Τ	Global warming will largely increase waste treatment CH ₄ emissions in Chinese Megacities:
2	insight from the first city scale CH ₄ concentration observation network in Hangzhou city,
3	China
4	
5	Cheng Hu ^{1,2} , Junqing Zhang ¹ , Bing Qi ^{3,4*} , Rongguang Du ^{3*} , Xiaofei Xu ⁴ , Haoyu Xiong ⁵ , Huili
6	Liu ¹ , Xinyue Ai ¹ , Yiyi Peng ¹ , Wei Xiao ²
7 8 9	¹ College of Biology and the Environment, Joint Center for sustainable Forestry in Southern China Nanjing Forestry University, Nanjing 210037, China ² Collaborative Innovation Center on Forecast and Evaluation of Meteorological Disaster.
10 11	(CIC-FEMD), Nanjing University of Information Science & Technology, Nanjing, China ³ Hangzhou meteorological bureau, Hangzhou 310051, China
12 13	⁴ Zhejiang Lin'an Atmospheric Background National Observation and Research Station, Hangzhou 311300, China
14	⁵ College of Environment, Zhejiang University of Technology, Hangzhou 311300, China
15	
16	
17	
18	
19	
20	
21	
22	*Compounding outhors: Ding Oi (kill 120@sing com) Dangeryong Dy (dra1000@162 com)
	*Corresponding authors: Bing Qi (bill_129@sina.com), Rongguang Du (drg1998@163.com).
23	
24	
25	
26	
27	
28	
29	To be submitted to: ACP
30	
31	
22	

Abstract:

33

34

35

36

37

38

39

40

41

42

43

44

45

46

47

48

49

50

51

52

53

54

55

56

57

58

59

60

61

Atmospheric CH₄ is the second largest anthropogenic contributor to global warming. However, its emissions, components, spatial-temporal variations and projected changes still remain large uncertain from city to national scales. CH₄ emissions from waste treatment (including solid waste landfills, solid waste incineration and sewage) account for >50% of total anthropogenic CH₄ emissions at city scale, and considering the high temperature sensitivity of CH₄ emission factors (EFs) for the biological processes-based sources such as waste treatment, large bias will be caused when estimating future CH₄ emissions under different global warming scenarios. Furthermore, the relationships between temperature and waste treatment CH₄ emissions have only been conducted in a few site-specific studies and lack the representativity for whole city, which contains various biophysical conditions and shows heterogeneous distribution. These above factors cause uncertainty in the evaluation of city scale CH₄ emissions (especially from waste treatments) and projected changes still remain unexplored. Here we conduct the first tower-based CH₄ observation network with three sites in Hangzhou city, which is located in developed Yangtze River Delta (YRD) area and ranks as one of the largest megacities in China. We found the a priori total annual anthropogenic CH₄ emissions and those from waste treatment were overestimated by 36.0% and 47.1% in Hangzhou city, respectively. In contrast, the total emissions in the larger region, such as Zhejiang province or the YRD area, were slightly underestimated by 7.0%. Emissions from waste treatment showed obvious seasonal patterns following air temperature. By using the linear relationship constructed between monthly waste treatment CH₄ emissions and air temperature, we find the waste treatment EFs increase by 38%~50% with temperature increases of 10°C. Together with projected temperature changes from four climate change scenarios, the global warming induced EFs in Hangzhou city will increase at the rates of 2.2%, 1.2%, 0.7% and 0.5% per decade for IPCC AR5 (International Peace Cooperation Center, the fifth assessment report) RCP (Representative Concentration Pathway)8.5, RCP6.0, RCP4.5 and RCP2.6 scenarios, respectively. And the EFs will finally increase by 17.6%, 9.6%, 5.6%, and 4.0% at the end of this century. Additionally, the derived relative changes in China also show high heterogeneity and indicate large uncertainty in projecting future national total CH_4 emissions. Hence, we strongly suggest the temperature-dependent EFs and the positive feedback between global warming and CH₄ emissions should be considered in future CH₄ emission projections and climate change models.

Keyword: CH₄ emissions, waste treatment, observation network, global warming

1. Introduction

62

63

64

65

66

67

68

69

70

71

72

73

74

75

76

77

78

79

80

81

82

83

As the second largest anthropogenic greenhouse gas, the reduction of CH₄ emissions is considered an effective way to mitigate future climate change on short timescales (Henne et al., 2016; Lin et al., 2021). Accurate estimation of CH₄ emissions from its main sources is the basis of policy making. However, recent studies find there still remain large uncertainties for its total emissions, components, spatial-temporal variations and projected changes at city scale especially for megacities in China (USPA 2013; Cai et al., 2018; Lin et al., 2021). CH₄ emission from waste treatment (mainly including sewage and solid waste by landfills and incineration) ranked as the world's third largest anthropogenic source after fuel exploitation and livestock, and was responsible for ~13% of global anthropogenic CH₄ emissions of 371 (±26) Tg a⁻¹ (Lu et al., 2021). It also ranked as the fourth largest anthropogenic source in China, the biggest anthropogenic CH₄ emitting country, and accounted for ~14% of national total anthropogenic emissions of 65 (±22) Tg a⁻¹ (Saunois et al., 2020; Lu et al., 2021; Chen et al., 2022). Furthermore, its contribution is even larger than 50% at city scale especially for megacities, where both active and closed household waste (including landfills and waste water systems) are located and found as super emitters (Williams et al., 2022; Maasakkers et al., 2022). A large number of Chinese landfills were constructed in suburbs more than 5-10 years ago and most landfills have no gas collection systems, with the urban area expanding in recent decades, the locations of many landfills are now within the urban scope (Zhejiang Statistical Yearbook 2018-2019). In addition, the decreasing area of the agricultural sector (rice paddies and husbandry) in megacities also makes their emissions negligible when compared with waste treatment. Therefore, accurate quantification of CH₄ emissions from waste treatment in urban area becomes increasingly important.

84

85

86

87

88

89

90

Although some progress has been made in measuring site scale CH₄ emissions from waste treatment, the estimated emissions still show large discrepancies due to many factors such as the amount of waste and its composition, relative proportions of landfills and incineration, degradable organic carbon ratio, CH₄ oxidation efficiency, and landfill gas collection, and meteorological conditions including temperature, water content, atmospheric pressure (Masuda et al., 2018; Cai et al., 2018; Zhao et al., 2019; Hua et al., 2022; Bian et al., 2022; Maasakkers et al., 2022; Kissas et

91 al., 2022).

Furthermore, CH₄ emissions from sewage and landfills result from microbial processes especially from methanogens, and their emission factors (EFs) are highly sensitive to temperature. These available studies were mainly conducted at some specific sites with measured EFs varying widely (Du et al., 2017; 2018; Cai et al., 2014; 2018; Zhao et al., 2019; NBSC, 2015; Wang et al., 2015; Florentino et al., 2010; Tolaymat et al., 2010; Hua et al., 2022). The lack and discrepancies of detailed information for all the above factors and their uncertainties have led to considerable difficulty in estimating CH₄ emissions for most-to-date inventories (Höglund-Isaksson, 2012; USEPA et al., 2013; Cai et al., 2018; Lin et al., 2021; Maasakkers et al., 2022).

China, the developing country with the largest anthropogenic CH₄ emissions, is expected to increase its emissions because of projected rapid economic development, urbanization and generated waste (Cai et al., 2018). The increase of waste treatment emissions in East China was also found as the second largest sector in driving national total anthropogenic CH₄ emissions since 2000 (Lin et al., 2021). In addition, the mitigation potential of waste treatment in developing countries is thought to be four times that of developed countries (USEPA, 2013). Therefore, mitigating CH₄ emissions from waste treatment in China is a robust and cost-effective way to reduce total national anthropogenic greenhouse gas emissions.

Many previous studies have estimated the waste treatment CH₄ emissions for China by both "bottom-up" and "top-down" approaches, with results varied by 2.5-fold from 4.3 to 10.4 Tg CH₄ yr⁻¹, and accounted for 8.1%~24.2% of national total anthropogenic CH₄ emissions (USEPA 2013; Peng et al., 2016; Miller et al., 2019; Lin et al., 2021; Lu et al., 2021; Chen et al., 2022). For these "bottom-up" approaches, the high uncertainties were directly attributed to omission of many small point sources and discrepancies of observed site-specific EFs, which varied largely by climate and management technology such as the efficiency of gas collection systems (Zhao et al., 2019; Hua et al., 2022). Previous studies most commonly used the EDGAR (Emission Database for Global Atmospheric Research) inventory, using the IPCC recommended default EF values of 15.0%

(Höglund-Isaksson, 2012; Lin et al., 2021; Bian et al., 2022), but this value is around 5-7 times of EFs used in China by Zhang and Chen et al. (2014). A recent study comparing waste treatment CH₄ emissions among different inventories also reported that the EDGAR v5.0 and CEDS (Community Emissions Data System) inventories were 21~153% higher than other inventories, and EDGAR v5.0 tended to assign more emissions in urban areas especially for provincial capitals. In addition, emissions from wastewater were found to be overestimated by higher emission factors or chemical oxygen demand (Peng et al., 2016; Lin et al., 2021).

127

128

129

130

131

132

133

134

135

136

137

138

139

140

141

142

143

144

145

146

147

148

120

121

122

123

124

125

126

And for the "top-down" atmospheric inversion approaches, a few studies constrained anthropogenic sources including waste treatment, where the most widely used concentrations were from satellite observations (Miller et al., 2019; Lu et al., 2021; Chen et al., 2022). The satellite observations have the advantage of easy data access and global coverage. But as already noted, the emissions constraint results are highly dependent on availability of observed concentrations, which are largely influenced by weather conditions and cloud coverage. As was illustrated in a recently published study by Chen et al. (2022), although the numbers of grid cell (0.25° ×0.3125°) based year-round satellite observations were more than 1000 in north China, the available numbers were less than 10 (even including grid cells without any observations) in most of central, west, east and south of China. Such sparse distribution of available data may not provide robust constrains on waste treatment emissions for some Chinese cities without enough observations, especially considering waste treatment is co-located with high population density megacities in the developed area of east and south of China. Furthermore, there should be large temperature induced monthly variations for waste treatment CH₄ emissions (Börjesson et al., 1997), but almost all satellite-based inversions were conducted at annual scale without seasonal variations. Besides, given the strong influence from atmospheric pressure on landfill CH₄ emissions (Kissas et al., 2022), satellite observations are too sparse to be up-scaled to estimate annual total because satellite observations are mostly available only on clear-sky conditions and cannot represent atmospheric pressure and CH₄ emissions on cloudy or rainy days. There was only one recent study which focused on urban waste treatment CH₄ emissions, it found annual CH₄ emissions from four cities were 1.4 to 2.6 times larger than inventories in India and Pakistan, where landfills contributed to 6~50% of total emissions and indicated large bias of our understanding of waste treatment CH₄ emissions (Maasakkers et al., 2022).

The tower-based atmospheric inversion approach, which is based on hourly atmospheric concentration observations within the planetary boundary layer, can be used independently to constrain CH₄ emissions and its main components. Besides, compared with "bottom-up" approaches, the "top-down" method can avoid using the factors that lead to large uncertainties in CH₄ emissions especially from waste treatment. And to our best knowledge, there are few tower-based observation inversion studies which focus on waste treatment emissions at city scale or much larger regional scales especially in China. Only one study in Los Angeles, U.S.A. used tower-based CH₄ concentration and found the influence of a landfill site closure on CH₄ emissions, which was not included in a priori inventory (Yadav et al., 2019). In addition, the influences of global warming on city scale (or higher regional scale) emissions are still unclear and have not been considered in future emission projections (USEPA 2013; Cai et al., 2018). In general, previous studies which predicted future waste treatment CH₄ emissions only used activity data changes, without considering climate change on the EFs. Considering the potential high sensitivity of waste treatment CH₄ emissions on the projected global warming, how these emissions will change with increasing temperature is still unknown, especially within megacities where more waste is generated and the urban heat island effect will lead to much stronger warming climate (Zhang et al., 2022).

Here, we established three tower-based CH₄ concentration observation sites in Hangzhou city, one of the largest megacities in China. To our best knowledge, it is the first city-scale tower-based CH₄ concentration observation network in China. We present our work on urban CH₄ emissions inversion and aim to (1) constrain CH₄ emissions from waste treatment alongside total anthropogenic emissions in Hangzhou city, (2) derive temperature sensitivity of waste treatment CH₄ emissions at city scale and quantify the projected emission changes in future climate change scenarios. One-year hourly CH₄ concentration observations from December 1st, 2020 to November 30th, 2021 were combined with atmospheric transport model and Bayesian inversion

approach to constrain monthly CH₄ emission inventories. The constructed relationship between monthly temperature and *posteriori* waste treatment CH₄ emissions will be used with future temperature projection to quantify how the EFs will change in different global warming scenarios.

181

182

183

184

185

186

187

188

189

190

191

192

193

194

195

196

197

198

199

200

178

179

180

2. Materials and Method

2.1 Tower-based CH₄ observation network and supplementary materials

The city of Hangzhou, which has a population of 12.2 million and area of 1.7×10^4 km² (core urban area of $8.3 \times 10^3 \,\mathrm{km}^2$), is the capital of Zhejiang province and located in the middle of East China (Figure 1a). As displayed in Figures S1-S2, the East China accounts for the majority of the national total population and waste treatment CH₄ emissions. Hangzhou city ranked in the top 10 megacities in China, with annual solid waste of around 5 million tons in 2021. The tower-based CH₄ concentration observation network includes three observation sites (Figure 1a-d), as (1) Hangzhou site (120.17° E, 30.23° N, 43.2 m a.s.l.), which is located in the core urban region; (2) Linan site (119.72° E, 30.30° N, 138.6 m a.s.l.), regional background site with no obvious emission sources within 10 km radius; (3) Damingshan site (119.00° E, 30.03° N, 1485.0 m a.s.l.), which is built on the top of a 1500 m mountain and represents background from much more diluted regional emission signals. The distance is around 50 km between Hangzhou site and Linan site, and around 150 km between Hangzhou site and Damingshan site. These three sites represent obvious gradients from east of densely populated area (Figure 1c-d) and anthropogenic emissions to west of much weaker anthropogenic influence and background conditions. Based on the wind direction for the three sites, there is not any obvious difference of seasonal wind direction patterns among them. The prevailing wind direction from October to February was from the north, which changed to east from February to May and then changed to south during the monsoon in summer.

201

202

203

204

205

206

The air inlet heights are 25 m above ground for the Hangzhou site, 53 m at Linan and 10 m at Damingshan, respectively. Atmospheric CH₄ concentrations at all three sites were continuously measured by cavity ring-down spectroscopy analyzer (model G2301 for Hangzhou site and G2401 for Linan site and Damingshan site; Picarro Inc., Sunnyvale, CA). To obtain high precision observations, two different standard gases were measured every 6 hours and a linear two-point fit

was used to calibrate observations, with the precision and accuracy of 2 ppb and 1 ppb. More details of the observation and calibration systems were described in Fang et al., (2014; 2022). Note that because of instrument issues at Damingshan site, there is a data gap in September-October, 2021. In general, 99.4%, 99.0%, 79.3% of hourly CH_4 observations were available in the whole year observation period for Hangzhou site, Linan site and Damingshan site, respectively. Meteorological observations at Hangzhou meteorological station were used to evaluate simulated meteorological fields, including air temperature at 2 m (T_{2m}), relative humidity (RH), downward solar radiation ($S\downarrow$), wind speed (WS) at 10 m height, and planetary boundary layer height (PBLH).

Note some previous studies of city scale greenhouse gas concentration observation networks chose sites at the edge of urban borders as background in emission inversion system (i.e. Indianapolis, U.S.A., Miles et al., (2017); Los Angeles, U.S.A., Verhulst et al., (2017); Washington, DC-Baltimore, U.S.A., Lopez-Coto et al., (2020); Paris, France, Lian et al., (2021)), but we chose to use five NOAA CH₄ background sites as the potential background, including UUM, TAP, YRO, YON and WLG site (Figure 1a), which were much further than the observations at Damingshan site. This strategy is based on following three reasons: (1) our footprint domain is much larger than Hangzhou city and these five sites are also located close to the edge of the model domain; (2) CH₄ concentrations within Hangzhou city will be influenced by seasonally varying monsoon and the monthly varying wind directions will lead to obvious changes of CH₄ background than only at Damingshan site; (3) our model setups can partition CH₄ enhancements from within Hangzhou city and other regions.

The projected climate data from four RCP (Representative Concentration Pathway) scenarios (RCP8.5, RCP6.0, RCP4.5 and RCP2.6) by MRI-CGCM3 model were downloaded from World Data Center for Climate (WDCC, https://www.wdc-climate.de/ui/), where annual air temperature at 2m was used from years 2021 to 2100. The most recent population density data for Hangzhou city is for the year 2019 and was downloaded from Chinese national resource and environmental science and data center.

2.2 WRF-STILT model setup

The WRF-STILT (WRF: Weather Research and Forecasting, version 4.2.2, and STILT: Stochastic Time-Inverted Lagrangian Transport) model was used to simulate hourly footprint and CH₄ enhancement, see more details in Hu et al. (2019; 2021). Domain setups are displayed in Figure 1a, with the outer nested domain (Domian-1, 27 km×27 km grid resolution) covering eastern and central China, and the inner domain (Domain-2, 9 km×9 km grid resolution) covering the YRD area. The physical schemes used in the WRF model are the same as in our previous studies for the YRD domain (Hu et al., 2019; 2021). The simulated CH₄ concentration is the sum of background and enhancement, where the enhancement is calculated by multiplying all CH₄ flux with hourly footprint that represents the sensitivity of the concentration changes to its regional sources/sinks with spatial resolution of 0.1°×0.1°. To better quantify CH₄ components at each site, CH₄ enhancements from different regions and sources are also tracked and separately simulated. Besides, we should note the CH₄ background is important in simulating CH₄ concentrations and atmospheric inversion. We will choose CH₄ background from the five background sites based on monthly footprint as discussed in Section 3.1.

The most recent inventory of Emission Database for Global Atmospheric Research (EDGAR v6.0), which has 20 categories, and WetCHARTs ensemble mean were used as the *a priori* anthropogenic and natural CH₄ emissions. We should note there are many CH₄ inventories for some developed regions and countries (i.e. France, U.S.A., Germany) with high spatial resolutions. The reasons to choose EDGAR as *a priori* anthropogenic emissions are: (1) for all available CH₄ inventories that covered China, the spatial resolution of EDGAR (0.1°×0.1°) is the highest, and it provides the most up-to date results; (2) most previous studies that constrain emissions by atmospheric inversion studies also chose EDGAR, and our results can be directly compared with previous studies; (3) the preliminary simulation of CH₄ concentrations showed generally good performance with observations, indicating its spatial distributions in Hangzhou city has relatively small bias even with a potentially large bias for magnitude, which will be constrained by our atmospheric inversion method.

The main sources of CH₄ emissions in Hangzhou city include SWD_LDF (solid waste landfills),

WWT (waste water handling), SWD_INC (solid waste incineration), PRO (all processes related to fuel exploitation from coal, oil, and natural gas, including extraction, transportation, refining, distribution as list in IPCC database (https://www.ipcc-nggip.iges.or.jp/EFDB/find_ef.php), RCO (energy for buildings, mainly containing nature gas escaping from household use) and AGS (agricultural soils). We found emissions from SWD_LDF, WWT and SWD_INC were simply assigned in the same locations in EDGAR inventory, and hence combined them as waste treatment. For the CH₄ emissions from wetland, we used WetCHARTs ensemble mean with spatial resolution of 0.5° at monthly average (Bloom et al., 2017). Considering WetCHARTs treats rice paddies (main source as AGS) as one wetland type, AGS in EDGAR was excluded and we assume WetCHARTs represent all wetland CH₄ emissions as natural wetland and rice paddies.

277

278

267

268

269

270

271

272

273

274

275

276

2.3 Bayesian inversion framework

- The Scale Factor Bayesian inversion (SFBI) approach was applied to interpret the atmospheric CH₄ concentration (or enhancement) variations in terms of quantitative constraint on all CH₄
- sources. The relationship between observed and simulated CH₄ concentrations (or enhancement)
- can be expressed as follows in Equation 1:

$$y = K\Gamma + \varepsilon \quad (1)$$

- Where y is the observed CH₄ concentration (or enhancement), K corresponds to simulated
- 285 enhancements from all categories, Γ is the state vector to be optimized and consists of *posteriori*
- SFs for corresponding categories in K, and ε is the observing system error.

- The optimal solution to derive *posteriori* SFs is to minimize a cost function $J(\Gamma)$, which represents
- the mismatch between CH₄ observations and simulations and the mismatch between *posteriori* and
- 290 *a priori* SFs (Miller et al., 2008; Griffis et al., 2017). The cost function $J(\Gamma)$ can be expressed as:

291
$$J(\Gamma) = \frac{1}{2} \left[(y - K\Gamma)^T S_e^{-1} (y - K\Gamma) + (\Gamma - \Gamma_a)^T S_a^{-1} (\Gamma - \Gamma_a) \right]$$
 (2)

- where S_e and S_a are the constructed error covariance matrices for observations and the *a priori*
- values, and S_e consists of measurement and model errors. Here each element in a priori SFs Γ_a
- is treated as 1. Therefore, the solution for obtaining the *posteriori* SFs is to solve $\nabla_{\Gamma} J(\Gamma) = 0$,
- and is given by,

296
$$\Gamma_{\text{post}} = (K^T S_e^{-1} K + S_a^{-1})^{-1} (K^T S_e^{-1} y + S_a^{-1} \Gamma_a)$$
 (3)

In the Bayesian inversion framework, we first need to give an estimate of the error covariance matrices and the state vector for the *a priori* and observational data. And following our previous studies conducted in East China (Hu et al., 2019; 2022). Uncertainties of 10%, 13% and 20% were assigned to the measurement errors (S_{obs}), the finite number of particles (500) released in the STILT model ($S_{particles}$) and uncertainty in meteorological fields (S_{met}), respectively.

A previous study derived uncertainties of CH₄ from waste treatment and other categories, which varied between 30% and 50%, these uncertainties were calculated mainly from activity data and EFs at the country scale on annual averages (Solazzo et al. 2021). We should also note CH₄ emissions uncertainty will largely increase as the study region size decreases, and, as stated above, the relative difference among different inventories can reach 150%. Considering the disaggregation of spatial distributions and temporal variations, CH₄ emission uncertainties can be much larger at urban and monthly scales. To provide robust constraints on CH₄ emissions in our study, we used three cases of *a priori* uncertainty combinations for different emissions in Bayesian inversion:

(1) the first case use three elements as wetland, waste treatment and all other anthropogenic sources, considering the larger seasonality of waste treatment, the uncertainties of 300% was used for waste treatment and 200% for other categories, (2) the second case have more detailed categories as wetland, waste treatment, fuel exploitation, energy for building, and the other anthropogenic sources, where the *a priori* uncertainty of 200% was used for each category, (3) the third case has the same categories as case 1 but uses a different *a priori* uncertainty for waste treatment of 200%. The averages of all three cases are used as final *posteriori* SFs and the largest difference between each of three cases is used as the final uncertainty.

3. Results

3.1 Atmospheric CH₄ observations

We first display the hourly CH₄ concentrations from our three tower-based sites and smoothed background at five sites by CCGCRV fitting method (Thoning et al., 1989) in Figure 2a. The

hourly observations at three towers show similar temporal variations but with different amplitudes. Observations at Hangzhou site vary between 2000 ppb and 2800 ppb, and were much larger than both Linan site and Damingshan site. Their monthly averages are also compared in Figure 2b, and results show the monthly CH₄ vary between lowest 2106.3 ppb in July and highest 2225.0 ppb in September (annual mean of 2159.9 ppb) at Hangzhou site, lowest 2023.3 ppb in July and highest 2132.0 ppb in September (annual mean of 2086.7 ppb) at Linan site, the lowest 1955.5 ppb in July and without observations in September at Damingshan site (annual mean of 2013.4±(3) ppb, where the uncertainty was calculated based on the assumption that monthly CH₄ concentration in September and October varies between August and November), respectively. The similar trends among the three sites can be explained by all three sites being dominated by similar atmospheric transport processes, such as synoptic process (i.e. monsoon) and seasonally changing wind directions as summarized above. But their surrounding emission sources are highly different, implying the emissions of Hangzhou site should be much larger than Linan and Damingshan sites.

Because the CH₄ background is important in concentration simulation and emission inversion, we also compare CH₄ background between five sites, where the annual averages at TAP, YON, RYO, WLG and UUM were 1989.8 ppb, 1850.1 ppb, 1982.7 ppb, 1973.4 ppb and 1984.2 ppb, respectively. We found the differences were generally within 20 ppb among TAP, RYO, WLG and UUM sites (Figure 2), but there was large difference between YON site and other four sites from May to August, which can reach to around 100 ppb. Note YON site is located in the south of East China Sea (Figure 1a), it can be influenced by monsoon with clean air flows from the South China Sea, which has many fewer CH₄ sources compared to air flows from East Asia. The CH₄ background at TAP site appeared slightly higher than other four sites because TAP site is located in the coast of South Korea and can be more easily polluted by anthropogenic emissions. Considering the large spatial difference between the CH₄ background sites, monthly air flows and source footprint will be used to identify backgrounds for our observation network, with details discussed in Supplementary Material (Section S1, Figure S3 and Table S1).

3.2 Concentration footprint and the *a priori* emissions

To illustrate the potential source regions of the three sites, annual averages of simulated footprints for each site are displayed in Figure 3a-c. The results show their footprint distributions were quite similar because of close distance, but we also notice there were obvious differences in the footprint strengths (i.e. the area covered by red color) with Hangzhou site > Linan site > Damingshan site. The reason why the footprint at the Damingshan site is the lowest can be explained that the observations were collected at 1500 m height, and it was not easy to receive emissions signals within boundary layer at that height. Besides, the Hangzhou site is located in the core urban area of Hangzhou city, and it will show significant diurnal variation in PBLH, especially since it has higher nighttime PBLH caused by anthropogenic heat and high buildings than grassland/farmland, which dominate Linan site and Damingshan site. Hence more air particles can remain within PBLH and generate stronger footprint.

The a priori EDGAR CH₄ emissions for total anthropogenic categories, waste treatment and its proportions are given in Figure 3d-f. Significant gradients are observed from higher emissions in the east to lower emissions in the west, which is consistent with our three tower-based sets of observations. And the CH₄ emissions for waste treatment indicated similar spatial distributions with urban land use and population density (Figure 1c-d). Moreover, waste treatment seems to emit CH₄ as area sources instead of point sources from waste treatment super plants. Although a few previous studies found limitations of EDGAR inventory to capture CH₄ emission patterns in some urban areas (Pak et al., 2021), here considering the fact that locations of landfills (Figure 1b-d), which is the largest anthropogenic CH₄ emitter in Hangzhou city, are very close to the core urban area and in high consistency with EDGAR, hence we believe the spatial patterns of EDGAR in study region to be reliable. We should note the Chinese government constructed waste separation stations in each city with density of one station for per 150~200 households (around 450~800 people), usually these waste separation stations are full with waste because domestic garbage can be generated every day, they do not have gas collection systems and can emit large quantity of CH₄ emissions caused by daily biomass waste as area sources (Tian et al., 2022). Besides, there is only one landfill that has gas collection systems, the reported gas collection

efficiency was less than 80%, which also indicates large quantity of CH₄ emissions will be directly emitted into the atmosphere and the emissions will be influenced by climate change. These above analyses also imply Hangzhou site can observe higher emissions from both waste treatment and total anthropogenic emissions, which will be discussed and quantified later.

3.3 Simulation of CH₄ concentrations and its components for three sites

Comparisons between observed and simulated daily CH₄ concentration averages are displayed in Figure 4a-c and hourly concentrations in Figure S4 for three sites. First, the hourly simulations in Figure S4 show high consistency when only comparing the temporal patterns with observations, indicating good performance of model transport simulations as confirmed in Figure S5 for evaluating meteorological fields. But the relative variations display obvious differences among the three sites for daily averages in Figure 4a-c. The mean bias (MB), root mean squared error (RMSE), and correlation coefficient (R) between daily observations and *a priori* simulations were 64.1 ppb, 129.2 ppb and 0.44, respectively, for Hangzhou site; and were -6.0 ppb, 57.1 ppb, 0.50 for Linan site, 36.2 ppb, 55.6 ppb, 0.54 for Damingshan site. As for the Hangzhou site, simulated CH₄ concentrations show obvious overestimation from October to April, and the overestimation is also found at Damingshan site. We found the simulations at the Linan site showed overall good agreement with observation, but still with slight overestimation from January to April and underestimation from May to September. Considering the source area contributions for the three sites are different, these differences among the three sites indicated the bias in CH₄ emission largely varied from Hangzhou city to larger regional scale.

To further quantify detailed contributions from different regions and categories to each tower site, CH₄ enhancements from different categories and source areas were also simulated separately for the three sites. As displayed in Figure 4d-e, the simulated *a priori* total enhancements at Hangzhou site, Linan site, and Damingshan site were 244.3 ppb, 100.8, and 69.0 ppb, respectively. We also found contributions by waste treatments dominated the total enhancements but with obvious differences among the three sites, which varied from the highest 64.2% at Hangzhou site to the lowest 41.4% at Damingshan site. We further calculated anthropogenic contributions from

Hangzhou city (excluding wetlands because of coarser spatial resolution for Hangzhou city) and other provinces, which were 158.4 ppb at Hangzhou site, 30.7 ppb at Linan site, and 10.1 ppb at Damingshan site, respectively. And they accounted for 69.3%, 34.0%, and 16.9% of total anthropogenic enhancements at corresponding sites. These results indicate the CH₄ observations at Hangzhou site, which is located at the core urban region, are more influenced by local emissions (mainly for waste treatment which will be discussed later) and contain much higher enhancements than the other two sites. The relative contributions from Hangzhou city to observations at the Hangzhou site, Linan site and Damingshan site were 158.4 ppb (69.3% to total CH₄ enhancement), 30.7 ppb (34.0% to total CH₄ enhancement), and 10.1 ppb (16.9% total CH₄ enhancement), respectively. The relative contributions from Zhejiang province to observations at the Hangzhou site, Linan site and Damingshan site were 181.7 ppb (79.5% to total CH₄ enhancement), 44.3 ppb (49.0% to total CH₄ enhancement), and 17.9 ppb (29.9% total CH₄ enhancement), respectively. These different values also imply that the observations at Linan and Damingshan sites can represent CH₄ emissions of much larger region as Zhejiang province or YRD area than Hangzhou city (Figure 4e), and Daminshan site.

The seasonally averaged diurnal variations for both observations and simulations are also displayed in Figure 5 for the three sites. Although many previous studies only used daytime observations and simulations to evaluate *a priori* emissions bias and constrain emissions (Sargent et al., 2018; Hu et al., 2022), these studies were based on the assumption that the diurnal scaling factors used for the *a priori* emissions are right (i.e. for anthropogenic CO₂), or the emissions do not have obvious diurnal variations (i.e. emissions from industries or manufacturing). As concluded above, the main CH₄ component in Hangzhou city was waste treatment (Figure 3f), which should be highly sensitive to temperature and indicates obvious diurnal and seasonal patterns (Mønster et al., 2019; Kumar et al., 2022). And total CH₄ emissions will be overestimated when using daytime emissions to represent all-day averages. Further, we found strong similarities of the diurnal variations between observations and simulations for the three sites, but there are still some discrepancies especially that the observations at Linan site were generally higher than simulations from spring to autumn for both all-day and midday averages.

Hence, our preliminary conclusions were that the *a priori* CH₄ emissions were generally overestimated for Hangzhou city but underestimated in the larger region of Zhejiang or YRD area. We also found simulations were higher than observations for all seasons at Damingshan site, and it can be explained by the complex topography around the Damingshan site, where elevations changed from 0 m to 1600 m within the site's grid cell of 9 km ($\sim 0.1^{\circ}$) as displayed in Figure 1b, and the mountain-valley wind patterns, PBLH changes can only be resolved with much higher spatial resolutions of < 1km. Hence the use of coarse resolutions (i.e. 9 km in this study) at the mountainous regions introduces large bias in simulating concentration and emission inversion, as also recently found in China for CO_2 as "aggregation error" (Agustí-Panareda et al., 2019; Wang et al., 2022), so observations at Damingshan site will not be used in emissions inversions in this study.

3.4 Constraints on anthropogenic CH₄ emissions

As displayed in Figures 3f, 5a and concluded in Section 3.3, simulations using a priori CH₄ emissions show obvious overestimation especially from October to April at Hangzhou site, and emissions were also overestimated in winter and underestimated from spring to autumn at Linan site. Note this bias can be attributed to a priori emissions or meteorological simulations. Our previous studies in YRD have evaluated the meteorological simulations by using the same physical parameterization schemes, which showed high consistency with observations (Hu et al., 2019; 2021; 2022; Huang et al., 2021). We also evaluated the meteorological simulations with observations and confirmed with good model performance (Figure S5). Note PBLH simulations are important in evaluating model performance, we only have four months of PBLH observations (one month in each season), these hourly PBLH observations were used to evaluate the general performance of WRF model. As displayed in Figure S6, it shows overall good performance for both daytime and nighttime PBLH variations. Furthermore, we found there no monthly variations in EDGAR v6.0 CH₄ emissions for waste treatment, which contributed 64.2% to annual CH₄ enhancement average and much higher in winter (Figure S7-S8). The CH₄ emissions from waste treatment are produced by the microbial process, which should be affected by meteorological conditions especially by seasonal temperature changes. Hence our assumption is that the bias in both its seasonality and annual average lead to large overestimation/underestimation in the simulated CH₄ concentration. Besides, bias in other anthropogenic emissions and wetlands can also partly contribute to the bias of the simulated CH₄ concentration.

473

474

475

476

477

478

479

480

481

482

483

484

485

486

487

488

489

490

491

470

471

472

To quantify the bias sources and constrain corresponding a priori emissions for Hangzhou city, we applied the scaling factor Bayesian inversion approach with three different cases as introduced in the Method section. Instead of only using daytime CH₄ observations to constrain a priori emissions, we choose to use all-day hourly data at Hangzhou site to constrain emissions for Hangzhou city, for the following three reasons: (1) the enhancements contributed by Hangzhou city at the Hangzhou site was 69.3%, and much larger than 34.0%, and 16.9% for Linan site and Damingshan site, respectively; (2) the waste treatment dominated anthropogenic CH₄ emissions in Hangzhou city, which is caused by biological process and should be temperature dependent. Since the observed temperature varied diurnally by 20 °C, the use of only daytime observations without considering diurnal CH₄ emissions will bring significant bias when using derived daytime emissions to represent all-day averages. The annual averages of daytime and all-day average concentrations were 2112.4 and 2156.0 ppb at Hangzhou site, respectively, the reason why higher emissions in daytime correspond to lower concentration than in all-day and nighttime is that lower PBLH in nighttime will leads to higher concentration, and more comparisons between daytime and all-day average concentrations are displayed in Figure 5 for three sites; (3) previous studies using daytime observations were mainly conducted for regions dominated by industry or energy production, which have much smaller diurnal variations than waste treatment as stated above (Mønster et al., 2019; Kumar et al., 2022).

492493

494

495

496

497

498

The derived monthly *posteriori* SFs for each emission source are displayed in Table 1 for Hangzhou city. The results show that the *posteriori* SFs for waste treatment are much smaller in winter and higher in summer, indicating obvious seasonality and the overestimation in winter was mainly contributed by waste treatment. The annual mean *posteriori* SFs for waste treatment vary between 0.50 and 0.56 in all three cases, illustrating overestimation at annual average for the *a priori* waste treatment emissions. Besides, the annual mean *posteriori* SFs vary between 0.87 and

0.94 for the rest of the total anthropogenic categories (excluding agricultural soil), and are 0.97 for PRO (fuel exploitation) and 0.91 for RCO (energy for building), respectively; the annual mean *posteriori* SF is 1.05 for wetland (including agricultural soil and natural wetland). These *posteriori* SFs for the rest anthropogenic categories and wetland indicate much smaller bias than waste treatment. The monthly *posteriori* SFs for PRO and RCO also illustrate obvious seasonal variations, but are still smaller than the *a priori* seasonality in the inventory (Figure S9). Although the evaluations of hourly PBLH simulations have illustrated good performance in both daytime and nighttime (Figure S6), we also conducted inversions by only using daytime observations to constrain CH₄ emissions. Considering results from Case 2 varied between Case 1 and Case 3, here we only display the results from Case 1 and Case 3 (Table S2), it shows similar seasonal variations as using all all-day observations. We notice the values are larger than later, which is reasonable because CH₄ emissions in daytime should be larger than all-day and nighttime emissions. In general, *posteriori* SFs by using all-day concentration observations will be used to represent total CH₄ emissions from monthly to annual scales.

To evaluate whether the *posteriori* SFs have significantly improved CH₄ emissions, we used these SFs to derive the *posteriori* emissions and re-simulated hourly concentrations in Figure 6 (and daily averages in Figure S9). Results show the hourly overestimation by using *a priori* emissions is largely reduced by using *posteriori* emissions when compared with observations in Figure 6a-b, and the regression slopes between daily averaged observations and simulations decrease from 1.51(±0.15) for *a priori* simulations to 0.85(±0.07) for *posteriori* simulations in Figure 6c. The mean bias (MB), root mean squared errors (RMSE), correlation coefficient (R) between daily observations and *a priori* simulations are 64.1 ppb, 129.2 ppb and 0.44, respectively, and these statistics change to -22.2 ppb, 72.3 ppb and 0.58 for *posteriori* simulations. These results indicate the *posteriori* SFs obviously decrease the bias in *a priori* emissions and are closer to observations, when considering there are no system biases in simulated monthly PBLH.

The comparisons of monthly CH₄ emissions between *a priori* and *posteriori* waste treatment and other anthropogenic sources (excluding agricultural soil) in Hangzhou city are displayed in

Figures 7a and S7. For the *a priori* inventory, there is not seasonal variations for waste treatment with constant monthly emissions of 8.67×10^3 t, and other anthropogenic sources show seasonality with much higher in winter (i.e. 5.22×10^3 t in January) than in summer (i.e. $3.06 \times$ 10³t in August). The seasonality in a priori EDGAR inventory is mainly dominated by RCO (Energy for buildings), with proportions to total anthropogenic emissions changing from the highest 22% in winter to lowest 8% in summer. Such information indicates the a priori inventory assigned more leaks from natural gas distribution infrastructure in winter than in summer. As discussed above, constant emissions from waste treatment should be wrong because of its large temperature sensitivity, and the observed monthly temperature difference between summer and winter was larger than 25°C in Hangzhou city in study period. After including the constraints from the observed concentrations, the posteriori emissions for waste treatment show obvious seasonality with highest emission in July $(7.66 \pm 0.09 \times 10^{3})$ t) and lowest emission in February $(2.20 \pm 0.87 \times 10^3 \text{ t})$. And emissions from other anthropogenic categories show much smaller seasonality (highest emission in January of $4.18 \pm 0.69 \times 10^3$ t and lowest emission in August of $2.88 \pm 0.15 \times 10^3$ t) than a priori emissions. In general, the annual emissions from waste treatment were 10.4×10^4 t in the a priori EDGAR inventory and decreased to 5.5 (± 0.6) $\times 10^4$ t for the posteriori emissions, a decrease of 47.1%. The a priori emissions from other anthropogenic sources were 4.5×10^4 t and only slightly decreases to $4.1 \ (\pm 0.3) \times 10^4$ t for the posteriori emissions, an 8.9% decrease. The proportion of waste treatment to total anthropogenic emissions decreases from a priori 69.3% to posteriori 57.3%. To summarize, the annual total anthropogenic CH₄ emission (excluding agricultural soil) decreases from 15.0×10^4 t to $9.6 (\pm 0.9) \times 10^4$ t, indicating overestimation of 36.0% in Hangzhou city for the *a priori* emissions.

550

551

552

553

554

555

556

528

529

530

531

532

533

534

535

536

537

538

539

540

541

542

543

544

545

546

547

548

549

However, as concluded above the observations and simulations at Linan site, which represents the much larger region of Zhejiang province or YRD area, data from that site indicated slightly different results that CH₄ simulations were underestimated from spring to autumn and overestimated in winter (Figure 4b and Figure 5e-h). Here we used the multiplicative scaling factor (MSF) method and observations at Linan site to derive SFs at seasonal scale (Sargent et al., 2018; He et al., 2020), where we used 10 ppb as the potential CH₄ background uncertainty in

winter, spring and autumn, and 20 ppb in summer, see details in the Supplementary Material (Section S2). The derived *posteriori* SFs were 0.87 (\pm 0.08), 1.07 (\pm 0.11), 1.19 (\pm 0.24), and 1.16 (\pm 0.11) for winter, spring, summer, and autumn, respectively. The results for the Linan site showed similar seasonal variations as found for Hangzhou city and was 1.07 (\pm 0.14) of *a priori* anthropogenic emissions for the annual average. Our observations at Hangzhou site and Linan site together indicate the *a priori* emissions were largely biased on both seasonal and annual scales, and the annual anthropogenic CH₄ emission was largely overestimated by 36.0% in Hangzhou city, but was underestimated by 7.0% in the larger region of Zhejiang province or YRD area.

565

566

567

568

569

570

571

572

573

574

575

576

577

578

579

580

581

582

583

584

585

557

558

559

560

561

562

563

564

3.5 Temperature sensitivity of waste treatment CH₄ EFs and projected changes

Although the derived posteriori monthly SFs on waste treatment reflected changes on emissions, considering the monthly activity data does not have obvious monthly changes, these SFs can mainly reflect relative variations of monthly EFs and contain meteorological dominated changes especially for temperature. To evaluate the temperature sensitivity of its EFs, we first calculated the normalized monthly SFs by dividing monthly SFs by annual averages (Tables 1 and S3), and quantified the relationship between observed T_{2m} and normalized SFs. Note decomposition of organic waste by methanogens mostly takes at depth within the landfills and temperature can be higher than at the surface, hence the temperature within landfills should be much more related to methanogens activities and CH₄ emissions than T_{2m}. However, considering (1) we do not have direct temperature observations under landfills, (2) T_{2m} can be used as indicator of methanogens activities, and (3) T_{2m} is commonly used meteorological data that can be provided for future RCP scenarios, hence the relationship between waste CH₄ emissions and T_{2m} is constructed and used to predict how will CH₄ EFs change in different climate scenarios. The normalized SFs illustrate significant linear relationship with monthly T_{2m} (Figure 7b), where the slopes imply that normalized SFs (and EFs) will increase by 38%~50% with temperature increase by 10°C at city scale. We also analyzed the temperature sensitivity by only using daytime CH₄ observations and simulations in Figure S10, it still shows strong linear relationship between normalized SFs and T_{2m}, with the slopes of 0.046 and 0.060. These results are in high consistency with using all-day observations of 0.038 and 0.050, indicating similar results of using 24 hours observations and only

using daytime observations, and less influence of simulated nighttime PBLH bias on corresponding temperature sensitivity.

We should note the precipitation, soil water content and atmospheric pressure can also have obvious influence on CH_4 emissions, and considering the fact that we have not conducted field measurement in landfills and landfills are usually covered by metal or plastic in China to avoid the spread of odors, hence reanalysis data cannot represent real soil water contents in these site scale landfills. Precipitation and atmospheric pressure show obvious linear relationship with temperature as displayed in Figure S11. They display positive linear relationship between precipitation (affect water content) and T_{2m} , and negative linear relationship between monthly averaged atmospheric pressure and T_{2m} . We also found negative relationship between atmospheric pressure and normalized SFs, and positive relationship between T_{2m} and normalized SFs (Figures 7b and S11). Considering air temperature always displays negative relationship with atmospheric pressure as warmer air temperature coincides with lighter air mass and lower atmospheric pressure in summer as displayed in Figure 11b, and colder air temperature coincides with heavier air mass and higher atmospheric pressure in winter. Hence, the temperature can be used to represent co-influence of both temperature and atmospheric pressure, and we only focus on the influence of temperature on CH_4 emissions and will add more supporting data in following studies.

Our findings for the high sensitivity of waste treatment CH₄ emissions to temperature also suggest a dramatic increase with the projection of future global warming trends. We further derived the T_{2m} trends for four different RCP scenarios as RCP8.0, RCP6.0, RCP4.5 and RCP2.6 (Figure 8a). The results show T_{2m} will increase by 0.50°C, 0.28°C, 0.16°C, 0.10°C per decade for Hangzhou city, respectively. These different warming trends also indicate distinct temperature-dominated influence on future CH₄ EFs and emissions from waste treatment. We then used the slopes from Figure 7b and annual temperature from 2021 to 2100 to derive relative changes of EFs in future 80 years, where observations for year 2021 were treated as the baseline year. As displayed in Figure 8b, the EFs in RCP8.5, RCP6.0, RCP4.5 and RCP2.6 scenarios will increase with the rates of 2.2%, 1.2%, 0.7% and 0.5% per decade, respectively. And CH₄ EFs for waste treatment will be

higher by 17.6%, 9.6%, 5.6%, and 4.0% at the end of this century.

616

617

618

619

620

621

622

623

624

625

626

627

628

629

630

631

632

633

634

635

636

637

638

615

The spatial distribution of T_{2m} trends for all of China is also displayed in Figure S12, which shows heterogeneous distribution across China for four global warming scenarios. Because East China has high population density, with the majority of the national population (Figure S1), and is responsible for the largest domestic garbage induced CH₄ emissions (Figure S2), these combined factors indicate considerable CH₄ emissions changes from waste treatment in such a temperature-sensitivity area. Considering that the temperature sensitivity of waste treatment CH₄ EFs is caused by microbial process at regional scales, the sensitivity can represent general conditions of different cities or landfills. And if we assume the derived temperature sensitivity (increase by 44% with temperature increases of 10°C on average) is applicable for China as a whole, especially for East China, the relative changes of waste treatment CH₄ EFs can be calculated by multiplying this value by air temperature trends. The spatial distribution of global warming induced EF changes at the end of this century is displayed Figure 9. For RCP2.6 scenario, EFs for waste treatment will slightly increase by 4.0-6.5% in the north eastern China and increase by 3.0-4.0% in south eastern China. The RCP6.0 also displayed heterogeneous changes in East China, with EFs in the north eastern China increasing by 10.5-13.0% and in south eastern China increasing by 9.0-10.5%. Relative changes in RCP4.5 and RCP8.5 are more homogeneous for East China, which indicates EFs will significantly increase by 5.0-7.5% and 17.5-19.5%, respectively. The largest changes will occur in West China for RCP8.5, with EFs increasing by >20.0%, but this area has low population density and CH₄ emissions, and therefore these effects of global warming can be ignored (Figure S12). Finally, we should note these derived relative changes are only caused by global warming, and the influence of activity data, management technology and other factors is not considered and out of the scope of this study.

639

640

641

642

643

4 Discussions and implications

Many previous studies have compared total CH₄ emissions and its components for different inventories and bottom-up methods, which illustrated large uncertainty and bias at city scale and these biases were much larger for waste treatment (Peng at al., 2016; Saunois et al., 2020; Lin et

al., 2021; Bian et al., 2022). A recent bottom-up research compared wastewater CH₄ EFs in China, which largely varied by four-fold in different provinces and the uncertainty in the same province were even two-fold larger than its average, implying considerable uncertainty in recent understanding of waste treatment EFs at regional scale (Hua et al., 2022). And for the national total emissions, waste treatment CH₄ emissions varied between 5 and 15 Tg a⁻¹ (Peng et al., 2016; EDGAR v6). There are also other atmospheric inversion studies in estimating China's CH₄ emissions (Hopkins et al., 2016; Hu et al., 2019; Huang et al., 2021; Miller et al., 2019; Lu el., 2021; Chen et al., 2022). These studies found large variations of national emissions for almost all inventories, which were mainly caused by fossil fuel exploitation, agricultural sector (livestock and rice paddies) and waste treatment. For the comparisons of waste treatment emissions, these satellite-based inversions also largely varied between 6 and 9 Tg a⁻¹ by 1.5-fold (Miller et al., 2019; Lu et al., 2021; Chen et al., 2022; Zhang et al., 2022).

The reported discrepancies between "bottom-up" and "top-down" approaches indicate large uncertainty in understanding China's national CH₄ emissions from waste treatment. And it is well known the uncertainties will increase from national scale to regional and city scales, also implying considerable uncertainties in city-scale emissions for inventories. But the atmospheric inversion approach for city scale waste treatment, which can act as an independent evaluation, is still rare not only for China but also globally. To our best knowledge, there is only one recent atmospheric inversion research focused on CH₄ emissions from city-scale waste treatment, which used satellite-based observation to constrain emissions from four cities in India and Pakistan, that concluded underestimation of landfills CH₄ emissions by 1.4 to 2.6 times for EDGAR inventory (Maasakkers et al., 2022). In our study, we found annual waste CH₄ emissions were overestimated by 47.1% for Hangzhou city, our findings are different from results in India and Pakistan. These differences indicate bias of waste treatment CH₄ emissions considerably varied in different countries and climate divisions. Our results highlight there is a large knowledge gap in understanding waste treatment emissions mechanisms and estimating urban waste treatment CH₄ emissions especially in China.

Different from fossil-type sources that have much smaller monthly variations, CH₄ emission from waste treatment is biological processes-based source and its EFs are highly sensitive to meteorological conditions especially for temperature. These factors lead to obvious bias in waste treatment CH₄ emissions not only for annual average but also for its seasonality. Besides, although there were a few studies that aimed to predict future CH₄ emissions from waste treatment, these studies were mainly based on activity data changes without considering the EFs variations caused by future global warming trends or only based on site-specific observations (USEPA 2013; Cai et al., 2018; Spokas et al., 2021). Of these three cited studies, USEPA (2013) and Cai et al. (2018) only predicted emissions changes due to changes in activity data and management technology. And the CH₄ emissions for year 2030 by Cai et al. (2018) was 23.5% lower than the USEPA (2013) estimation, which was caused by the consideration of new policies and low-carbon policy scenarios. Spokas et al. (2021) modeled the CH₄ emission changes with increasing air temperature, where CH₄ emissions did not show obvious changes even with temperature increasing by ~5°C by the end of year 2100. To our best knowledge, there are no inventories that considered the temperature-induced changes on both seasonal variations and annual trends of CH₄ emissions. Hence, it is still unclear in all inventories how EFs will change with different global warming scenarios at city scale.

A few observation-based measurements were conducted for waste treatment but only at some specific sites with large discrepancies of EFs (Du et al., 2017; 2018; Cai et al., 2018; Zhao et al., 2019; NBSC, 2015; Wang et al., 2015; Florentino et al., 2010; Tolaymat et al., 2010; Cai et al., 2014; 2018). And only one of our previous studies used year-round atmospheric CH₄ observations to constrain regional scale CH₄ emissions at Nanjing city in YRD area (Huang et al., 2021), where it found much higher emissions of the landfilling waste in summer than in winter: CH₄ emissions in July were around four times those in February. But there is no study that has quantified the temperature sensitivity of waste CH₄ emissions at city scale or much larger regional scales. These two studies in different cities confirmed temperature as the dominant factor that drives seasonal variations of waste treatment CH₄ emissions. Hence our study appears as the first one that estimated city scale waste treatment CH₄ emissions, its temperature sensitivity and projected

changes in different global warming scenarios. Our findings for the large sensitivity to temperature indicate the monthly scaling factors should be considered to better represent CH₄ emissions and simulate atmospheric CH₄ concentrations.

We also note that the predictions of future climate changes are mainly based on different emitting intensity of greenhouse gas, and CH₄ contributed around 20% of direct anthropogenic radiative forcing (Seto et al., 2014). The CH₄ emissions in different global warming scenarios were mainly calculated by predicting energy use data without considering the changes of EFs. In this study, we found there should be large positive feedback between global warming and CH₄ emissions, especially in the RCP 8.0 scenario where global warming induced CH₄ emissions from waste treatment will increase by 17.6%. Hence the projected emissions from waste treatments and other biological process based sources, together with positive feedback between temperature and their emissions are strongly suggested in future climate change models. Besides, it is well known that CH₄ concentration simulations are essential for modeling air pollutions (e.g. O₃, NO_x, and CO) especially in the stratosphere (Isaksen et al., 2011; Kaiho et al., 2013). Considering that waste treatment CH₄ emissions accounted for ~25% of total anthropogenic emissions (EDGAR v6.0) in East China where severe air pollution frequently occurred, we also believe the coupling of temperature-dependent CH₄ emissions and the monthly scaling factors on CH₄ emissions can improve air pollution modeling in East China.

We should note that new technology and other meteorological variables can also influence waste treatment CH₄ emissions. The main reason to only use temperature in this study is that we only constrained the emissions at monthly scale in one year, and derived twelve datasets of *posteriori* CH₄ emissions. Besides, temperature is considered to be the main factor in controlling monthly and annual variations of waste treatment CH₄ emissions, and can be used to represent the co-influence of other meteorological parameters such as atmospheric pressure. We will use multiple years' CH₄ concentration to quantify the influence of new technology and other meteorological variables on waste treatment CH₄ emissions in our following study, and we suggest that other tracers (e.g. ethane, ¹⁴CH₄) are also important to separate CH₄ emissions from biological

and fossil CH₄ emissions.

732

733

734

735

736

737

738

739

740

741

731

5 Summary and Conclusions

To better evaluate bias for city scale anthropogenic CH₄ emissions and understand the sensitivity of temperature on waste treatment CH₄ emissions, we used a three tower-based atmospheric CH₄ observation network in Hangzhou city, which is located in the developed YRD region and one of the top 10 megacities in China. One-year hourly atmospheric CH₄ observations were presented from December 2020 to November 2021. We then applied a scaling factor Bayesian inversion method to constrain monthly anthropogenic CH₄ emissions and its components (especially for waste treatments) in Hangzhou city, and also used multiplicative scaling factor method for broader Zhejiang province and YRD area at seasonal scale.

742

743

744

745

746

747

748

749

750

751

752

753

754

755

756

757

758

759

To the best of our knowledge, our study is the first tower-based CH₄ observation network in China. We found obvious seasonal bias of simulated CH₄ concentrations at the core urban area of Hangzhou city, which was mainly caused by bias of waste treatment at both annual and monthly scales. The derived posteriori CH₄ emissions display obvious seasonal variations with peak in summer and trough in winter, which was mainly contributed by waste treatment; the a priori annual waste treatment CH₄ emission in Hangzhou city was 10.4×10⁴ t and decreased to 5.5 $(\pm 0.6) \times 10^4$ t for the *posteriori* emissions, a decrease of 47.1%. Besides, the total anthropogenic CH₄ emissions (excluding agricultural soil) decreased from 15.0×10^4 t to $9.6(\pm 0.9) \times 10^4$ t, indicating overestimation of 36.0% for the whole year of 2021. Observations at Linan site imply that the annual CH₄ emissions was slightly underestimated by 7.0% for the larger region of Zhejiang province or YRD area, which was different from the case of Hangzhou city. Additionally, the posteriori monthly CH₄ emissions from waste treatment illustrate significant linear relationship with air temperature, with regression slopes indicating an increase of 38%~50% when temperature increases by 10°C. Finally, we found the waste treatment CH₄ EFs for Hangzhou city will increase by 17.6%, 9.6%, 5.6%, and 4.0% by the end of this century for RCP8.0, RCP6.0, RCP4.5 and RCP2.6 scenarios, respectively. The derived relative changes for whole China also showed high heterogeneity and indicate large uncertainty in projecting future national total CH₄

- 760 emissions. This study is also the first one that mainly focuses on city scale temperature sensitivity
- of waste treatment CH₄ emissions from the perspective of atmospheric inversion approach. And
- based on above results, we strongly suggest the temperature-dependent EFs should be coupled in
- both recent CH₄ inventories and future CH₄ emission projections.

- 765 Data availability: The atmospheric CH₄ observations data can be requested from Cheng Hu and
- 766 Bing Qi. STILT model is downloaded from http://www.stilt-model.org/, the EDGAR inventory is
- from https://edgar.jrc.ec.europa.eu/, and the projected climate data were downloaded from World
- Data Center for Climate (WDCC, https://www.wdc-climate.de/ui/).

769

- 770 Acknowledgement: Cheng Hu is supported by the National Natural Science foundation of China
- 771 (grant no. 42105117) and Natural Science Foundation of Jiangsu Province (grant no. BK20200802).
- Wei Xiao is supported by the National Key R&D Program of China (grants 2020YFA0607501 &
- 773 2019YFA0607202). This work is also supported by Zhejiang Provincial Basic Public Welfare Research
- Project (LGF22D050004). We sincerely thank the detailed comments from two anonymous reviewers.
- We also want to express our thanks to Prof. Timothy J. Griffis from University of Minnesota, who
- provided many important suggestions and support for this study.

777

- 778 **Author contribution:** Cheng Hu and Bing Qi designed the study. Cheng Hu performed the model
- simulation, data analysis and wrote and revised the paper; Bing Qi and Rongguang Du conducted CH₄
- 780 concentration observation and meteorological data collection, and all co-authors contributed to the
- 781 data/figures preparation and analysis.
- Declaration of competing interests: The authors declare that they have no conflict of interest.

783

- 784 **References:**
- Agustí-Panareda, A., Diamantakis, M., Massart, S., Chevallier, F., Muñoz-Sabater, J., Barré, J., Curcoll, R.,
- Engelen, R., Langerock, B., Law, R. M., Loh, Z., Morguí, J. A., Parrington, M., Peuch, V.-H., Ramonet, M., Roehl,
- 787 C., Vermeulen, A. T., Warneke, T., and Wunch, D.: Modelling CO₂ weather why horizontal resolution matters,
- 788 Atmos. Chem. Phys., 19, 7347–7376, https://doi.org/10.5194/acp-19-7347-2019, 2019.

789

- Bian R., Zhang T., Zhao F., et al. Greenhouse gas emissions from waste sectors in China during 2006–2019:
- 791 Implications for carbon mitigation. Process. Saf. Environ., 161:488-497, 2022.

- 793 Bloom, A. A., Bowman, K. W., Lee, M., Turner, A. J., Schroeder, R., Worden, J. R., Weidner, R., McDonald, K. C.,
- and Jacob, D. J.: A global wetland methane emissions and uncertainty dataset for atmospheric chemical transport
- 795 models (WetCHARTs version 1.0), Geosci. Model Dev., 10, 2141-2156,
- $796 \qquad https://doi.org/10.5194/gmd-10-2141-2017, \, 2017.$

- 797 Börjesson G, Svensson BH. Seasonal and Diurnal Methane Emissions From a Landfill and Their Regulation By
- 798 Methane Oxidation. Waste Management & Research. 1997;15(1):33-54. doi:10.1177/0734242X9701500104

- 800 Cai, B., J. Liu, X. Zeng, D. Cao, L. Liu, Y. Zhou, Z. Zhang, Estimation of CH₄ emission from landfill in China
- based on point emission sources. Adv. Clim. Change Res. 5, 81–91, 2014.

802

- and related environmental co-benefits, Sci. Adv., 4, eaar8400, https://doi.org/10.1126/sciadv.aar8400, 2018.

805

- 806 Chen, Z., Jacob, D. J., Nesser, H., Sulprizio, M. P., Lorente, A., Varon, D. J., Lu, X., Shen, L., Qu, Z., Penn, E., and
- 807 Yu, X.: Methane emissions from China: a high-resolution inversion of TROPOMI satellite observations, Atmos.
- 808 Chem. Phys., 22, 10809–10826, https://doi.org/10.5194/acp-22-10809-2022, 2022.

809

- Du, M., Peng, C., Wang, X., Chen, H., Wang, M., and Zhu, Q.: Quantification of methane emissions from
- municipal solid waste landfills in China during the past decade, Renew. Sust. Energ. Rev., 78, 272–279, 2017.

812

- Du, M., Zhu, Q., Wang, X., Li, P., Yang, B., Chen, H., Wang, M., Zhou, X., and Peng, C.: Estimates and
- predictions of methane emissions from wastewater in China from 2000 to 2020, Earths Future, 6, 252–263, 2018.

815

- Fang S.X., R.G. Du, B. Qi. et al., Variation of carbon dioxide mole fraction at a typical urban area in the Yangtze
- 817 River Delta, China. Atmos. Res, 265, 105884, 2022.

818

- 819 Florentino, Cruz., B. De La, and M. A. Barlaz., Estimation of waste component-specific landfill decay rates using
- laboratory-scale decomposition data. Environ. Sci. Technol. 44, 4722–4728, 2010.

821

- 822 Griffis, T. J., Chen, Z., Baker, J. M., Wood, J. D., Millet, D. B., Lee, X., et al., Nitrous oxide emissions are
- 823 enhanced in a warmer and wetter world. P. Natl. Acad. Sci. USA, 114(45), 12081-12085.
- 824 https://doi.org/10.1073/pnas.1704552114, 2017.
- He, J., Naik, V., Horowitz, L. W., Dlugokencky, E., and Thoning, K.: Investigation of the global methane budget
- 826 over 1980–2017 using GFDL-AM4.1, Atmos. Chem. Phys., 2020, 20, 805–827,
- 827 https://doi.org/10.5194/acp-20-805-2020.

828

- Henne, S., Brunner, D., Oney, B., Leuenberger, M., Eugster, W., Bamberger, I., Meinhardt, F., Steinbacher, M., and
- 830 Emmenegger, L.: Validation of the Swiss methane emission inventory by atmospheric observations and inverse
- 831 modelling, Atmos. Chem. Phys., 16, 3683–3710, https://doi.org/10.5194/acp-16-3683-2016, 2016.

832

- Hopkins, F. M., Kort, E. A., Bush, S. E., Ehleringer, J. R., Lai, C.- T., Blake, D. R., & Randerson, J. T. Spatial
- patterns and source attribution of urban methane in the Los Angeles Basin. J. Geophys. Res-Atmos., 121, 2490-
- 835 2507, 2016.

836

- Höglund-Isaksson, L.: Global anthropogenic methane emissions 2005–2030: technical mitigation potentials and
- 838 costs, Atmos. Chem. Phys., 12, 9079–9096, https://doi.org/10.5194/acp-12-9079-2012, 2012.

- 840 Hua, H., Jiang, S., Yuan, Z., Liu, X., Zhang, Y., & Cai, Z. Advancing greenhouse gas emission factors for
- 841 municipal wastewater treatment plants in China. Environ. Pollut., 295, 118648.
- 842 https://doi.org/10.1016/j.envpol.2021.118648, 2022.

- $Hu\ C,\ Griffis,\ T.\ J.,\ Liu,\ S.,\ Xiao,\ W.,\ Hu,\ N.,\ Huang,\ W.,\ Yang,\ D.,\ Lee,\ X.,\ Anthropogenic\ methane\ emission\ and$
- its partitioning for the Yangtze River Delta region of China. J. Geophys.l Res-Biogeo., 124(5): 1148-1170, 2019.

846

- Hu, C., Xu, J., Liu, C., Chen, Y., Yang, D., Huang, W., Deng, L., Liu, S., Griffis, T. J., and Lee, X.: Anthropogenic
- 848 and natural controls on atmospheric δ13C-CO2 variations in the Yangtze River delta: insights from a carbon
- 849 isotope modeling framework, Atmos. Chem. Phys., 21, 10015–10037, https://doi.org/10.5194/acp-21-10015-2021,
- 850 2021.

851

- Hu, C., Griffis, T.J., Xia, L., Xiao, W., Liu, C., Xiao, Q., Huang, X., Yang, Y., Zhang, L., Hou, B., Anthropogenic
- 853 CO₂ emission reduction during the COVID-19 pandemic in Nanchang City, China, Environ. Pollut., 309, 119767,
- doi: https://doi.org/10.1016/j.envpol.2022.119767, 2022.
- Huang, W. J., T. J. Griffis, C. Hu, W. Xiao, and X. H. Lee. Seasonal variations of CH₄ emissions in the
- 856 Yangtze River Delta region of China are driven by agricultural activities. Adv. Atmos. Sci., 38(9), 1537–1551,
- 857 https://doi.org/10.1007/s00376-021-0383-9, 2021.

858

- Isaksen I S, Gauss M, Myhre G, Anthony W, Katey M and Ruppel C 2011 Strong atmospheric chemistry feedback
- to climate warming from Arctic methane emissions. Global Biogeochem. Cy. 25 GB2002, 2011.

861

- 862 Kumar, P.; Broquet, G.; Caldow, C.; et al. Near-field atmospheric inversions for the localization and quantification
- 863 Of controlled methane releases using stationary and mobile measurements. Q. J. R. Meteorol. Soc. 2022, 148,
- 864 1886-1912

865

- 866 Kissas K, Ibrom A, Kjeldsen P, et al. Methane emission dynamics from a Danish landfill: The effect of changes
- in barometric pressure. Waste Management, 2022, 138:234-242.

868

- 869 Lian, J., Bréon, F.-M., Broquet, G., Lauvaux, T., Zheng, B., Ramonet, M., Xueref-Remy, I., Kotthaus, S.,
- Haeffelin, M., and Ciais, P.: Sensitivity to the sources of uncertainties in the modeling of atmospheric CO₂
- concentration within and in the vicinity of Paris, Atmos. Chem. Phys., 21, 10707–10726,
- 872 https://doi.org/10.5194/acp-21-10707-2021, 2021.

873

- 874 Lin, X., Zhang, W., Crippa, M., Peng, S., Han, P., Zeng, N., Yu, L., and Wang, G.: A comparative study of
- anthropogenic CH₄ emissions over China based on the ensembles of bottom-up inventories, Earth Syst. Sci. Data,
- 876 13, 1073–1088, https://doi.org/10.5194/essd-13-1073-2021, 2021.

877

- 878 Lopez-Coto, I., Ren, X., Salmon, O. E., Karion, A., Shepson, P. B., Dickerson, R. R., Stein, A., Prasad, K., and
- Whetstone, J. R.: Wintertime CO₂, CH₄, and CO Emissions Estimation for the Washington, DC-Baltimore
- Metropolitan Area Using an Inverse Modeling Technique, Environmental Science and Technology, 54, 2606–2614,
- 881 https://doi.org/10.1021/acs.est.9b06619, 2020.

882

Lou, Z., Cai, B.F., Zhu, N., Zhao, Y., Geng, Y., Yu, B., Chen, W., Greenhouse gas emission inventories from waste

- 884 sector in China during 1949-2013 and its miti- gation potential. J. Clean. Prod. 157, 118-124.
- 885 https://doi.org/10.1016/j.jclepro. 2017.04.135, 2017.

- Lu, X., Jacob, D. J., Zhang, Y., Maasakkers, J. D., Sulprizio, M. P., Shen, L., Qu, Z., Scarpelli, T. R., Nesser, H.,
- 888 Yantosca, R. M., Sheng, J., Andrews, A., Parker, R. J., Boesch, H., Bloom, A. A., and Ma, S.: Global methane
- budget and trend, 2010-2017: com- plementarity of inverse analyses using in situ (GLOBALVIEW- plus CH4
- ObsPack) and satellite (GOSAT) observations, At- mos. Chem. Phys., 21, 4637–4657, https://doi.org/10.5194/acp-
- 891 21-4637-2021, 2021.

892

- 893 Kaiho K., Koga S. Impacts of a massive release of methane and hydrogen sulfide on oxygen and ozone during the
- 894 late Permian mass extinction. Global Planetary Change, 107:91-101,
- https://doi.org/10.1016/j.gloplacha.2013.04.004, 2013.

896

- Maasakkers, J. D., Varon, D. J., Elfarsdóttir, A., McKeever, J., Jervis, D., Mahapatra, G., Pandey, S., Lorente, A.,
- Borsdorff, T., Foorthuis, L. R., Schuit, B. J., Tol, P., van Kempen, T. A., van Hees, R., & Aben, I. Using satellites to
- 899 uncover large methane emissions from landfills. Sci. Adv. 8, eabn9683, 10.
- 900 https://doi.org/10.1126/sciadv.abn9683, 2022.

901

- 902 Masuda, S., Sano, I., Hojo, T., Li, Y., Nishimura, O., The comparison of greenhouse gas emissions in sewage
- treatment plants with different treatment processes. Chemosphere 193, 581–590, 2018.

904

- 905 Miles, N. L., Richardson, S. J., Lauvaux, T., Davis, K. J., Balashov, N. V., Deng, A., Turnbull, J. C., Sweeney, C.,
- Gurney, K. R., Patarasuk, R., Razlivanov, I., Cambaliza, M. O. L. and Shepson, P. B.: Quantification of urban
- atmospheric boundary layer greenhouse gas dry mole fraction enhancements in the dormant season: Results from
- the Indianapolis Flux Experiment (INFLUX), Elem Sci Anth, 5, 27, doi:10.1525/elementa.127, 2017.

909

- 910 Miller, S. M., Matross, D. M., Andrews, A. E., Millet, D. B., Longo, M., Gottlieb, E. W., Hirsch, A. I., Gerbig, C.,
- 911 Lin, J. C., Daube, B. C., Hudman, R. C., Dias, P. L. S., Chow, V. Y., and Wofsy, S. C.: Sources of carbon monoxide
- 912 and formaldehyde in North America determined from high-resolution atmospheric data, Atmos. Chem. Phys., 8,
- 913 7673–7696, https://doi.org/10.5194/acp-8-7673-2008, 2008.

914

- 915 Miller, S. M., Michalak, A. M., Detmers, R. G., Hasekamp, O. P., Bruhwiler, L. M. P., & Schwietzke, S. China's
- 916 coal mine methane regulations have not curbed growing emissions. Nature Communications, 10(1), 303-308.
- 917 https://doi.org/10.1038/s41467-018-07891-7, 2019.
- 918 Mønster, J., Kjeldsen, P. and Scheutz, C. (2019) Methodologies for measuring fugitive methane emissions from
- 919 landfills a review. In Waste Management., 87, 835–859. https://doi.org/10.1016/j.wasman.2018.12.047.
- 920 National Bureau of Statistics of China (NBSC), China Statistical Yearbook (China Statistics Press, 2015) (in
- 921 Chinese).

922

- 923 Pak N M, Heerah S, Zhang J, et al. The Facility Level and Area Methane Emissions inventory for the Greater
- Toronto Area (FLAME-GTA)[J]. Atmospheric Environment, 2021, 252(9):118319.

925

926 Peng, S., Piao, S., Bousquet, P., Ciais, P., Li, B., Lin, X., Tao, S., Wang, Z., Zhang, Y., and Zhou, F.: Inventory of

- anthropogenic methane emissions in mainland China from 1980 to 2010, Atmos. Chem. Phys., 16, 14545–14562,
- 928 https://doi.org/10.5194/acp-16-14545-2016, 2016.

- 930 Sargent, M., Barrera, Y., Nehrkorn, T., Hutyra, L. R., Gately, C. K., Mckain, K., Sweeney, C., Hegarty, J.,
- 931 Hardiman, B., Steven C. Wofsy, S. C.: Anthropogenic and biogenic CO₂ fluxes in the Boston urban region, P. Natl.
- 932 Acad. Sci. USA., 115(40), https://doi.org/10.1073/pnas.1803715115, 2018.

933

- 934 Saunois, M., Stavert, A. R., Poulter, B., et al., The Global Methane Budget 2000–2017, Earth Syst. Sci. Data, 12,
- 935 1561–1623, https://doi.org/10.5194/essd-12-1561-2020, 2020.

936

- 937 Seto, K. C. hakal, S. Bigio, A. Blanco, H. elgado, G. C. ewar, Huang, L. Inaba, A. Kansal, A. Lwasa, S. cahon, J.
- 938 Iler, B. urakami, J. Nagendra, H. amaswami, A. Humansettlements, infrastructure and spatial planning.Climate
- Change 2014:Mitigation of Climate Change. IPCC Working Group III Contribution to AR5; Cambridge University
- 940 Press, 2014; Chapter 12.

941

- 942 Solazzo, E., Crippa, M., Guizzardi, D., Muntean, M., Choulga, M., and Janssens-Maenhout, G.: Uncertainties in
- 943 the Emissions Database for Global Atmospheric Research (EDGAR) emission inventory of greenhouse gases,
- 944 Atmos. Chem. Phys., 21, 5655–5683, https://doi.org/10.5194/acp-21-5655-2021, 2021.

945

- 946 Spokas, K.A., et al. 2021. Modeling landfill CH4 emissions: CALMIM international fieldvalidation, using
- 947 CALMIM to simulate management strategies, current and futureclimate scenarios. Elem Sci Anth, 9: 1.
- 948 https://doi.org/10.1525/elementa.2020.00050Do, 2020.

949

- Tolaymat, T., M., R. B. Green, G. R. Hater, M. A. Barlaz, P. Black, D. Bronson, J. Powell, Evaluation of landfill
- 951 gas decay constant for municipal solid waste landfills operated as bioreactors. J. Air Waste Manage. Assoc. 60, 91–
- 952 97, 2010.

953

- Thoning, K. W., Tans, P. P., and Komhyr, W. D.: Atmospheric carbon dioxide at Mauna Loa observatory 2.
- 955 Analysis of the NOAA/GMCC data, 1974–1985, J. Geophys. Res.-Atmos., 94, 8549–
- 956 8565, https://doi.org/10.1029/JD094iD06p08549, 1989.
- 957 Tian, J., Gong, Y., Li, Y., Chen, X., Zhang, L., & Sun, Y. (2022). Can policy implementation increase public waste
- sorting behavior? The comparison between regions with and without waste sorting policy implementation in China.
- Journal of Cleaner Production, 132401.

- 961 United States Environmental Protection Agency (USEPA), Global Mitigation of Non-CO₂ Greenhouse Gases
- 962 2010-2030 (United States Environmental Protection Agency Office of Atmospheric Programs (6207J),
- 963 EPA-430-R-13-011, 2013);
- 964 www.epa.gov/sites/production/files/2016-07/documents/mac report 2014-exec summ.compressed.pdf
- Verhulst, K. R., Karion, A., Kim, J., Salameh, P. K., Keeling, R. F., Newman, S., Miller, J., Sloop, C., Pongetti, T.,
- Rao, P., Wong, C., Hopkins, F. M., Yadav, V., Weiss, R. F., Duren, R. M. and Miller, C. E.: Carbon dioxide and
- 967 methane measurements from the Los Angeles Megacity Carbon Project Part 1: calibration, urban enhancements,
- 968 and uncertainty 10 estimates, Atmos. Chem. Phys., 17(13), 8313–8341, doi:10.5194/acp-17-8313-2017, 2017

- Wang, X., A. S. Nagpure, J. F. DeCarolis, M. A. Barlaz, Characterization of uncertainty in estimation of methane
- ollection from select U.S. landfills. Environ. Sci. Technol. 49, 1545–1551, 2015.

- 972 Wang, Y., Wang, X., Wang, K. et al. The size of the land carbon sink in China. Nature, E7-E9.
- 973 https://doi.org/10.1038/s41586-021-04255-y, 2022.

974

- 975 Williams, J. P., Ars, S., Vogel, F., Regehr, A., & Kang, M. (2022). Differentiating and Mitigating Methane
- 976 Emissions from Fugitive Leaks from Natural Gas Distribution, Historic Landfills, and Manholes in Montréal,
- 977 Canada. Environmental Science & Technology. https://doi.org/10.1021/acs.est.2c06254

978

- 979 Yadav, V., Duren, R., Mueller, K., Verhulst, K. R., Nehrkorn, T., and Kim, Jet., Spatio-temporally resolved
- 980 methane fluxes from the Los Angeles megacity J. Geophys. Res. Atmos. 124, 5131–5148 (2019).

981

- Zhao, X., Jin, X., Guo, W., Zhang, C., Shan, Y., Du, M., Tillotson, M., Yang, H., Liao, X., and Li, Y.: China's urban
- methane emissions from municipal wastewater treatment plant, Earths Future, 7, 480–490, 2019.

984

- 285 Zhao, Z., Bian, R., Zhao, F., Chai, X., Implications of municipal solid waste disposal methods in China on
- 986 greenhouse gas emissions. Renew. Sust. Energ. Rev. 39 (3). https://doi.org/10.1002/ep.13372, 2019.

987

- 288 Zhang, B. and Chen, G.: China's CH₄ and CO₂ emissions: Bottomup estimation and comparative analysis, Ecol.
- 989 Indic., 47, 112–122, https://doi.org/10.1016/j.ecolind.2014.01.022, 2014.

990

- Zhang, K., Lee, X., Schultz, N. M., Huang, Q., Liu, Z., Chu, H., Zhao, L., & He, C. A global dataset on subgrid
- 992 land surface climate (2015-2100) from the Community Earth System Model. Geosci. Data J., 1-12.
- 993 https://doi.org/10.1002/gdj3.153, 2022.
- Zhang Y., Fang S., Chen J., Lin Y., Chen Y., Liang R., Jiang K., Parker R., Boesch H., Steinbacher M., Sheng J.,
- 995 Lu X., Shaojie Song, Shushi Peng: Observed Changes in China's Methane Emissions Linked to Policy Drivers,
- Proceedings of the National Academy of Sciences, 119, e2202742119, 2022.
- Povincial Bureau of Statistics, Survey Office of the National Bureau of Statistics in Zhejiang, Zhejiang
- 998 Statistical Yearbook 2018-2019 (China Statistics Press, Beijing, China, 2019)

999

1000

1001

1002

1003

1004

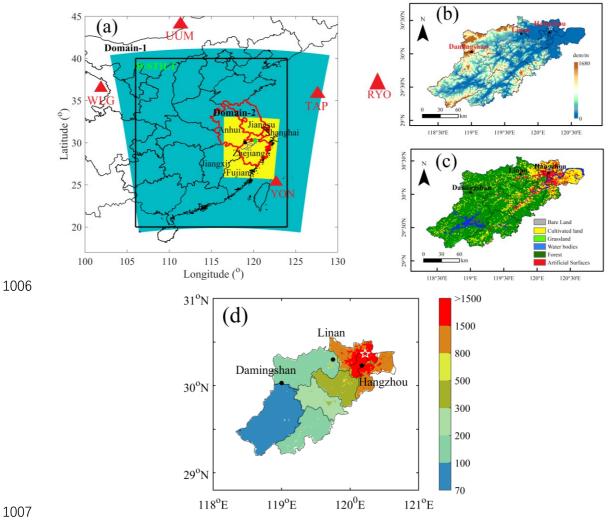


Figure 1. (a) WRF-STILT model domain setups, three CH₄ concentration observation sites in Hangzhou city, and five CH₄ background sites, note the green, red and black dots represent locations for Hangzhou site, Linan site and Damingshan site, respectively, Yangtze River Delta regions is displayed in red boundary, back rectangle represents domain in STILT model, (b) geophysical height within Hangzhou city, (c) land surface categories in Hangzhou city, and (d) population density in Hangzhou city for year 2019, units: person per km², the location of landfills in Hangzhou city is displayed with white star.

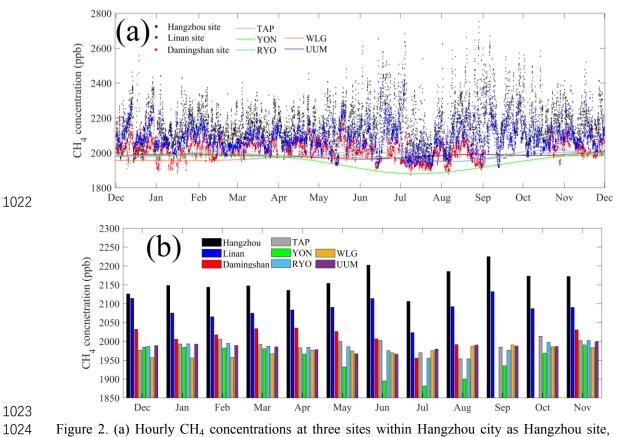


Figure 2. (a) Hourly CH₄ concentrations at three sites within Hangzhou city as Hangzhou site, Linan site, and Damingshan site, and fitting CH₄ background based on CCGCRV regression method at five background sites as TAP, YON, RYO, WLG and UUM, (b) monthly mean of CH₄ concentrations for above eight sites. Note the CH₄ background is smoothed by using CCGCRV fitting method on weekly or hourly observations, which can filter large fluctuations caused by sudden and unidentified sources

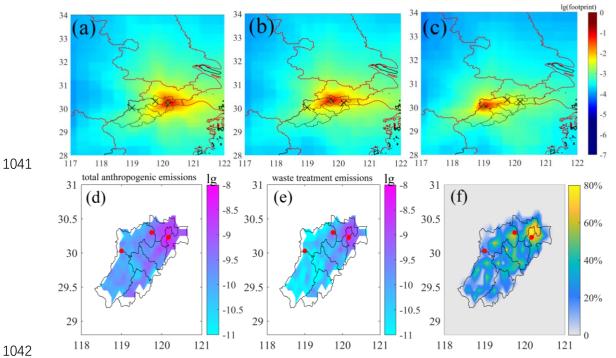
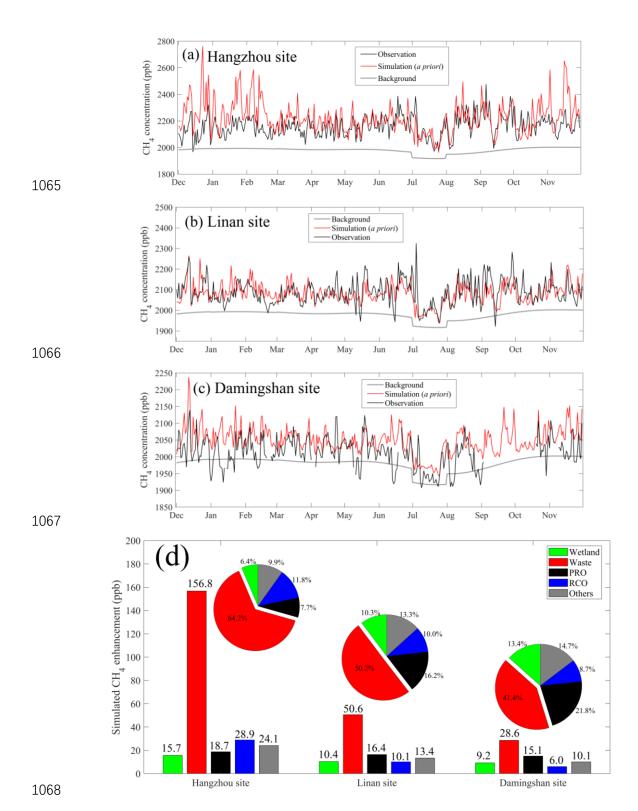


Figure 3. Annual averages of simulated footprint for (a) Hangzhou site, (b) Linan site, and (c) Damingshan site, where the green symbol "×" indicates receptor location in each pannel, (d) total anthropogenic CH₄ emissions in EDGAR v6.0 inventory, (e) waste treatment CH₄ emissions in EDGAR v6.0 inventory, and (f) proportions of waste treatment to total anthropogenic CH₄ emissions, red dot represents three sites, units for footprint: ppm m² s mol⁻¹, units for emissions: kg m⁻² s⁻¹. The divisions in Hangzhou city are different districts.



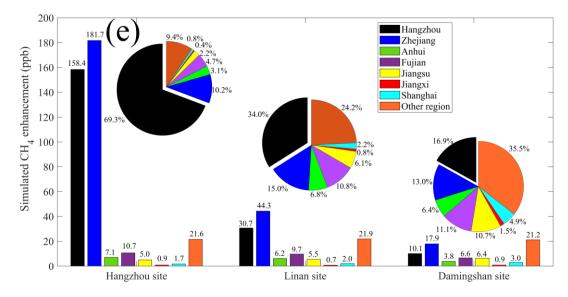


Figure 4. Comparisons between daily CH₄ observations and simulations for (a) Hangzhou site, (b) Linan site, (c) Damingshan site, (d) simulated CH₄ enhancements from main emission categories (e) simulated anthropogenic CH₄ enhancement from different regions and its proportions. Note the blue color for the bar charts include all contributions from "Zhejiang", including "Hangzhou"; and the blue regions in the pie charts represent rest regions of "Zhejiang minus Hangzhou".

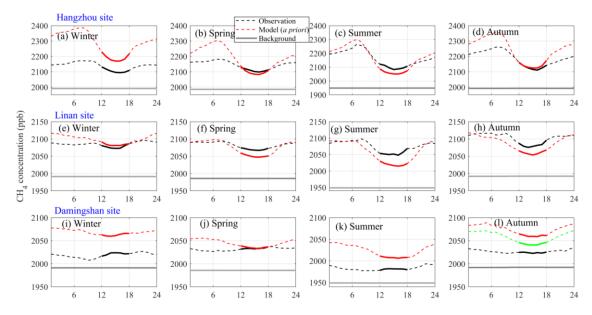


Figure 5. Seasonal averaged diurnal variations for Hangzhou site in (a) winter, (b) spring, (c) summer, (d) autumn, and Linan site in (e) winter, (f) spring, (g) summer, (h) autumn, and Damingshan site in (i) winter, (j) spring, (k) summer, (l) autumn; Note because of two months of data gap in Autumn for Damingshan site, the green line is for all September-November simulations, red line only represent simulation of corresponding period for available observation data, and bold lines represents data between 12:00 and 18:00.

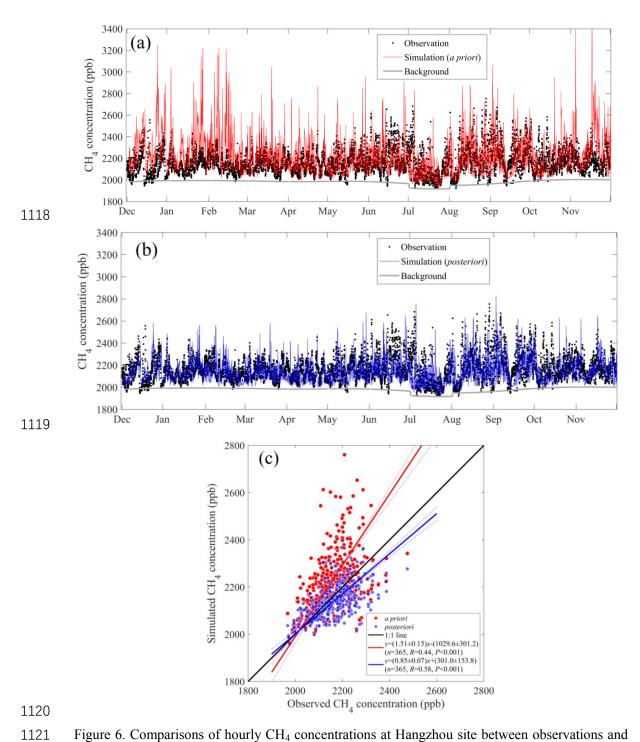


Figure 6. Comparisons of hourly CH₄ concentrations at Hangzhou site between observations and simulations by using (a) *a priori* and (b) *posteriori* emissions, (c) scatter plots of daily CH₄ averages by using *a priori* and *posteriori* emissions.

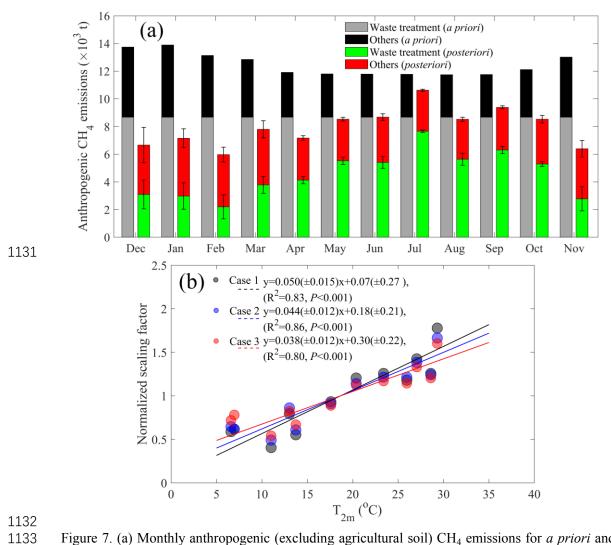


Figure 7. (a) Monthly anthropogenic (excluding agricultural soil) CH₄ emissions for *a priori* and *posteriori* emissions for Hangzhou city, (b) relationship between the monthly *posteriori* CH₄ emissions and temperature for the three cases discussed in section 2.3 of this text.

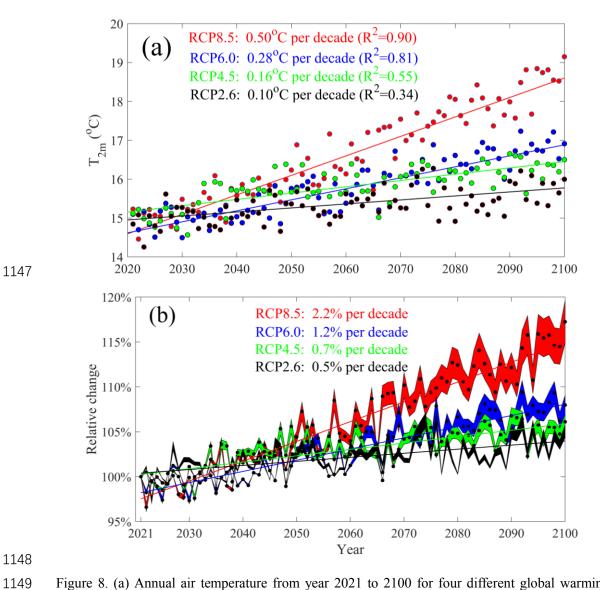


Figure 8. (a) Annual air temperature from year 2021 to 2100 for four different global warming scenarios for Hangzhou city, (b) the projected relative change of waste treatment CH₄ emissions (or EFs) for Hangzhou city, note the shading indicates extent of three cases.

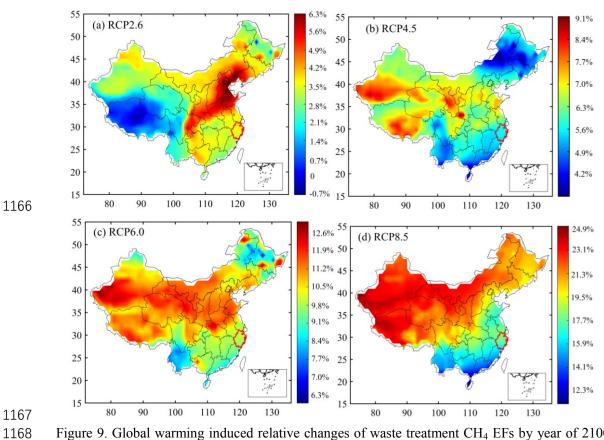


Figure 9. Global warming induced relative changes of waste treatment CH₄ EFs by year of 2100 for (a) RCP2.6, (b) RCP4.5, (c) RCP6.0, and (d) RCP8.5 scenarios. Note the red boundary is Zhejiang province.

Table 1. The *posteriori* SFs for different categories in three cases for Hangzhou city, where wetland: natural and agricultural wetland, Waste: waste treatment, PRO: fuel exploitation, RCO: energy for building, Others: the rest anthropogenic emissions. Note Case 1: 3 categories, and 300% uncertainty for waste treatment; Case 2: 5 categories; Case 3: 3 categories, and 200% uncertainty for waste treatment.

	Case 1			Case 2					Case 3		
Mont											
h	Wetland	Waste	Others	Wetland	Waste	PRO	RCO	Others	Wetland	Waste	Others
1	1.00	0.29	0.83	1.00	0.34	0.90	0.80	0.93	1.00	0.40	0.72
2	1.00	0.20	0.89	1.00	0.26	0.97	0.83	0.93	1.00	0.30	0.77
3	1.03	0.39	1.04	1.02	0.46	1.07	0.80	0.97	1.02	0.46	0.95
4	1.10	0.46	0.96	1.08	0.48	1.01	0.95	0.93	1.08	0.49	0.91
5	1.12	0.62	0.99	1.10	0.64	1.06	0.97	0.92	1.11	0.65	0.95
6	1.22	0.59	1.09	1.18	0.64	1.05	0.97	1.03	1.18	0.64	1.05
7	1.10	0.88	0.96	1.09	0.88	1.00	1.00	0.94	1.09	0.89	0.94
8	1.05	0.62	0.95	1.01	0.66	0.99	0.97	0.95	1.01	0.67	0.91
9	1.04	0.71	1.01	1.02	0.73	0.96	0.98	1.04	1.02	0.74	0.98
10	1.06	0.60	0.94	1.06	0.61	0.92	0.96	1.00	1.06	0.62	0.90
11	1.01	0.27	0.86	1.00	0.32	0.91	0.85	0.93	1.00	0.37	0.75
12	1.00	0.31	0.70	1.00	0.33	0.75	0.79	0.91	1.00	0.43	0.58