% of all the data. Thus, the model's good performance at  
275 the validation sites demonstrated its utility in data-scarce regions. Furthermore, with  
276 the incorporation of PINN, the PINN-ResMLP<sub>T</sub> showed a better understanding of the  
277 real target-feature relationships with more interpretable prediction results. The model  
278 performance was further improved, as indicated by the highest R<sup>2</sup> values across all the  
279 sites (Figure 2).

280 Next, we also validated the PINN-ResMLP<sub>T</sub> model by applying it to predict isoprene  
281 concentrations at the overseas sites (Figure 1). The model was pre-trained with the  
282 complete dataset from all the sites in China, which was further fine-tuned with 70% of  
283 the data at the individual target sites and validated with the remaining data. Compared  
284 to the suboptimal model, the PINN-ResMLP<sub>T</sub> model significantly improved the  
285 prediction of isoprene, especially at New Delhi and Manaus, with the increase in R<sup>2</sup>  
286 (decrease by NMAE) by 0.07 (25%) and 0.08 (13%), respectively (Figure 2). This  
287 demonstrated the model's broad applicability. Moreover, the model outperforms many  
288 other methods in predicting isoprene. For instance, the root mean square error of the  
289 PINN-ResMLP<sub>T</sub> at Manaus was 0.17 ppbv, compared to 0.95 ppbv for the early attempt  
290 in Cross-track Infrared Sounder retrieval (Fu et al., 2019). This superior performance  
291 establishes the PINN-ResMLP<sub>T</sub> as our best choice. In fact, we would expect more  
292 accurate predictions if the model were pre-trained by a wider range of field  
293 measurement data from various regions.

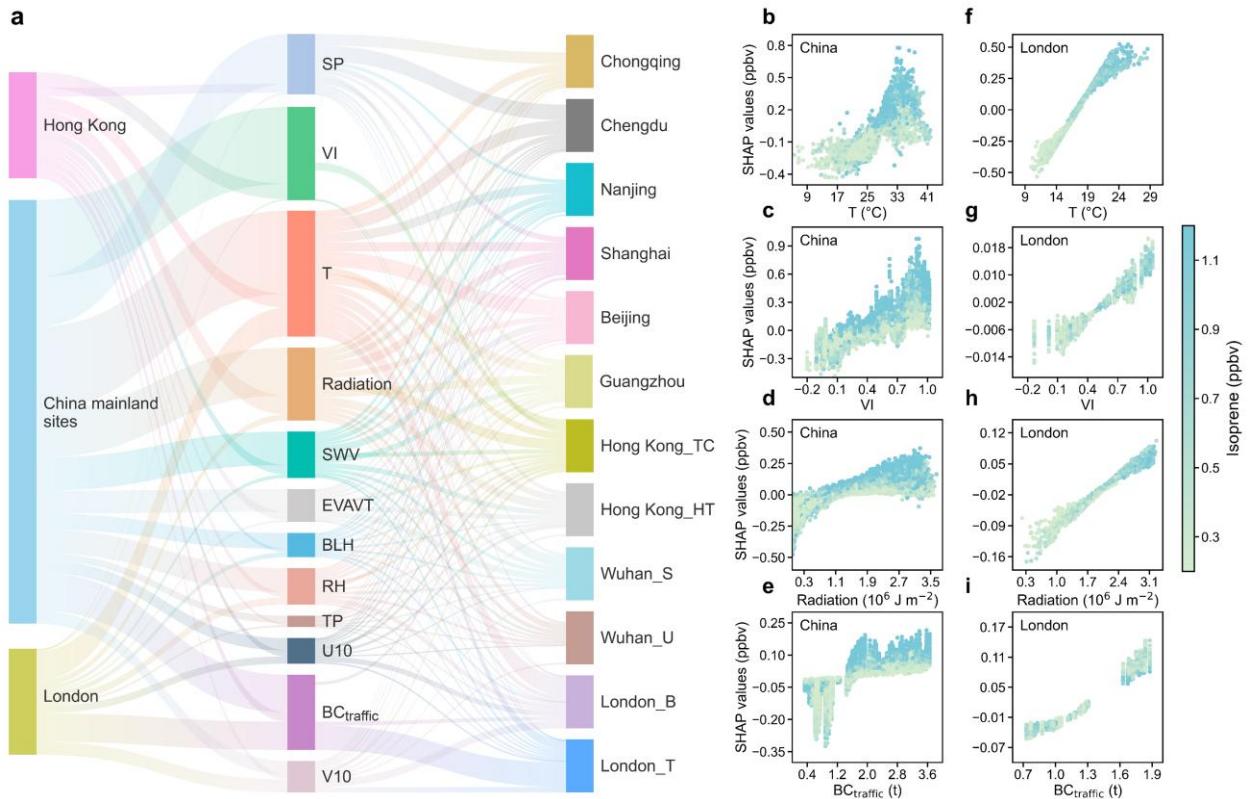


302 **3.2 Main Factors Influencing Urban Isoprene Concentrations**

303 Further, a feature importance method based on the ~~the SHapley Additive exPlanations~~  
304 (~~SHAP~~) values was employed to explore the prediction results (Figure 3). While we  
305 prefer to present it for individual sites, the feature importance of VI and BC<sub>traffic</sub> was  
306 not calculated for the Chinese sites except a suburban site in Hong Kong (HK\_TC), due  
307 to the low temporal resolutions of VI and BC<sub>traffic</sub> data and the short isoprene  
308 observation periods. Here, we discuss the drivers of short-term (2-4 years) and long-  
309 term (over 10 years) isoprene concentration variations, separately.

310 With the VI and BC<sub>traffic</sub> remaining relatively stable in the short term, the model  
311 indicated that temperature, radiation, surface pressure, and soil water vapor were the  
312 most significant drivers of short-term isoprene variations, and their average relative  
313 importance was 18.8%, 11.9%, 11.3%, and 8.1% across the all the Mainland China sites,  
314 respectively. In addition, evaporation from vegetation transpiration and relative  
315 humidity also played significant roles in affecting isoprene concentrations at the Wuhan  
316 suburban site and Beijing urban site, respectively. The model also effectively captured  
317 the target-feature relationships. In China, the predicted isoprene concentrations  
318 increased with temperature below  $\sim 35^{\circ}\text{C}$ , above which a decline occurred at some sites.  
319 A typical example was the response in Chongqing with frequent occurrence of high  
320 temperature extremes (Figure S4). High temperatures suppress vegetation emissions  
321 due to a reduction in enzyme activity and substrate availability while accelerating the  
322 photochemical oxidation loss of isoprene. A similar pattern was observed in the  
323 response of isoprene concentrations to radiation in China, especially at extremely high  
324 levels. Such nonlinear responses are critical to the parameterization of isoprene  
325 emissions in numerical models. Notably, recent studies have revealed substantial  
326 uncertainties in the MEGAN model's performance under extreme heat conditions  
327 (Wang et al., 2024a). In contrast, our data-driven machine learning approach effectively  
328 captures these complex, nonlinear relationships between isoprene concentrations and  
329 environmental predictors, offering a promising pathway to refine and optimize  
330 parameterization schemes in chemical transport models. In addition to the close

331 relationships with temperature, solar radiation affects the hydroxyl radical  
 332 concentrations and therefore can significantly increase chemical loss of isoprene. In  
 333 contrast, these phenomena were not observed in London, because of lower temperatures  
 334 and weaker solar radiations. Overall, our transfer learning model reasonably reflected  
 335 the isoprene-meteorology relationships.



336 **Figure 3.** Modeling explainable results at each site based on SHAP value. **(a)** Feature  
 337 importance for isoprene concentrations at individual sites. **(b-i)** The SHAP dependence plots of  
 338 major influencing variables averaged at the Chinese sites **(b-e)** and London sites **(f-i)**.  
 339

340 Further, the long-term isoprene observations in London and Hong Kong offer an  
 341 opportunity to examine how the evolutions of VI and BC<sub>traffic</sub> affected isoprene  
 342 variations on a climatic scale. As two prominent international cities, London and Hong  
 343 Kong are characterized by distinct climatic zones: London experiences a mid-latitude  
 344 temperate maritime climate, whereas Hong Kong is influenced by a low-latitude  
 345 subtropical monsoon climate. This climatic differentiation is reflected in their  
 346 predominant vegetation types, with temperate deciduous trees being prevalent in

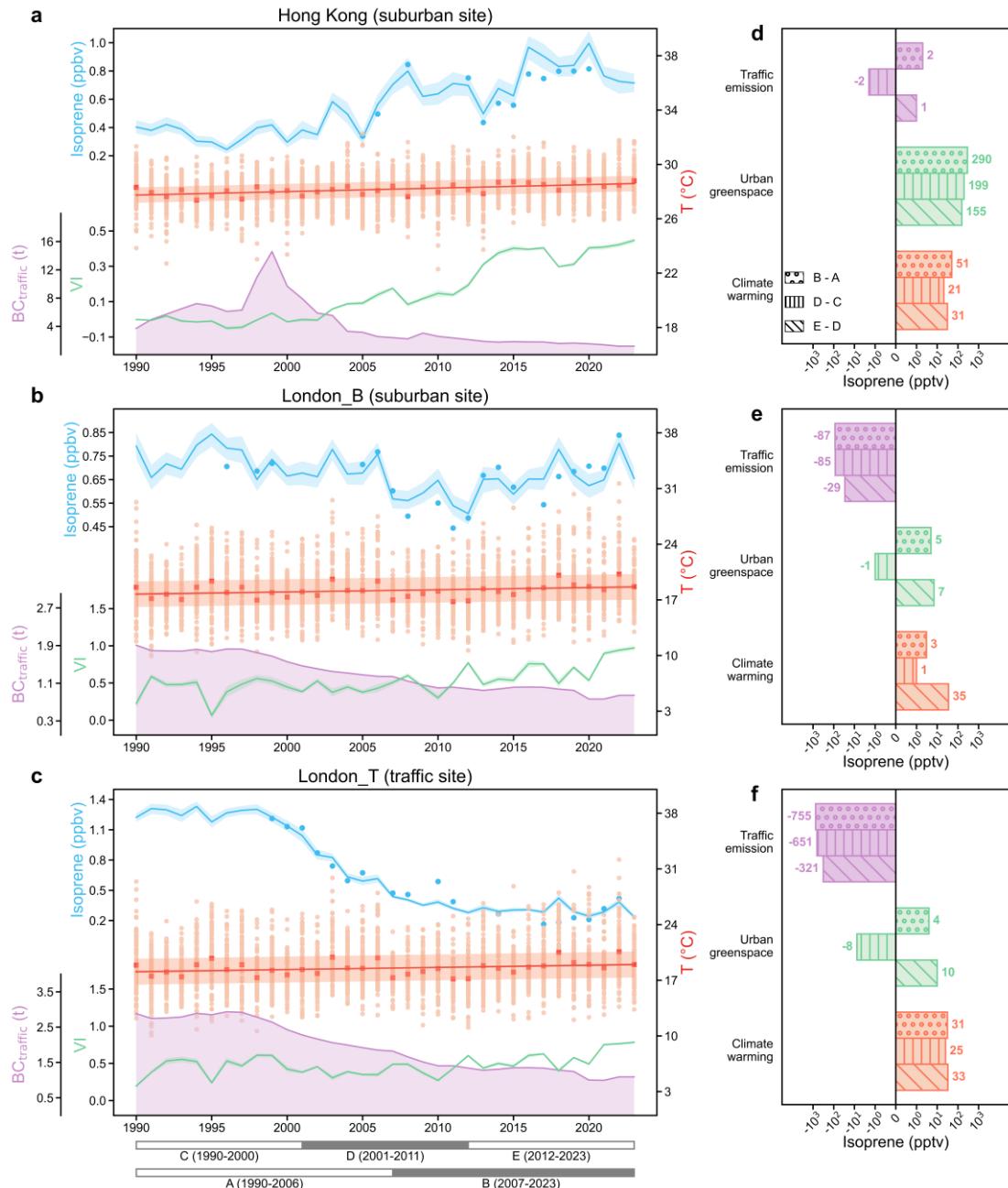
347 London and evergreen broad-leaved trees dominating the landscape in Hong Kong.  
348 Furthermore, the trajectories of urbanization and air pollution management have  
349 evolved differently in each city, shaped by their unique environmental and socio-  
350 economic contexts. Here, we focus on the results at two London sites and a Hong Kong  
351 site, where long term data was available. Radiation, VI and temperature were the most  
352 predominant influencing factors at the suburban site in Hong Kong. In contrast, the  
353 relative importance of VI was low at the two London sites. Over the period of 2000–  
354 2021, Hong Kong has experienced a notable increase in NDVI ( $2.1 \times 10^{-3}$  year $^{-1}$ ) and  
355 LAI ( $2.8 \times 10^{-2}$  year $^{-1}$ ), while the rate was much lower in London, i.e.,  $0.9 \times 10^{-3}$  year $^{-1}$   
356 for NDVI and  $0.9 \times 10^{-2}$  year $^{-1}$  for LAI. Additionally, the significant difference in VI  
357 importance between Hong Kong and London might also be attributed to the different  
358 strength of vegetation emissions across latitudes (Guenther et al., 2006; Guenther et al.,  
359 2012). Instead, BC<sub>traffic</sub> (temperature) ranked the first at the traffic (suburban) site in  
360 London, followed by other meteorological factors (Figure 3). While the meteorological  
361 impacts were not surprising, isoprene correlated well with the BC<sub>traffic</sub> emissions and  
362 observed benzene at the London traffic site (Figure S5), thereby the high relative  
363 importance of BC<sub>traffic</sub>. This is consistent with the previous studies on traffic emissions  
364 of isoprene in London (Borbon et al., 2001; Von Schneidemesser et al., 2011). As  
365 constrained using the PINN, the SHAP values for isoprene concentrations of VI and  
366 BC<sub>traffic</sub> showed a monotonic increasing trend.

367

368 **3.3 Factors Driving Long-term Trends of Isoprene**

369 The model was also used to build the time series data of daytime isoprene  
370 concentrations at a daily resolution over a climatically relevant period (1990–2023) at  
371 the three sites with long-term but incomplete isoprene data. The comparison between  
372 the geographically distinctive London and Hong Kong offers a rare opportunity to  
373 examine the different drivers of isoprene trends. As shown in Figure 4, the predicted  
374 isoprene concentrations were in good agreement with the observations, with the  $R^2$  of  
375 0.68–0.83 and NMAE of 21%–27%. It's worth noting that the observational data was  
376 missing for 50%–67% of the dates at the three sites. This underscored the model's  
377 effectiveness in retrieving historical isoprene concentrations from limited observation  
378 data.

379 Over the past 34 years, the isoprene concentrations at the Hong Kong site have shown  
380 an increasing trend with the rate of 18.1 pptv year<sup>-1</sup>, as have the temperature and VI. In  
381 contrast, traffic emissions have been significantly reduced since 1998, due to the  
382 effective human interventions. The trend of the predicted isoprene correlated strongly  
383 with the SHAP values of VI ( $R = 0.95$ ) and moderately with that of temperature ( $R =$   
384 0.63). By fixing the variables one by one, we determined the variations in factor  
385 contributions to isoprene concentrations, which were then compared between different  
386 time periods. It was found that urban greenspace emerged as the dominant factor  
387 impacting Hong Kong's isoprene levels, causing a rise in isoprene concentrations of  
388 290 pptv between the last 17 years and the first 17 years. Meanwhile, the contribution  
389 of climate warming was 51 pptv, while the traffic contribution was minor. Moreover,  
390 without changes in urban greenspace, the coefficient of variation (CV) of annual  
391 average isoprene concentrations would decrease by 70.5%, in comparison to the  
392 decrease of 12.0% and 6.0% in absence of changes in climate warming and traffic  
393 emissions, respectively. This reiterated the significant impacts of urban greenspace on  
394 the variations and trends of isoprene concentrations.



395

396 **Figure 4.** Long-term trends of the summertime isoprene and the drivers. (a-c) Variations of  
397 isoprene concentrations (blue lines for simulated, blue dots for observed), temperature (T),  
398 urban greenspace (VI) and traffic emissions (BC<sub>traffic</sub>) in Hong Kong (a) and London (b and c).  
399 The red dots and red line represent temperature and the fitted trend for the mean temperature,  
400 respectively. The bands represent 95% confidence intervals. (d-f) Changes in isoprene  
401 concentrations caused by climate warming, urban greenspace and traffic emissions in Hong  
402 Kong (d) and London (e and f) during different periods.

403 In contrast, the isoprene concentrations in London were lower in the last 17 years.  
404 Climate warming would have increased the isoprene concentrations by 31 pptv  
405 compared to those in the first 17 years at the traffic site, while the impact at the suburban  
406 site and the effects of urban greenspace at both sites were negligible. Interestingly,  
407 traffic emissions accounted for 87 pptv and 755 pptv of isoprene reduction at the  
408 suburban site and the traffic site, respectively. This was likely attributed to stringent  
409 traffic emission controls, as indicated by the significant downward trend of  $BC_{\text{traffic}}$ .  
410 The effect was more pronounced during the first two thirds of the study period (1990–  
411 2011). Specifically, the traffic-related isoprene reduction was 85 pptv from the first  
412 (1990–2000) to the second decade (2001–2011) at the suburban site, in comparison to  
413 the 29 pptv between the second and the last decade (2012–2023). Actually, the observed  
414 isoprene concentration correlated moderately ( $R = 0.60$ ) with  $BC_{\text{traffic}}$  from 1990 to  
415 2011 based on their annual averages. This suggests that traffic emission controls also  
416 affected isoprene concentrations even in non-urban areas. Despite the higher VI in  
417 London, the increasing rate ( $1.2 \text{ year}^{-1}$  at the suburban site and  $0.7 \text{ year}^{-1}$  at the traffic  
418 site) was lower than that in Hong Kong ( $1.6 \text{ year}^{-1}$ ). Additionally, the weak effects of  
419 urban greenspace might be also due to the relatively low emission strengths of high-  
420 latitude vegetation (Guenther et al., 2006). Moreover, the impact of climate warming  
421 became evident in the last decade (2012–2023) at the suburban site in London and,  
422 together with urban greenspace, reversed the isoprene reduction that would otherwise  
423 have been achieved by traffic emission controls. This aligned with the accelerated  
424 temperature rise from 2011 onwards (Figure S6), which was also reported elsewhere  
425 (Cao et al., 2021). From the perspective of variations in annual isoprene concentrations,  
426 the CV at the traffic site primarily resulted from changes in traffic emissions. At the  
427 suburban site, it would decrease by 32.4% and 14.0% if temperature and traffic  
428 emissions did not change.

429 Overall, our results demonstrate a tale of two cities: similarities and differences in  
430 drivers of long-term isoprene trends. Temperature-driven increases in isoprene  
431 concentrations were revealed in both cities, especially in the last decade, underscoring

432 the universal impacts of climate warming on vegetation emissions. However, the  
433 disparities in green space changes and probably different biogenic isoprene emission  
434 strengths between the two cities led to the different effects of VI on isoprene variations.  
435 Additionally, the isoprene variations over the 34 years have been more influenced by  
436 traffic emissions in London, although both cities have implemented stringent vehicle  
437 emission controls. While the reasons remain to be explored, we did not identify any  
438 correlation between the observed isoprene and  $BC_{\text{traffic}}$  (or benzene) in Hong Kong,  
439 even at a traffic site (Figure S5).

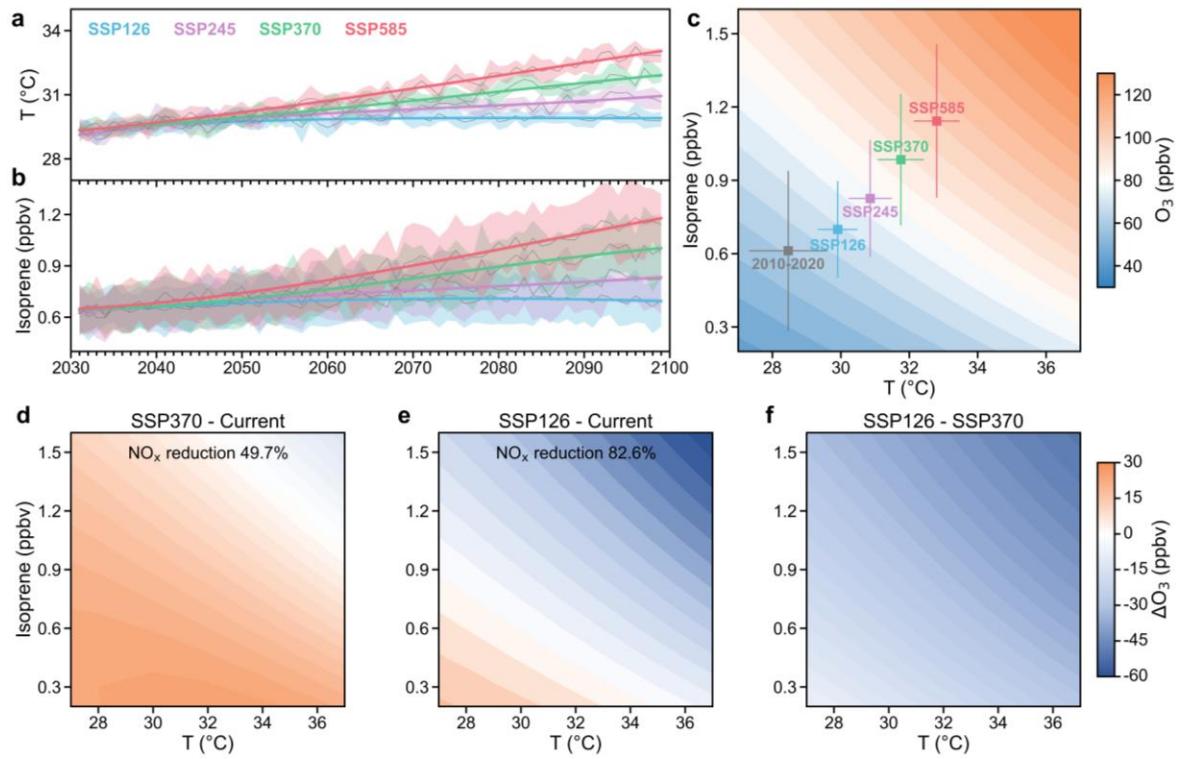
440

441 **3.4 Future Projections for Isoprene and O<sub>3</sub> in Hong Kong**

442 A significant issue associated with increasing isoprene levels in a warming climate is  
443 the potential for elevated ground-level O<sub>3</sub> pollution (Xi et al., 2025b; Wang et al.,  
444 2024c). We used the temperature from the latest CMIP6 multi-model ensemble  
445 forecasts to predict isoprene concentrations from 2030 to 2100 in Hong Kong under  
446 four IPCC's shared socio-economic pathway (SSP) scenarios, while the other factors  
447 were kept constant. As shown in Figure 5a, the temperature is expected to increase by  
448 0.71–3.60 °C from 2030 to 2100. The model indicated that the daytime average  
449 concentration of isoprene would increase by 87–530 pptv (15%–87%) by 2100 (Figure  
450 5b). The changes are on the same magnitude as the previous estimates that isoprene  
451 emissions will increase by 21%–57% by the end of this century relative to the 1990–  
452 2010 levels (Cao et al., 2021; Sanderson et al., 2003).

453 Further, we simulated the O<sub>3</sub>-isoprene-temperature relationships in Hong Kong (as an  
454 example) using future temperatures and isoprene concentrations while fixing the other  
455 air pollutants and meteorological conditions at present levels. The simulated O<sub>3</sub>  
456 increased markedly with the rise in temperature and isoprene concentrations (Figure  
457 5c-5f). The O<sub>3</sub> concentration would increase by up to 1.7 folds by 2100 under the  
458 SSP585 scenario of temperatures and isoprene. An increase in the combined risk of heat  
459 and O<sub>3</sub> exposure could be expected. To explore the approach of alleviating the adverse  
460 impact of O<sub>3</sub>-isoprene-temperature synergy, we proposed additional scenarios by  
461 cutting anthropogenic NO<sub>x</sub> emissions. With the NO<sub>x</sub> reduction from the current to  
462 different SSPs levels, the O<sub>3</sub> concentrations would increase and decrease under low and  
463 high isoprene-temperature conditions, respectively (Figure 5d-5f). This inconsistent  
464 variation is due to the evolution of O<sub>3</sub> formation regime with the rising temperatures  
465 and isoprene. It is worth noting that more ambitious NO<sub>x</sub> reduction would result in  
466 greater O<sub>3</sub> benefits. For example, O<sub>3</sub> would decrease in a much wider range of  
467 temperature and isoprene when NO<sub>x</sub> is reduced under SSP126. The O<sub>3</sub> growth by 2100  
468 would be only 1.2 folds in the SSP585 scenario of temperatures and isoprene. Therefore,

469 substantial reduction in anthropogenic  $\text{NO}_x$  would effectively address the synergy  
 470 between temperature, isoprene and  $\text{O}_3$ .



471  
 472 **Figure 5.** Projected temperature, isoprene and O<sub>3</sub>-isoprene-temperature relationships under  
 473 different climate scenarios. **(a-b)** Projections of the summertime daytime air temperature **(a)**  
 474 and isoprene concentrations **(b)** during 2030-2100 in Hong Kong. The shaded areas represent  
 475 the 25th to 75th percentile of the estimated isoprene concentration and temperature for each  
 476 SSP. **(c-f)** Responses of simulated O<sub>3</sub> concentrations to temperature and isoprene under  
 477 abundant-NO<sub>x</sub> **(c)** and reduced-NO<sub>x</sub> **(d-f)** conditions. The squares represent the projected O<sub>3</sub>  
 478 concentrations at specific temperatures and isoprene levels, with error bars indicating the  
 479 standard deviation of isoprene concentrations and temperatures.

480

481 **4 Conclusions**

482 As one of the most reactive and abundant VOC, isoprene plays a significant role in  
483 shaping urban air quality. We developed an explainable deep transfer learning  
484 framework to predict isoprene concentrations and elucidate the underlying drivers of  
485 their variability. Our model outperformed conventional approaches, effectively  
486 capturing the spatial heterogeneity of isoprene concentrations through localized fine-  
487 tuning. Leveraging this framework, we quantified the relative importance of factors  
488 influencing isoprene concentrations across numerous sites in China and internationally.  
489 The contrasting cases of Hong Kong and London highlight how isoprene dynamics  
490 were shaped by distinct local drivers, underscoring the need to tailor air quality  
491 management strategies to specific urban contexts. Despite the anticipated increase in  
492 biogenic emissions in a warming climate, our findings caution against reducing urban  
493 greenspace solely based on isoprene-related concerns. Instead, mitigating global  
494 warming emerges as a crucial strategy for managing isoprene's air quality impacts, as  
495 evidenced by the strong isoprene–temperature relationships observed. For  $O_3$   
496 abatement, coordinated control of  $NO_x$  emissions appears effective in reducing the  
497 contribution of isoprene to  $O_3$  formation. Moreover, the differing responses of isoprene  
498 to VI between Hong Kong and London suggest that informed tree species selection can  
499 serve as an alternative urban planning measure. Traffic emissions may also remain a  
500 significant source of urban isoprene in cities lacking stringent vehicle emission controls  
501 and should be addressed accordingly. Overall, this study provides novel insights into  
502 isoprene emissions and chemistry, air quality impacts, and practical mitigation  
503 strategies. Nonetheless, limitations persist, particularly regarding the comprehensive  
504 representation of emissions and chemical loss processes, which are discussed in Text  
505 S2.

506

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515 **Open Research**

516 Meteorology data for 1990-2023 at each site were obtained from the hourly ERA5  
517 reanalysis dataset (Hersbach et al., 2023); The CMIP6 model outputs can be accessed  
518 at <https://pcmdi.llnl.gov/CMIP6/>; The NDVI data from 1990-2022 are available at  
519 <https://doi.org/10.3334/ORNLDAA/2187>; The GLASS LAI V5 and V6 products are  
520 downloaded from <https://www.glass.hku.hk/download.html> and  
521 <https://www.geodata.cn/main/>, respectively.

522 **Author contributions**

523 X.L. and N.W. conceived the study. S.L. developed the methodology. Data collection  
524 was performed by S.L., N.W., X.L., Z.S., X.H., T.L., H.W., M.L., J.G., N.C., G.S., Y.Z.,  
525 C.P., Z.L., C.T., and X.L. Data analysis was conducted by S.L., X.L., and N.W. N.W.  
526 and X.L. led the investigation and supervision. Visualization was completed by S.L.  
527 and N.W. The original draft was written by S.L., N.W., and X.L. All authors, including  
528 F.Y., Z.S., X.H., and A.G., contributed to reviewing and editing the manuscript.

529 **Competing Interests**

530 The authors declare that they have no known competing financial interests or personal  
531 relationships that could have appeared to influence the work.

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