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To cite this article: Bo Zheng *et al* 2018 *Environ. Res. Lett.* **13** 044007

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LETTER

OPEN ACCESS

RECEIVED

10 October 2017

REVISED

20 February 2018

ACCEPTED FOR PUBLICATION

28 February 2018

PUBLISHED

23 March 2018

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Rapid decline in carbon monoxide emissions and export from East Asia between years 2005 and 2016

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Keywords: carbon monoxide, East Asia, emissions, decline

Supplementary material for this article is available [online](#)

Abstract

Measurements of Pollution in the Troposphere (MOPITT) satellite and ground-based carbon monoxide (CO) measurements both suggest a widespread downward trend in CO concentrations over East Asia during the period 2005–2016. This negative trend is inconsistent with global bottom-up inventories of CO emissions, which show a small increase or stable emissions in this region. We try to reconcile the observed CO trend with emission inventories using an atmospheric inversion of the MOPITT CO data that estimates emissions from primary sources, secondary production, and chemical sinks of CO. The atmospheric inversion indicates a $\sim -2\% \text{ yr}^{-1}$ decrease in emissions from primary sources in East Asia from 2005–2016. The decreasing emissions are mainly caused by source reductions in China. The regional MEIC inventory for China is the only bottom up estimate consistent with the inversion-diagnosed decrease of CO emissions. According to the MEIC data, decreasing CO emissions from four main sectors (iron and steel industries, residential sources, gasoline-powered vehicles, and construction materials industries) in China explain 76% of the inversion-based trend of East Asian CO emissions. This result suggests that global inventories underestimate the recent decrease of CO emission factors in China which occurred despite increasing consumption of carbon-based fuels, and is driven by rapid technological changes with improved combustion efficiency and emission control measures.

1. Introduction

Carbon monoxide (CO) is produced by the incomplete combustion of carbon-based fuels and atmospheric oxidation of hydrocarbons. It is the dominant sink of the hydroxyl radical (OH), which controls the oxidizing power of the troposphere, and hence influences the lifetime of most atmospheric pollutants and reactive greenhouse gases. Each reaction of CO with OH radicals has a theoretical maximum yield of one ozone (O_3) and one carbon dioxide (CO_2) molecule, which results in an indirect positive radiative forcing around 0.2 W m^{-2} (Myhre *et al* 2013).

The Measurements of Pollution in the Troposphere (MOPITT) space-borne instrument has been measuring tropospheric CO since 2000, and shows a decreasing global trend $\sim -1\% \text{ yr}^{-1}$ in CO total column, with stronger trends (-1.42 to $-1.60\% \text{ yr}^{-1}$) identified over Europe, the United States, and East Asia for 2000–2010 (Worden *et al* 2013). Measurements of surface concentrations confirm similar declining trends (Yoon and Pozzer 2014). The reduction of CO pollution is consistent with bottom-up emission inventories indicating reduced emissions in Europe and the United States, and chemical transport model simulations driven by those inventories reproduce the

negative trends of observed CO concentrations (Yoon and Pozzer 2014, Strode *et al* 2016). This suggests that emission inventories over these two regions successfully track the progress of emission reduction due to pollution control.

However, global inventory data show increasing emissions over East Asia, which cannot be reconciled with observed CO concentrations (Strode *et al* 2016). A downward trend $\sim -1.6\% \text{ yr}^{-1}$ in CO concentrations is observed by MOPITT (Worden *et al* 2013) over East Asia for years 2000–2010, but model simulations using the time-dependent MACCity inventory (Granier *et al* 2011) show upward trends. Strode *et al* (2016) analyzed possible model biases that contributed to the model-observation discrepancy, and found that an overestimate of O_3 associated with less O_3 break-up forming less OH played an important role. While chemistry model biases examined by Strode *et al* (2016) cannot explain all aspects of the inconsistencies, their study also questioned the increasing CO emission trends of the MACCity inventory. Yet other global bottom-up inventories also show increasing CO emissions ($0.9\%–2.9\% \text{ yr}^{-1}$) from East Asia over the past decade (Granier *et al* 2011, Crippa *et al* 2016, Zhong *et al* 2017). Anthropogenic sources are the major sources of CO in East Asia. The increasing emissions from East Asia reported by global bottom-up inventories are mainly caused by growing anthropogenic sources, which include industrial boilers, residential stoves, iron and steel production, and motor vehicles. Conversely, top-down atmospheric inversion-based emission estimates assimilating CO observations (Tohjima *et al* 2014, Yumimoto *et al* 2014, Yin *et al* 2015, Jiang *et al* 2017) all yield negative CO emission trends (-2.0 to $-3.2\% \text{ yr}^{-1}$, for 2005–2015), though the inversion approach cannot attribute the driving forces.

In the present study, we reevaluate the 2005–2016 trends of CO concentrations over East Asia, and analyze the underlying drivers of CO changes. Here East Asia refers to the geographical area covering Mainland China, Hong Kong, Macau, Taiwan, Japan, Mongolia, North Korea, and South Korea. Our goal is to understand the inconsistencies between observed and modeled CO trends over East Asia reported by previous studies. We first investigate the tropospheric CO column from the MOPITT version 7 product (Deeter *et al* 2017). Then, we use a Bayesian inversion technique (Chevallier *et al* 2005) jointly assimilating observations of the main species involved in the oxidation chain of hydrocarbons to estimate the sources and sinks of atmospheric CO. Both the diversity of assimilated data and the treatment of the uncertainty in CO chemical production and sinks are important features of our method that permit the reliable estimation of the trends in CO emissions over East Asia.

Below, we first describe the tools used to analyze the sources and sinks of atmospheric CO: the MOPITT satellite measurements, the atmospheric

inversion model that assimilates these data to provide optimized surface CO emissions, secondary CO production, and CO destruction by OH in the atmosphere, and the inventories used for comparison with inversion emissions. Then the observed trends in MOPITT data are analyzed, inversion CO emissions trends are discussed and compared with inventories.

We show that the declining trend in CO concentrations over East Asia of $-0.41 \pm 0.09\% \text{ yr}^{-1}$ for 2005–2016 ($P < 0.001$, 95% confidence interval, two-tailed) can be explained by a $-2.51 \pm 0.94\% \text{ yr}^{-1}$ ($P < 0.001$) decrease in CO emissions from primary sources in this region, which outweighs increasing secondary CO production ($1.56 \pm 0.56\% \text{ yr}^{-1}$, $P < 0.001$) due to the rising CH_4 concentrations and NMVOC emissions. Global bottom-up emission inventories fail to reproduce the negative trend of CO emissions probably because they underestimate the strength of emissions control in China, whereas the detailed inventory of Multi-resolution Emission Inventory for China (MEIC, www.meicmodel.org/) matches the top-down inversions well. The MEIC data is further analyzed to investigate sectors and emission factors that drive the decreasing CO emissions in China, which accounts for 84% of the CO emissions decrease during 2005–2016 in East Asia.

2. Methods and Data

2.1. MOPITT Version 7 CO

The MOPITT instrument was launched aboard the EOS-Terra satellite platform in December 1999 and began reporting data in March 2000 (Deeter *et al* 2003). It measures CO column on the global scale, which means the number of CO molecules between the MOPITT instrument and the Earth's surface per area of the surface (i.e. molecules cm^{-2}). The MOPITT retrieval products have been improved continuously, as confirmed by independent validation data, since 2000. Retrieval product improvements are the result of radiative transfer model enhancements, updated *a priori* information, and bias corrections (Deeter *et al* 2010, 2013, 2014).

In this study, we use MOPITT Version 7 (V7) level 2 total column retrievals from the multispectral TIR-NIR product. First V7 products were released in August 2016. As demonstrated by comparisons with CO in-situ vertical profiles measured from aircraft over North America, MOPITT V7 products exhibit much improved error characteristics (Deeter *et al* 2017). In contrast with the previous V6 product, for example, the overall biases for V7 are a few percent or less at all levels for the TIR-only, NIR-only and TIR-NIR products. For the period from 2000–2015, analysis of the long-term bias trends (i.e. bias drift) for the V7 TIR-NIR product indicates a negative bias drift for the lower-troposphere (e.g. $-1.04\% \text{ yr}^{-1}$ at 800 hPa) and an opposing positive bias drift for upper-tropospheric

retrieval levels (e.g. $1.15\% \text{ yr}^{-1}$ at 400 hPa). However, due to the opposite effects of bias drift in the lower and upper troposphere, the reported bias drift for the TIR-NIR total column product is nearly negligible (e.g. a relative bias drift of less than $0.1\% \text{ yr}^{-1}$, thus much smaller than the CO trend in East Asia). To exclude retrievals with low information content, we use only satellite retrievals with solar zenith angle less than 70° , surface pressure greater than 900 hPa, and latitude within 65°S – 65°N (Fortems-Cheiney *et al* 2011, Yin *et al* 2015).

2.2. Atmospheric inversion

We use a variational Bayesian approach (Chevallier *et al* 2005) to estimate CO emissions from primary sources, secondary production, and chemical sinks of CO for 2005–2016 (detailed in text S2–7). Technically, the Bayesian inference can be solved as a variational optimization problem by minimizing the following cost function:

$$J(\mathbf{x}) = (\mathbf{x} - \mathbf{x}^b)^T \mathbf{B}^{-1}(\mathbf{x} - \mathbf{x}^b) + (\mathbf{H}(\mathbf{x}) - \mathbf{y})^T \mathbf{R}^{-1}(\mathbf{H}(\mathbf{x}) - \mathbf{y})$$

The variables that we seek to estimate are assembled into the state vector \mathbf{x} . Through optimization we obtain the optimal \mathbf{x} given *a priori* guess \mathbf{x}^b and observation vector \mathbf{y} , for which the error statistics are represented by covariance matrices \mathbf{B} and \mathbf{R} , respectively. \mathbf{x} and \mathbf{y} relate to each other through the forward model H that can be simply understood as an operator calculating \mathbf{y} as a function of \mathbf{x} . In this study, the Bayesian framework is used in the hydrocarbon oxidation chain, consisting of methane (CH_4), formaldehyde (HCHO), CO, CO_2 as well as intermediate species, with chain reactions driven by OH among all chemical species (Pison *et al* 2009). Methyl chloroform (MCF) is also included to constrain OH concentrations.

The state vector \mathbf{x} contains OH concentrations, emission fluxes of CH_4 , HCHO , CO and MCF, and the initial concentration fields of these four species. The *a priori* CO sources include MACCity anthropogenic emissions (Granier *et al* 2011, downloaded from <http://eccad.sedoo.fr>), GFED 4s biomass burning emissions (van der Werf *et al* 2017, downloaded from www.globalfiredata.org/index.html), MEGAN biogenic emissions (Sindelarova *et al* 2014, downloaded from <http://eccad.sedoo.fr>), and POET oceanic emissions (Olivier *et al* 2003, Granier *et al* 2005, downloaded from <http://eccad.sedoo.fr>). These datasets are selected as *a priori* emissions input because they represent the most up-to-date period of CO emission fluxes freely available on global scales. The MACCity dataset is the only available dataset that provides monthly anthropogenic emissions that cover 2005–2016 (data after 2010 are emission projections). The GFED4s dataset is the latest data on biomass burning emissions that achieve high accuracy (van der Werf *et al* 2017), and the MEGAN and POET dataset are

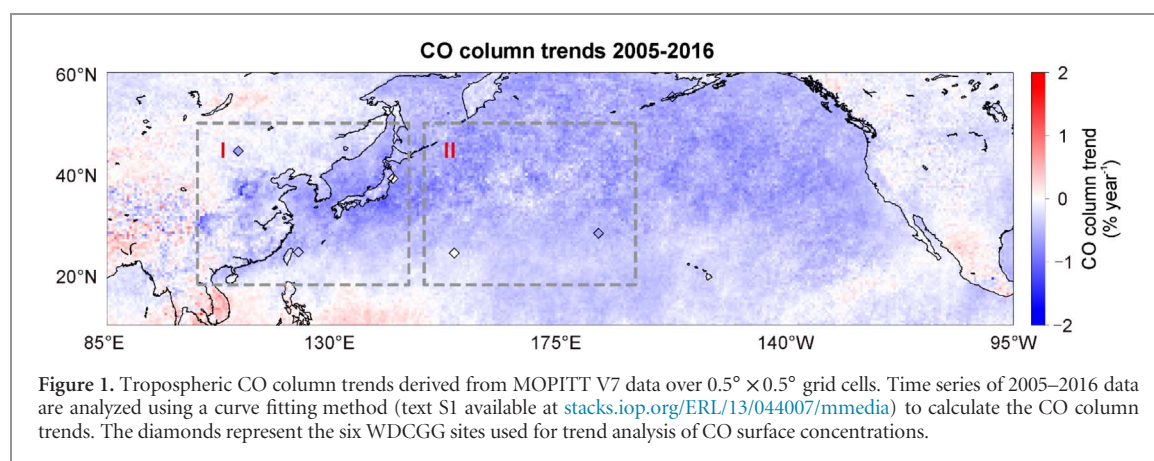
both the latest emissions data. The *a priori* information for the other variables and covariance matrix \mathbf{B} follow the configurations of Yin *et al* (2015, 2016).

The observation vector \mathbf{y} consists of satellite CO and HCHO tropospheric columns, and surface concentrations of CH_4 and MCF from in-situ networks. We use CO column retrievals from the MOPITT V7 product (Deeter *et al* 2017), HCHO column retrievals from the Ozone Monitoring Instrument V003 product (González *et al* 2015), CH_4 and MCF surface air-sample measurements from the World Data Centre for Greenhouse Gases dataset (WDCGG, <http://ds.data.jma.go.jp/gmd/wdogg/>). The forward model H is LMDz-SACS ($1.9^\circ \text{ lat} \times 3.75^\circ \text{ lon} \times 39$ vertical layers), a 3D transport model with a simplified chemistry scheme. LMDz is a general circulation model (<http://lmdz.lmd.jussieu.fr/>) which is nudged towards the European Centre for Medium-Range Weather Forecasts analyses for horizontal winds. The LMDz model is coupled with SACS module, a simplified chemistry module for the oxidation chain of hydrocarbons, which is developed by Pison *et al* (2009) on the basis of the Interaction with Chemistry and Aerosols (INCA) full chemistry model (Hauglustaine *et al* 2004). The LMDz-SACS model is described in Text S2. Details of \mathbf{y} , \mathbf{H} and covariance matrix \mathbf{R} refer to Yin *et al* (2015, 2016).

The inversion solves for emission fluxes of CH_4 , CO and MCF in each surface grid cell ($1.9^\circ \text{ lat} \times 3.75^\circ \text{ lon}$) of the transport model over eight day periods (detailed in Text S3). This inversion system has been much used and evaluated in the optimization for sources of CH_4 , CO and HCHO at both global and regional scales (Chevallier *et al* 2009, Pison *et al* 2009, Fortems-Cheiney *et al* 2009, 2011, 2012, Yin *et al* 2015, 2016). We also collected emission estimates from two regional inversion systems including China for comparison. They are from Tohjima *et al* (2014) (assimilating surface observations for 1999–2010) and Yumimoto *et al* (2014) (assimilating MOPITT Version 5 data for 2005–2010).

2.3. Bottom-up inventories

We use a regional emission inventory, the MEIC version 1.2 data (www.meicmodel.org/), to analyze the drivers of long-run CO emissions in China, which represents 90% of East Asian CO emissions. The MEIC model is a technology-based emission inventory framework developed by Tsinghua University (Zheng *et al* 2014, Liu *et al* 2015). The main emission sources are identified and quantified through the product of activity data and time-dependent emission factors, which are estimated by technology turnover models that track the penetration of different combustion technology used in emission source sectors. As emissions rates depend on combustion technology, this method can calculate dynamic emission factors that reflect technological changes over time (Zhang *et al* 2009, Lei *et al* 2011, Liu *et al* 2015). The MEIC database



provides time series of emission estimates for China spanning from 1990–2015. We use MEIC sectoral changes of CO emissions to investigate the drivers behind the variations of inversion-based emissions. To compare with the MEIC data and our inversion results, we also collected data from six bottom-up inventories including PKU (2005–2014) (Zhong *et al* 2017), REAS v2.1 (2005–2008) (Kurokawa *et al* 2013), EDGAR v4.3 (2005–2010) (Crippa *et al* 2016), MIX (2006, 2008 and 2010) (Li *et al* 2017), and the data from Xia *et al* (2016) (2005–2014) and Zhao *et al* (2012) (2005–2009).

3. Observed CO trends over East Asia

MOPITT observes a substantial decrease in tropospheric CO column over East Asia from 2005–2016 (figure 1). Geographically, the pattern of decreasing linear trends fitted to the data is not evenly distributed. Western China that is generally upwind of the heavily industrialized areas of China to the east, presents relatively weak CO trends. Larger decreases generally occur in industrialized areas with higher CO levels, such as in eastern China (-1 to $-2\% \text{ yr}^{-1}$). MOPITT also observes a strong decrease in CO ($\sim -1\% \text{ yr}^{-1}$) off the coast of East Asia (figure 1, box II), and similar decreases over the eastern Pacific, suggestive of reduced export of CO from East Asia via the prevailing westerlies.

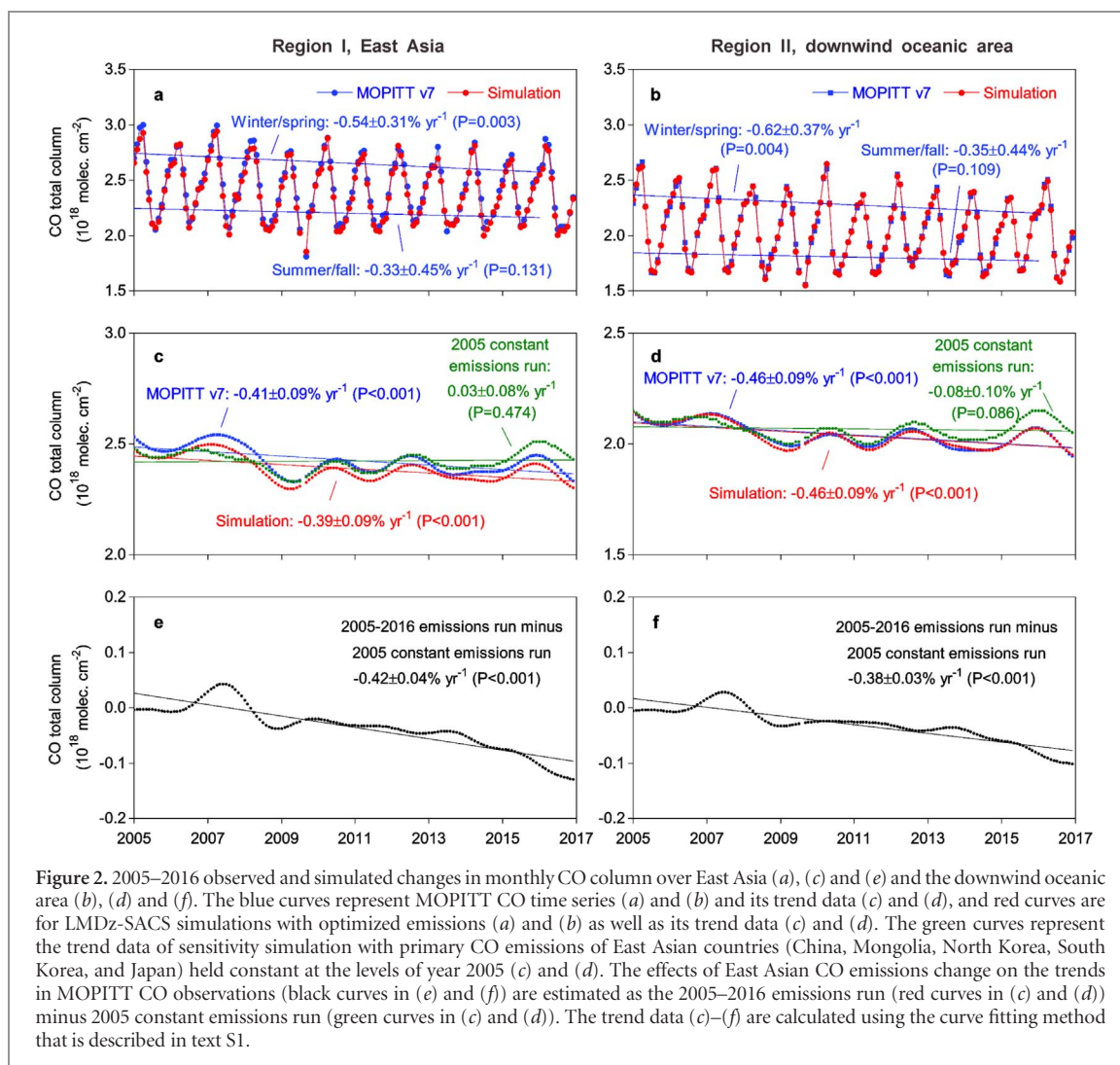
Monthly average observations reveal the detailed temporal evolution of CO column (blue curves in figures 2(a) and (b)). The peak column CO in winter/spring shows a large, significant decrease ($P=0.003$) of $-0.54 \pm 0.31\% \text{ yr}^{-1}$ over East Asia (figure 1, box I), while that in summer/fall exhibits a small, insignificant decrease ($P=0.131$) of $-0.33 \pm 0.45\% \text{ yr}^{-1}$. The de-seasonalized monthly CO column shows a medium, significant decrease ($P < 0.001$) of $-0.41 \pm 0.09\% \text{ yr}^{-1}$ (blue curves in figure 2(c)). The downwind oceanic region of East Asian continent (figure 1, box II) exhibits similar decreasing linear trends (blue curves in figures 2(b) and (d)). We also investigate surface concentrations of CO from the WDCGG dataset,

which includes six remote observing stations (the diamonds in figure 1) within East Asia and its downwind areas. De-seasonalized monthly surface observations show a significant negative trend ($P < 0.001$) of $-0.46 \pm 0.14\% \text{ yr}^{-1}$ for 2005–2016, driven by a significant decrease ($P=0.008$) of $-1.00 \pm 0.67\% \text{ yr}^{-1}$ in winter/spring. The observed decrease of mean annual surface concentrations ($-0.46 \pm 0.14\% \text{ yr}^{-1}$, $P < 0.001$) is similar to the linear trend of CO column ($-0.41 \pm 0.09\% \text{ yr}^{-1}$, $P < 0.001$) observed by MOPITT. We note that the time period selected for the trend calculation is also important due to significant variability in global CO from large biomass burning episodes such as the boreal fires in 2002, 2003 (Yurganov *et al* 2005) and the El Nino driven fires in Indonesia in September–October 2015 (Field *et al* 2016, Yin *et al* 2016).

4. Inverse analysis of CO sources trends

The results from the LMDz-SACS inversion assimilating MOPITT CO and other related tracer measurement show a linear reduction of -2000 to $-4000 \text{ kg km}^{-2} \text{ yr}^{-1}$ in high emission areas such as East China, South Korea and Japan for 2005–2016 (figures 3(a) and (b)). The anthropogenic sources drive the downward trend because biomass burning contributes less (figure 3(c)). We also see a linear growth of $< 2000 \text{ kg km}^{-2} \text{ yr}^{-1}$ in CO emissions over the areas adjacent to South Central Siberia due to increased fire activities (figures 3(b) and (c)). However, compared with the widespread decline in emissions over high emission areas, the slight increase of wildfires has little effect on net emissions due to their small size.

Inversion-based estimates show a decreasing trend of $-2.51 \pm 0.94\% \text{ yr}^{-1}$ for CO emissions in East Asia ($P < 0.001$) with respect to 2005 ($-5.07 \text{ Tg year}^{-1}$) spanning from 2005–2016 (the red solid curve in figure 3(d)). This negative trend is not present in the *a priori* emission fluxes used in the LMDz-SACS inversion (the black solid curve in figure 3(d)). For China specifically, the inversion infers a decrease of $-2.16 \pm 3.40\% \text{ yr}^{-1}$ ($P=0.152$) for 2005–2010, which

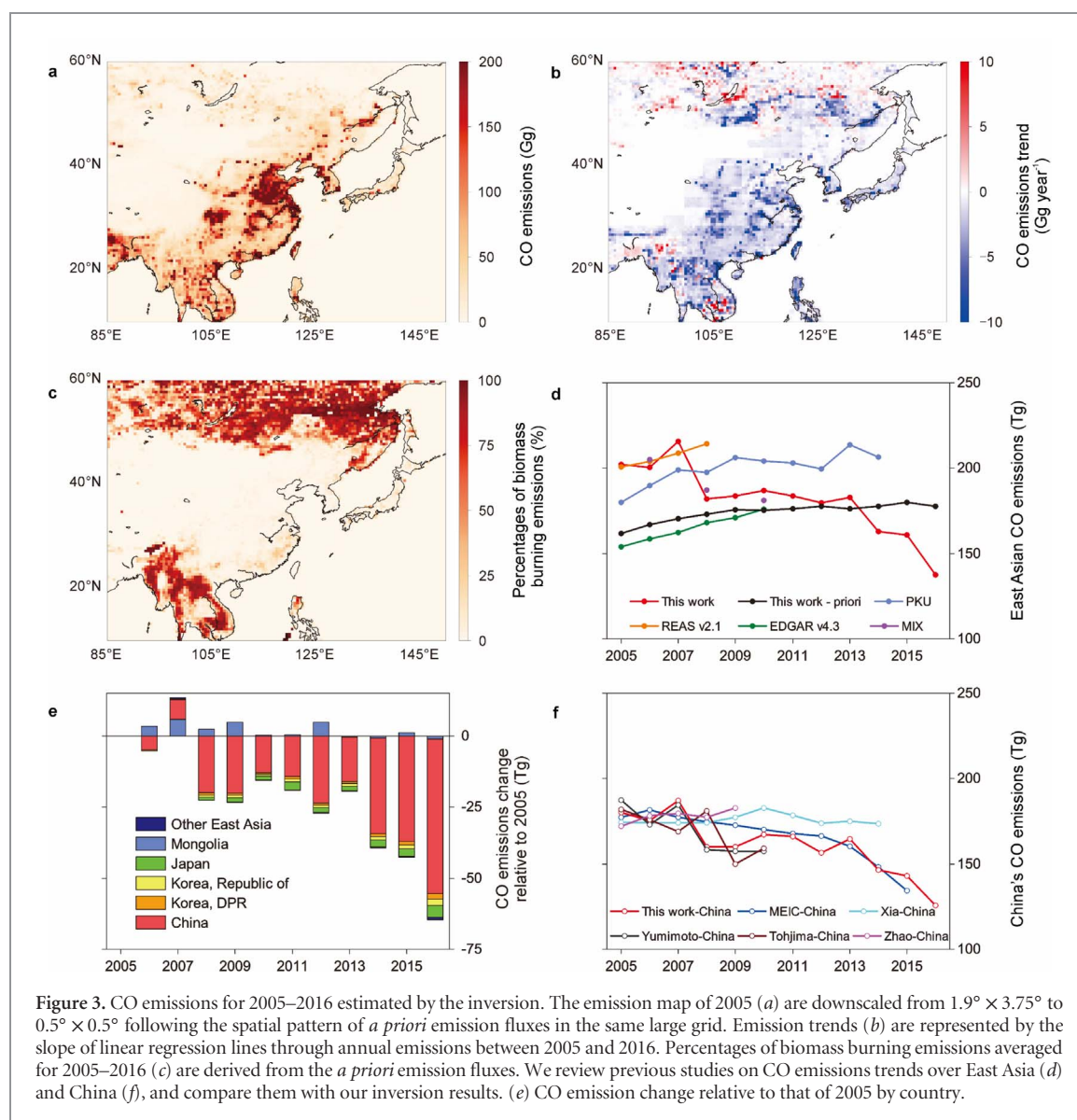


is close to the regional inversions of Tohjima *et al* (2014) and Yumimoto *et al* (2014) that show a linear emission decrease of $\sim -2\% \text{ yr}^{-1}$ in China for the same period. Overall, our inversion results for East Asia change in three successive phases: Phase I (2005–2007, a slight increase), Phase II (2008–2013, a drop by 15.55% in 2008 and then a slight decrease of $-0.18 \pm 0.91\% \text{ yr}^{-1}$, $P=0.621$), and Phase III (2014–2016, an accelerated decline). The abrupt drop of CO emissions in 2008 is coincident with a sharp decrease in MOPITT CO column in the second half of this year (figure 2(c)) as reported by Witte *et al* (2009), Yurganov *et al* (2010), Worden *et al* (2012) and Strode *et al* (2016). Witte *et al* (2009) showed CO reductions of 12% at 700 hPa using the MOPITT data that was attributed to rapid and strict controls on pollutant emissions in East China for the Beijing Olympics in August–September 2008. Yurganov *et al* (2010) estimated that the upper limit of monthly drop of CO column could reach 30% at the end of 2008 due to reduced activities during the economic recession. Therefore, one may think that the abrupt drop of 2008 CO emissions over East Asia was probably caused by stringent pollution control for the Beijing

Olympics followed by the subsequent sharp slowdown in the GDP growth and stalled industrial production.

In Phase I, the inversion-based emissions agree well with REAS v2.1 and MIX bottom-up inventories, and exhibit similar growth to other bottom-up inventories. In Phase II, our inversion emissions are consistent with the MIX data for the years of 2008 and 2010, but are distinct from the PKU data, MACCity data and EDGARv4.3. These three global inventories fail to reproduce both the large and abrupt fall of emissions in 2008 and the slight decrease after that. In Phase III, the inversion estimate of this work shows an accelerated decreasing trend in emissions, inconsistent with the rising emissions of MACCity. The global bottom-up inventories mentioned above fail to reflect the evolution of emissions over East Asia during Phases II and III, thus they cannot capture the declining CO trend for 2005–2016. Over all three phases, China is responsible for 84% of the decrease (the red bar in figure 3(e)), suggestive of the dominant role that China plays in the trend and variability of East Asian CO emissions.

We further evaluate the inversion emissions over China (the red curve in figure 3(f)) against the MEIC inventory (the blue curve in figure 3(f)).



The MEIC data shows a decreasing linear trend of $-2.16 \pm 0.79\% \text{ yr}^{-1}$ ($P < 0.001$) which agrees well with our inversion estimates ($-1.86 \pm 0.92\% \text{ yr}^{-1}$, $P = 0.001$) for 2005–2015, while the bottom-up emission inventories developed by Xia *et al* (2016) and Zhao *et al* (2012) both show flattening emissions. Still, none of these regional bottom-up inventories can capture the abrupt drop of CO emissions in 2008.

5. Drivers of the decline in CO concentrations and emissions

We use the LMDz-SACS forward model to check the contributions of primary CO emissions trends ($-2.51 \pm 0.94\% \text{ yr}^{-1}$, $P < 0.001$) to the observed 2005–2016 trend of CO concentrations ($-0.41 \pm 0.09\% \text{ yr}^{-1}$, $P < 0.001$). Using the optimized emissions, optimized OH concentrations, and initial concentration fields inferred from the inversion, the forward model reproduces the absolute magnitude and temporal

evolution of MOPITT observations well (red curves in figures 2(a)–(d)), indicating that the inversion fits the CO concentrations well.

To quantify the impacts of East Asian CO source change on the trends in MOPITT CO observations, we conduct a sensitivity simulation with primary CO emissions held constant at the levels of year 2005 in East Asian countries (green curves in figures 2(c) and (d)) and other factors—emissions in other regions, OH concentrations, and initial concentration fields—being variable and taken from the inversion. The modeling results for this sensitivity simulation show a positive, insignificant trend of CO column ($0.03 \pm 0.08\% \text{ yr}^{-1}$, $P = 0.474$) over East Asia (figure 2(c)), indicating that the other factors in the model do not explain the decrease of CO concentrations, and thus that only a reduction of primary CO emissions in East Asia can match the MOPITT trends ($-0.41 \pm 0.09\% \text{ yr}^{-1}$, $P < 0.001$). These results are consistent with the findings of Strode *et al* (2016), whose simulation with constant CO emissions showed a positive trend in

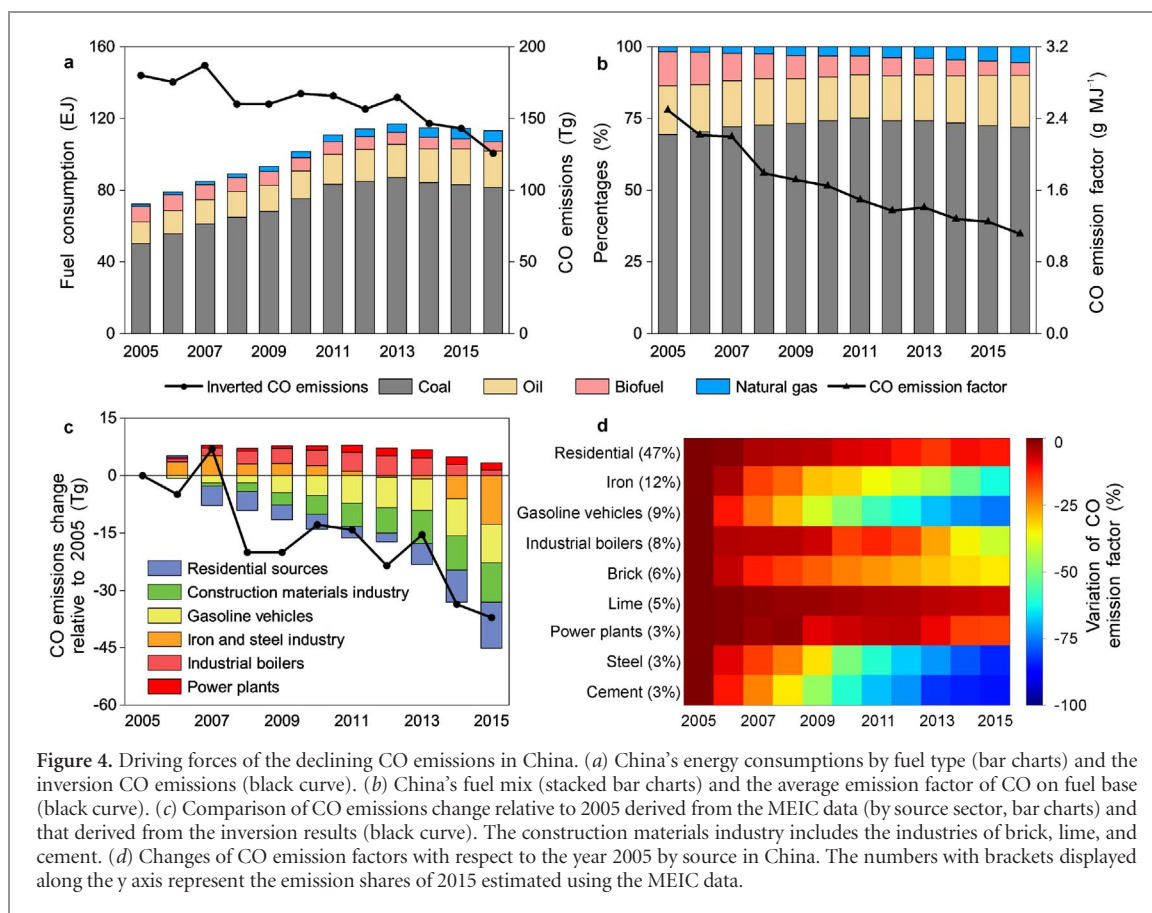


Figure 4. Driving forces of the declining CO emissions in China. (a) China's energy consumptions by fuel type (bar charts) and the inversion CO emissions (black curve). (b) China's fuel mix (stacked bar charts) and the average emission factor of CO on fuel base (black curve). (c) Comparison of CO emissions change relative to 2005 derived from the MEIC data (by source sector, bar charts) and that derived from the inversion results (black curve). The construction materials industry includes the industries of brick, lime, and cement. (d) Changes of CO emission factors with respect to the year 2005 by source in China. The numbers with brackets displayed along the y axis represent the emission shares of 2015 estimated using the MEIC data.

CO concentrations for 2000–2010 over East China. Worden *et al* (2013) also suggests a decrease of primary CO emissions over East China to match the declining CO concentrations in this region for 2000–2011.

We subtract the model run driven by constant 2005 emissions from the simulation with variable 2005–2016 emissions (black curves in figures 2(e) and (f)). The 2005–2016 emission update results in a decrease of $-0.42 \pm 0.04\% \text{ yr}^{-1}$ ($P < 0.001$) in CO column (figure 2(e) that matches the MOPITT observations well ($-0.41 \pm 0.09\% \text{ yr}^{-1}$, $P < 0.001$, figure 2(c)). The same holds true in areas downwind (figures 2(d) and (f)). These results show that primary emissions change over East Asian countries can account for the entire declining trend of CO concentrations observed in this region.

As China dominates the East Asian emission budget, and because of the good match between inversion-based emission estimates and the independent bottom-up MEIC inventory, we use the MEIC data to understand the drivers behind the declining trends of Chinese emissions (figure 4). In MEIC, China's CO emissions keep falling because decreasing emission factors totally offset the increasing use of carbon fuels (figures 4(a) and (b)). Generally, all important source sectors (e.g. residential, iron, gasoline-powered vehicles, and industrial boilers) are improving combustion efficiency and strengthening air pollution control in the last decade, which ultimately leads to the steady decline in emission factors

spanning from -7% to -86% across sources for 2005–2015 (figures 4(c) and (d)).

Four source sectors dominate the downward trends of China's emissions, including iron and steel industries, residential sources, gasoline-powered vehicles, and construction materials industries (figure 4(c)). These four sectors are responsible for 92% of China's emissions cut, and can explain 76% of the emissions decrease for East Asia. In iron and steel industries, CO is an unavoidable byproduct released from blast furnace and basic oxygen furnace. Industry operators have reduced gas leaks, so the amount of CO emitted to the atmosphere is decreasing (-62% for iron production and -84% for steel making in 2005–2015 estimated by the MEIC data). Residential sources contribute half of CO emissions in China due to the extensive use of low-efficiency fuels and stoves. China has started to promote the use of clean stoves and phase out traditional biofuels (i.e. wood and crop residual) since 1990s (World Bank 2013), which helps lower CO emission rates of the residential sector (-12% for 2005–2015). The emissions from gasoline-powered vehicles are controlled successfully with the increasingly more stringent emission standards from stage 2 (equivalent to Euro 2/II standards) to stage 5 (equivalent to Euro 5/V standards) implemented since 2005 (Wu *et al* 2017). Consequently, the fleet average CO emission factor has been reduced by 76% from 2005–2015 according to the MEIC emission factors. The construction materials industries reduce

emissions through using high efficiency kilns. For example, the cement industry replaces low-efficiency shaft kilns with a new type of rotary kiln, called the new dry process in China, in the last decade (Lei *et al* 2011). The percentage of cement produced by the new kilns increase from 44% in 2005 to 99% in 2015, which reduces the CO emission factor by 86%. These four sectors show a much larger decline of emissions since 2013, because China accelerated the air pollution control at the end of 2013 to fight against severe haze pollution (State Council of the People's Republic of China 2013).

6. Conclusions and implications

In this study, we have analyzed the main sources of inconsistency between 2005–2016 CO trends from MOPITT column observations and from global emission inventories in East Asia. The most important findings are that (1) the decreasing linear trend of $-0.41 \pm 0.09\% \text{ yr}^{-1}$ ($P < 0.001$) in CO concentrations over East Asia is due to a $-2.51 \pm 0.94\% \text{ yr}^{-1}$ ($P < 0.001$) decrease in emissions from primary sources in this region, that is a cumulative decline of -32% from 2005 to 2016 and (2) 76% of the emissions decline over East Asia can be explained by emissions control of four source sectors in China, i.e. iron and steel industries, residential sources, gasoline-powered vehicles, and construction materials industries. This emission decrease is enough to counterbalance the effect of rising concentrations of CH_4 ($0.38 \pm 0.01\% \text{ yr}^{-1}$, derived from WDCGG observations) and increasing emissions of NMVOC ($4.59 \pm 0.44\% \text{ yr}^{-1}$, estimated by MEIC data) in East Asia, that increase the secondary CO formation at a rate of $1.56 \pm 0.56\% \text{ yr}^{-1}$ ($P < 0.001$) according to our multispecies inversion (Text S7). Global bottom-up emission inventories were less successful in capturing the negative emission trends than the MEIC inventory, probably because they underestimate the strength of emissions control in East Asia, especially in China. The MACCity inventory since 2010 are emission projection data so they cannot reflect the recent trends in China CO emissions, and changes in emission factors. As a fast-growing economy with rapid technology changes, emission factors vary so fast that capturing the CO emissions variation remains a big challenge. Moreover, multi-instrument space-borne observations also verified recent (2005–2015) reductions in air pollution loadings of sulfur dioxide, nitrogen dioxide and aerosols over East Asia (Krotkov *et al* 2016, Liu *et al* 2016, Zhang *et al* 2017, Zhao *et al* 2017). It is difficult to predict these changes using models and conservative estimates. Our research method incorporating observations, inverse modeling and technology based bottom-up inventory provides an opportunity to better understand what happened recently as well as the underlying drivers. Though all data

and methods are subject to their own uncertainties, the final results are considered robust given the consistency between observations from MOPITT satellite and ground-based CO measurements (section 3), between our inversion analysis and other inversion results (section 4), and between our inversion results and the latest bottom-up emissions data (section 5).

Acknowledgments

Data to support this research are available upon request to the corresponding author Bo Zheng (bo.zheng@lsce.ipsl.fr). We acknowledge the data providers of NCAR MOPITT for satellite CO retrievals, SAO OMI for CH_2O retrievals, and WDCGG for CH_4 , CO, and MCF surface air-sample measurements. We also thank F Marabelle for computer support at LSCE.

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References

- Chevallier F, Fisher M, Peylin P, Serrar S, Bousquet P, Bréon F M, Chédin A and Ciais P 2005 Inferring CO_2 sources and sinks from satellite observations: method and application to TOVS data *J. Geophys. Res. Atmos.* **110** D24309
- Chevallier F, Fortems A, Bousquet P, Pison I, Szopa S, Devaux M and Hauglustaine D A 2009 African CO emissions between years 2000 and 2006 as estimated from MOPITT observations *Biogeosciences* **6** 103–11
- Crippa M, Janssens-Maenhout G, Dentener F, Guizzardi D, Sindelarova K, Muntean M, Van Dingenen R and Granier C 2016 Forty years of improvements in European air quality: regional policy-industry interactions with global impacts *Atmos. Chem. Phys.* **16** 3825–41
- Deeter M N *et al* 2003 Operational carbon monoxide retrieval algorithm and selected results for the MOPITT instrument *J. Geophys. Res. Atmos.* **108** 4399
- Deeter M N *et al* 2010 The MOPITT version 4 CO product: algorithm enhancements, validation, and long-term stability *J. Geophys. Res. Atmos.* **115** D07306
- Deeter M N, Martínez-Alonso S, Edwards D P, Emmons L K, Gille J C, Worden H M, Pittman J V, Daube B C and Wofsy S C 2013 Validation of MOPITT Version 5 thermal-infrared, near-infrared, and multispectral carbon monoxide profile retrievals for 2000–2011 *J. Geophys. Res. Atmos.* **118** 6710–25
- Deeter M N, Martínez-Alonso S, Edwards D P, Emmons L K, Gille J C, Worden H M, Sweeney C, Pittman J V, Daube B C and Wofsy S C 2014 The MOPITT Version 6 product: algorithm enhancements and validation *Atmos. Meas. Tech.* **7** 3623–32
- Deeter M N, Edwards D P, Francis G L, Gille J C, Martínez-Alonso S, Worden H M and Sweeney C 2017 A climate-scale satellite record for carbon monoxide: the MOPITT Version 7 product *Atmos. Meas. Tech.* **10** 2533–55
- Field R D *et al* 2016 Indonesian fire activity and smoke pollution in 2015 show persistent nonlinear sensitivity to El Niño-induced drought *Proc. Natl Acad. Sci.* **113** 9204–9
- Fortems-Cheiney A, Chevallier F, Pison I, Bousquet P, Carouge C, Clerbaux C, Coheur P F, George M, Hurtmans D and Szopa S 2009 On the capability of IASI measurements to inform about CO surface emissions *Atmos. Chem. Phys.* **9** 8735–43

- Fortems-Cheiney A, Chevallier F, Pison I, Bousquet P, Szopa S, Deeter M N and Clerbaux C 2011 Ten years of CO emissions as seen from measurements of pollution in the Troposphere (MOPIIT) *J. Geophys. Res. Atmos.* **116** D05304
- Fortems-Cheiney A, Chevallier F, Pison I, Bousquet P, Saunio M, Szopa S, Cressot C, Kurosu T P, Chance K and Fried A 2012 The formaldehyde budget as seen by a global-scale multi-constraint and multi-species inversion system *Atmos. Chem. Phys.* **12** 6699–721
- González Abad G, Liu X, Chance K, Wang H, Kurosu T P and Suleiman R 2015 Updated Smithsonian Astrophysical Observatory Ozone Monitoring Instrument (SAO OMI) formaldehyde retrieval *Atmos. Meas. Tech.* **8** 19–32
- Granier C, Lamarque J F, Mieville A, Muller J F, Olivier J, Orlando J, Peters J, Petron G, Tyndall G and Wallens S 2005 POET a database of surface emissions of ozone precursors
- Granier C *et al* 2011 Evolution of anthropogenic and biomass burning emissions of air pollutants at global and regional scales during the 1980–2010 period *Clim. Change* **109** 163–90
- Hauglustaine D A, Hourdin F, Jourdain L, Filiberti M A, Walters S, Lamarque J F and Holland E A 2004 Interactive chemistry in the Laboratoire de Météorologie dynamique general circulation model: description and background tropospheric chemistry evaluation *J. Geophys. Res. Atmos.* **109** D04314
- Jiang Z, Worden J R, Worden H, Deeter M, Jones D B A, Arellano A F and Henze D K 2017 A 15 year record of CO emissions constrained by MOPITT CO observations *Atmos. Chem. Phys.* **17** 4565–83
- Krotkov N A *et al* 2016 Aura OMI observations of regional SO₂ and NO₂ pollution changes from 2005 to 2015 *Atmos. Chem. Phys.* **16** 4605–29
- Kurokawa J, Ohara T, Morikawa T, Hanayama S, Janssens-Maenhout G, Fukui T, Kawashima K and Akimoto H 2013 Emissions of air pollutants and greenhouse gases over Asian regions during 2000–2008: regional emission inventory in Asia (REAS) version 2 *Atmos. Chem. Phys.* **13** 11019–58
- Lei Y, Zhang Q, Nielsen C and He K 2011 An inventory of primary air pollutants and CO₂ emissions from cement production in China, 1990–2020 *Atmos. Environ.* **45** 147–54
- Lei Y, Zhang Q, He K B and Streets D G 2011 Primary anthropogenic aerosol emission trends for China, 1990–2005 *Atmos. Chem. Phys.* **11** 931–54
- Li M *et al* 2017 MIX: a mosaic Asian anthropogenic emission inventory under the international collaboration framework of the MICS-Asia and HTAP *Atmos. Chem. Phys.* **17** 935–63
- Liu F, Zhang Q, Tong D, Zheng B, Li M, Huo H and He K B 2015 High-resolution inventory of technologies, activities, and emissions of coal-fired power plants in China from 1990 to 2010 *Atmos. Chem. Phys.* **15** 13299–317
- Liu F *et al* 2016 Recent reduction in NO_x emissions over China: synthesis of satellite observations and emission inventories *Environ. Res. Lett.* **11** 114002
- Myhre G *et al* 2013 Anthropogenic and natural radiative forcing, in: climate change 2013: the physical science basis *Contribution of Working Group I to the Fifth Assessment Report of the Intergovernmental Panel on Climate Change* (Cambridge: Cambridge University Press)
- Olivier J, Peters J, Granier C, Petron G, Muller J F and Wallens S 2003 Present and future surface emissions of atmospheric compounds *POET Report #2* EU project EVK2–1999–00011
- Pison I, Bousquet P, Chevallier F, Szopa S and Hauglustaine D 2009 Multi-species inversion of CH₄, CO and H₂ emissions from surface measurements *Atmos. Chem. Phys.* **9** 5281–97
- Sindelarova K, Granier C, Bouarar I, Guenther A, Tilmes S, Stavrakou T, Müller J F, Kuhn U, Stefani P and Knorr W 2014 Global data set of biogenic VOC emissions calculated by the MEGAN model over the last 30 years *Atmos. Chem. Phys.* **14** 9317–41
- State Council of the People's Republic of China 2013 Air Pollution Prevention and Control Action Plan (www.gov.cn/jzwgk/2013-09/12/content_2486773.htm)
- Strode S A *et al* 2016 Interpreting space-based trends in carbon monoxide with multiple models *Atmos. Chem. Phys.* **16** 7285–94
- Tohijima Y, Kubo M, Minejima C, Mukai H, Tanimoto H, Ganshin A, Maksyutov S, Katsumata K, Machida T and Kita K 2014 Temporal changes in the emissions of CH₄ and CO from China estimated from CH₄/CO₂ and CO/CO₂ correlations observed at Hateruma Island *Atmos. Chem. Phys.* **14** 1663–77
- van der Werf G R *et al* 2017 Global fire emissions estimates during 1997–2016 *Earth Syst. Sci. Data* **9** 697–720
- Witte J C, Schoeberl M R, Douglass A R, Gleason J F, Krotkov N A, Gille J C, Pickering K E and Livesey N 2009 Satellite observations of changes in air quality during the 2008 Beijing Olympics and Paralympics *Geophys. Res. Lett.* **36** L17803
- Worden H M, Cheng Y, Pfister G, Carmichael G R, Zhang Q, Streets D G, Deeter M, Edwards D P, Gille J C and Worden J R 2012 Satellite-based estimates of reduced CO and CO₂ emissions due to traffic restrictions during the 2008 Beijing Olympics *Geophys. Res. Lett.* **39** L14802
- Worden H M *et al* 2013 Decadal record of satellite carbon monoxide observations *Atmos. Chem. Phys.* **13** 837–50
- World Bank 2013 China: accelerating household access to clean cooking and heating *East Asia and Pacific Clean Stove Initiative Series* (Washington, DC: World Bank)
- Wu Y, Zhang S, Hao J, Liu H, Wu X, Hu J, Walsh M P, Wallington T J, Zhang K M and Stevanovic S 2017 On-road vehicle emissions and their control in China: a review and outlook *Sci. Total Environ.* **574** 332–49
- Xia Y, Zhao Y and Nielsen C P 2016 Benefits of China's efforts in gaseous pollutant control indicated by the bottom-up emissions and satellite observations 2000–2014 *Atmos. Environ.* **136** 43–53
- Yin Y, Chevallier F, Ciais P, Broquet G, Fortems-Cheiney A, Pison I and Saunio M 2015 Decadal trends in global CO emissions as seen by MOPITT *Atmos. Chem. Phys.* **15** 13433–51
- Yin Y *et al* 2016 Variability of fire carbon emissions in equatorial Asia and its nonlinear sensitivity to El Niño *Geophys. Res. Lett.* **43** 472–9
- Yoon J and Pozzer A 2014 Model-simulated trend of surface carbon monoxide for the 2001–2010 decade *Atmos. Chem. Phys.* **14** 10465–82
- Yumimoto K, Uno I and Itahashi S 2014 Long-term inverse modeling of Chinese CO emission from satellite observations *Environ. Pollut.* **195** 308–18
- Yurganov L N *et al* 2005 Increased Northern Hemispheric carbon monoxide burden in the troposphere in 2002 and 2003 detected from the ground and from space *Atmos. Chem. Phys.* **5** 563–73
- Yurganov L, McMillan W, Grechko E and Dzhola A 2010 Analysis of global and regional CO burdens measured from space between 2000 and 2009 and validated by ground-based solar tracking spectrometers *Atmos. Chem. Phys.* **10** 3479–94
- Zhang J, Reid J S, Alfaro-Contreras R and Xian P 2017 Has China been exporting less particulate air pollution over the past decade? *Geophys. Res. Lett.* **44** 2941–8
- Zhang Q *et al* 2009 Asian emissions in 2006 for the NASA INTEX-B mission *Atmos. Chem. Phys.* **9** 5131–53
- Zhao B *et al* 2017 Decadal-scale trends in regional aerosol particle properties and their linkage to emission changes *Environ. Res. Lett.* **12** 054021
- Zhao Y, Nielsen C P, McElroy M B, Zhang L and Zhang J 2012 CO emissions in China: uncertainties and implications of improved energy efficiency and emission control *Atmos. Environ.* **49** 103–13
- Zheng B, Huo H, Zhang Q, Yao Z L, Wang X T, Yang X F, Liu H and He K B 2014 High-resolution mapping of vehicle emissions in China in 2008 *Atmos. Chem. Phys.* **14** 9787–805
- Zhong Q, Huang Y, Shen H, Chen Y, Chen H, Huang T, Zeng E Y and Tao S 2017 Global estimates of carbon monoxide emissions from 1960 to 2013 *Environ. Sci. Pollut. Res.* **24** 864–73