Response of winter fine particulate matter concentrations to emission and meteorology changes in North China

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Abstract

The winter haze is a growing problem in North China, but the causes have not been well understood. The chemistry version of the Weather Research and Forecasting model (WRF-Chem) was applied in North China to examine how the $PM_{2.5}$ concentrations change in response to changes in emissions (sulfur dioxide (SO_2), black carbon (BC), organic carbon (OC), ammonia (NH_3), and nitrogen oxides (NO_x)), as well as meteorology (temperature, relative humidity (RH), and wind speeds) changes in winter. From 1960 to 2010, the dramatic changes in emissions lead to +260% increases in sulfate, +320% increases in nitrate, +300% increases in ammonium, +160% increases in BC and 50% increases in OC. The responses of $PM_{2.5}$ to individual emission specie indicate that the simultaneous increases in SO_2, NH_3 and NO_x emissions dominated the increases in $PM_{2.5}$ concentrations. $PM_{2.5}$ shows more notable increases in response to changes in SO_2 and NH_3 as compared to increases in response to changes in NO_x emissions. In addition, OC also accounts for a large fraction in $PM_{2.5}$ changes. These results provide some implications for haze pollution control. The responses of $PM_{2.5}$ concentrations to temperature increases are dominated by changes in wind fields and mixing heights. $PM_{2.5}$ shows relatively smaller changes in response to temperature increases and RH decreases, compared to changes in response to changes in wind speed and aerosol feedbacks. From 1960 to 2010, aerosol feedbacks have been significantly enhanced, due to higher aerosol loadings. The discussions in this study indicate that dramatic changes in emissions are the main cause of increasing haze events in North China, and long-term trends in atmospheric circulations maybe another important cause since $PM_{2.5}$ is shown to be substantially affected by wind speed and aerosol feedbacks. More studies are necessary to get a better understanding of the aerosol-circulation interactions.

1 Introduction

$PM_{2.5}$ (particulate matter with diameter equal to or less than 2.5μm) is a main air pollution concern due to its adverse effects on public health (Gao et al., 2015; Pope et al., 2009). Pope et al. (2009) estimated that a decrease of 10μg $PM_{2.5}$ is related to about 0.6 year mean life expectancy increase. $PM_{2.5}$ is also associated with visibility reduction and regional climate (Cheung et al., 2005). Many cities in North China are experiencing severe haze pollution with exceedingly high $PM_{2.5}$ concentrations. In January 2010, a regional haze occurred in North China and maximum hourly $PM_{2.5}$ concentration in Tianjin was over 400μg/m^3 (Zhao et al., 2013). In January 2013, another unprecedented haze event happened, and the daily $PM_{2.5}$ concentrations in some areas of Beijing and Shijiazhuang reached over 500μg/m^3 (L. T. Wang et al., 2014), and instantaneous $PM_{2.5}$ concentration at some urban measurement sites were over 1000μg/m^3 (Zheng et al., 2015).

It is well known that particulate matter levels are strongly influenced by emissions and meteorological conditions (Steiner et al., 2006). The PM in the atmosphere can be directly emitted from sources like wildfires, combustion, wind-blown dust, and sea-salt, or formed from emitted gases through secondary aerosol formation mechanisms. Meteorology affects PM levels via changing emissions, chemical reactions, transport and deposition processes (Mu and Liao, 2014). For example, increasing wildfire emission in North America is mainly caused by warmer temperatures and precipitation changes (Dawson et al., 2014), and increased temperature leads to higher biogenic emissions, which are important precursors of secondary organic aerosols (Dawson et al., 2014; Heald et al., 2008; Jacob and Winner, 2009). Increasing temperature also increases sulfate concentration due to the temperature dependence of SO_2 oxidation and resulting higher SO_2 oxidation rates (Aw and Kleeman, 2003; Dawson et al., 2007) and semi-volatile aerosols may decrease due to evaporation under higher temperature (Sheehan and Bowman, 2001; Dawson et al., 2007; Tsigaridis and Kanakidou, 2007). Higher relative humidity (RH) favors the formation of nitrate and increasing precipitation decreases all PM species via wet scavenging (Dawson et al., 2007; Tai et al., 2010). Furthermore, increasing clouds promote in-cloud sulfate production (Tai et al., 2010) and changes in wind speed and mixing height determines the dilution of primary and secondary PM (Jimenez-Guerrero et al., 2012; Megaritis et al., 2014; Pay et al., 2012).

With rapid economic and industrial developments, emissions in China have grown during the past years. It is estimated that NO_x emissions in China increased by 70% from 1995 to 2004 (Zhang et al., 2007), Black Carbon (BC) by ~50% from 2000 to 2010 (Lu et al., 2011), Organic Carbon (OC) by ~30% from 2000 to 2010 (Lu et al., 2011), and SO_2 by ~60% from 2000 to 2006 (Lu et al., 2011). Apart from emission changes, it was observed that the winter is warming up in China, especially in the northern part (Guo et al., 2013; Hu et al., 2003; Ren et al., 2012). In addition, wind speed in North China has lowered (Shi et al., 2015; Wang et al., 2004) and RH has decreased in China (Song et al., 2012; Wang et al., 2004).

Many studies have investigated the impacts of emission changes on aerosol formation (Aksoyoglu et al., 2011; Andreani-Aksoyoglu et al., 2008; Megaritis et al., 2013; Tsimpidi et al., 2012a; Tsimpidi et al., 2012b) and the effects of climate/meteorology changes on $PM_{2.5}$ concentrations (Dawson et al., 2007; Megaritis et al., 2013; Megaritis et al., 2014; Tagaris et al., 2007; Tai et al., 2012a; Tai et al., 2012b) in Europe and in the United States. The haze pollution is growing in China, especially in North China, but the causes of the growth are not well understood. For haze pollution in China, it has been reported that aerosol feedbacks that change radiation and temperature can worsen pollution (Gao et al., 2016; Petäjä et al., 2016; Xing et al., 2015c; Zhang et al., 2015). In addition, the connections between haze and meteorological conditions have been established in many former studies (Fu et al., 2014; Jia et al., 2015; Leng et al., 2015; C. Li et al., 2015; Wang and Chen, 2016; Yang et al., 2016; X. Y. Zhang et al., 2015; Zhang et al., 2016). However, the roles of the large emission changes during the last 4 to 5 decades and the observed meteorology changes in North China are not known.

The main objective of this study is to investigate the responses of $PM_{2.5}$ and its major species to changes in emissions, including SO_2, BC, OC, NO_x and NH_3, and to temperature, RH and wind speed changes in North China region. Winter haze in North China has a large contribution from secondary inorganic aerosols and secondary inorganic aerosols are influenced by emissions, temperature and RH. The models used in previous studies of emissions and meteorology perturbations referenced above are all offline models, which are not capable of considering the feedbacks of changing meteorology on other meteorological variables, and the impacts of aerosols on meteorology. However, as pointed by Gao et al. (2016) and J. Wang et al. (2014) aerosol feedbacks should not be neglected when modeling aerosol in China. In this study, we consider aerosol feedbacks using the fully online coupled WRF-Chem model.

This paper is organized as follows. First, the WRF-Chem model, model settings and domain settings are briefly described and then in the next section, emission changes from 1960 to 2010 and accordingly $PM_{2.5}$ changes are discussed. After that, the responses of $PM_{2.5}$ to changes in each emission species are analyzed. At last, the impacts of temperature, RH and wind speed changes on $PM_{2.5}$ are analyzed and discussed.

2 Methodology

2.1 WRF-Chem model

The WRF-Chem model is the chemistry version of the Weather Research and Forecasting model, which is fully online coupled that allows gases and aerosols simulations at the same time as meteorology simulations. In this study, we used a configuration that includes direct and indirect feedbacks. The gas phase mechanism used in this study is the Carbon Bond Mechanism version Z (CBM-Z), which includes 67 species and 164 reactions (Zaveri and Peters, 1999; Zaveri et al., 2008). The gas-particle partitioning module used is the MOSAIC module, which considers all important aerosol components, such as sulfate, nitrate, ammonium, BC, and OC (Zaveri et al., 2008). Eight size bins version of MOSAIC was used and the aerosol sizes ranged from 0.039μm to 10μm. Wind-blown dust was modeled online using the AFWA scheme. Two nested domains with 81km and 27km horizontal grid resolutions from outer to innermost and 27 vertical grids were used (Figure S1 in supplementary material). Analysis nudging of meteorology variables was used for the outer domain. The model physics configurations generally follow the settings in Gao et al. (2016). Inputs into the model include meteorological boundary and initial conditions (BCs and ICs) from NCEP FNL 1 °×1 °data and chemical boundary and initial conditions from MOZART model simulations (Emmons et al., 2010). Chemical BCs and ICs are not changing along with the sensitivity simulations, but the studied domain (innermost domain) takes boundary conditions from outer main, where emissions are perturbed. The anthropogenic emission inventory used is the MACCity (MACC/CityZEN EU projects) emissions dataset, which provides monthly CO, NO_x, SO_2, VOC, BC, OC, and NH_3 emissions from different sectors for years between 1960 and 2020 (Granier et al., 2011). We compared the MACCity emission inventory for 2010 (Granier et al., 2011) with MIX emission inventory for 2010 (M. Li et al., 2015) in the China region, and the magnitudes of emissions in China from these two datasets are very close. For example, the SO_2 emissions in China in 2010 were estimated to be 28663 Gg in the MIX emission inventory, and were 26876.3 Gg in the MACCity emission inventory. Simulations for evaluating roles of emission changes were conducted using emissions for year 1960 and year 2010. We assigned emissions to the first 6 layers from surface based on sectors. For example, emissions from large point sources (such as chimneys) were assigned to higher layers. Biogenic emissions were estimated online using the MEGAN model (Guenther et al., 2006). The simulation period was January 2010 and five days in previous month were modeled as spin-up to overcome the influences of initial conditions.

2.2 Sensitivity experiments

We explored the sensitivities of the winter time haze event in 2010 to changes in emissions and meteorology features through a series of simulations using 1960 and 2010 emission baselines. Specifically, the influences of emission changes of SO_2, BC, OC, NH_3, and NO_x, and meteorology (temperature, RH and wind speeds) changes on $PM_{2.5}$ and its major species were evaluated using a series of simulations. They are listed and explained in Table 1. All base simulations use meteorology of January 2010. CTL case uses emissions for year 1960 and EMI2010 case uses emissions for year 2010. SO_2, NH_3, and NO_x emissions were perturbed separately from 1960 to 2010 (i.e., SO_2-2010 NH_3-2010 NO_x-2010 cases). In the CTL_NF and EMI2010_NF cases, aerosol-radiation interactions are excluded based on emissions for year 1960 and 2010. It was pointed out surface air temperature in North China increased at the rate of 0.36 ℃ per decade (Guo et al., 2013), the linear trends coefficient of relative humidity anomaly in North China is about -0.60% per decade (Wang et al., 2004), and national mean wind speed decreased 16% in the recent 50 years (Wang et al., 2004). To estimate the impacts of changes in temperature, RH and wind speed that happened in the past several decades, we decreased temperature by 2 degrees, increased RH by 10%, and increased wind speeds by 20%, to reflect conditions of early decades (CTL_T2, CTL_RH10, CTL_WS20, EMI2010_T2, EMI2010_RH10, and EMI2010_WS20 cases). These were conducted by perturbing the initial and boundary conditions of these individual meteorological variables.

At different vertical heights, emission and meteorological variables were uniformly perturbed. The changes of $PM_{2.5}$ and its major components due to perturbations in emissions and meteorology are analyzed for the North China region. The North China region is defined using the innermost domain (shown in Figure S1) and the statistics of changes are calculated within this domain for the January 2010 month.

2.3 Model Verification

The WRF-Chem model performance has been evaluated using multiple observations, including surface meteorological, chemical and optical data, and satellite data in Gao et al. (2016). The model was shown to capture the variations of surface temperature, RH, while wind speed was slightly overestimated (Gao et al., 2016), which has been reported as a common problem of current WRF-Chem model under low wind speed conditions. The Root Mean Square Error (RMSE) of temperature were all less than 3.2K and RMSEs of RH varied from 6.4 to 11.1%. The RMSE of wind speeds were below the proposed criteria (2m/s) (Emery et al., 2001) at the Beijing, Tianjin and Baoding stations, but larger than that criteria at the Chengde station. The time series of simulated surface $PM_{2.5}$, NO_2, and SO_2 showed good agreement with observations as did simulated aerosol optical depth (AOD) (Gao et al., 2016). Mean Fractional Bias (MFB) ranged from -21.8% to 0.4% and Mean Fractional Error (MFE) ranged from 26.3% to 50.7% when comparing against $PM_{2.5}$ observations (Gao et al., 2016). In addition, the comparison between model results and satellite found that the vertical distribution of aerosol and horizontal distribution were captured well by the model (Gao et al., 2016). Compared with observed $PM_{2.5}$ composition, sulfate and OC were underestimated and nitrate was overestimated by the model (Gao et al., 2016). The underestimation of sulfate may be due to underestimation of SO_2 gas phase oxidation, errors in aqueous-phase chemistry, and/or missing heterogeneous sulfate formation (Gao et al., 2016).

3 Results and Discussion

3.1 $PM_{2.5}$ sensitivity to emission changes from 1960 to 2010

The emission changes of SO_2, NO_x, NH_3, BC and OC and resulting impacts on $PM_{2.5}$ from 1960 to 2010 were examined based on the MACCity dataset for years 1960 and 2010. Figure 1(a-e) displays SO_2, NO_x, NH_3, BC and OC emissions for 1960 and Figure 1(f-j) shows the changes from 1960 to 2010. Populated regions of North China, such as urban Beijing, urban Tianjin, and urban Shijiazhuang, exhibit large emissions of SO_2, NO_x, NH_3, BC and OC in 1960. However, NH_3 emissions exhibit different spatial distribution patterns from SO_2, NO_x, BC and OC emissions, because NH_3 is mainly associated with agriculture while SO_2, NO_x, BC and OC are mainly related with industrial and residential activities. From 1960 to 2010, SO_2, NO_x, NH_3, BC and OC increased over the entire North China domain and markedly increased in the Jing-Jin-Ji city cluster. In general, the domain averaged SO_2 emissions and NO_x emissions in North China increased by ~220% and ~990% from 1960 to 2010, respectively. The domain averaged NH_3 emissions in North China increased by ~390% from 1960 to 2010, but the most significant increases occurred not in the Jing-Jin-Ji city cluster, but in Inner Mongolia. Unlike NH_3 emissions, BC emissions increased the most in urban Beijing from 1960 to 2010. This is because residential sources are the biggest contributor to BC in winter (Li et al., 2016) and the population in urban Beijing sharply increased with rapid urbanization. From 1960 to 2010, the mean BC emissions in North China increased by ~154%. Similar to BC emissions, OC emissions increased substantially in the center of Beijing, and the domain averaged increasing ratio is about 54% from 1960 to 2010. The enhancements of SO_2, NO_x, NH_3, BC and OC emissions in North China are expected to result in substantial increase in regional $PM_{2.5}$ concentrations.

Figure 2 shows the simulated monthly mean concentrations of $PM_{2.5}$ and its major components (sulfate, nitrate, ammonium, BC and OC) based on emissions for year 1960. As listed in Table 2, the domain averaged concentrations of sulfate, nitrate, ammonium, BC, OC, and $PM_{2.5}$ are 1.9, 0.8, 0.8, 1.5, 4.6, and 19.2μg/m^3, respectively. For year 1960, $PM_{2.5}$ concentrations are mainly dominated by sulfate, OC and natural dust (the difference between $PM_{2.5}$ and the sum of sulfate, nitrate, ammonium, BC, OC). Figure 3 displays the changes of sulfate, nitrate, ammonium, BC, OC, and $PM_{2.5}$ due to changes in SO_2, NO_x, BC and OC emissions from 1960 to 2010. The predicted monthly mean concentrations of $PM_{2.5}$ components and $PM_{2.5}$ increase everywhere over the entire domain due to emission changes resulting from the rapid urbanization and industrialization from 1960 to 2010 (Figure 3(a-f)). As listed in Table 2, the predicted monthly domain mean sulfate increases the largest (5.0 μg/m^3), followed by nitrate (2.6 μg/m^3) and OC (2.5 μg/m^3).

From 1960 to 2010, the predicted BC increased by ~157% and OC increased by ~54% due to 153% increase in BC emissions and 52% increase in OC emissions. The nearly linear response of both BC and OC aerosols to their emissions is due to the omission of a secondary organic aerosol formation in the chosen CBMZ/MOSAIC mechanism. Thus, both of them were treated as primary aerosols in these simulations. Our previous analyses indicate that SOA contribution in this time period was small (Gao et al., 2016). The domain mean $PM_{2.5}$ concentrations increased by 14.7μg/m^3 and the domain maximum increase is about 45μg/m^3 (Figure 3(f) and Table 2).

To quantify how much of the changes in Figure 3 are from the impacts of boundary conditions, we simulated another case with the innermost domain emissions fixed in 1960 and the outer domain emissions changed from 1960 to 2010. The impacts of boundary conditions mostly occur around the south boundary and show nearly no impact on $PM_{2.5}$ in Beijing (shown in Figure S2), which are consistent with the continuous weak southerly winds during the study period (Gao et al., 2016). On domain average, the impacts of boundary conditions result in 5.0ug/m^3 increase in the study domain, accounting for about 33.9% of the total changes in $PM_{2.5}$."

To explore how emission changes can affect haze days, we calculated the number of haze days in urban Beijing for the CTL and EMI_2010 cases, using daily mean thresholds of 35 and 75μg/m^3 (China National Ambient Air Quality Grade I and Grade II Standard, L. T. Wang et al., 2014). In urban Beijing, there are 4 days when daily mean $PM_{2.5}$ concentrations are above 35μg/m^3, and 0 days with daily mean $PM_{2.5}$ concentrations above 75μg/m^3 for the CTL case. For the EMI_2010 case, these two numbers increase to 15 and 8, indicating that the large increases in emissions over the past several decades have significantly affected haze occurrences in Beijing.

3.2 Sensitivity to changes in individual emission species

The results discussed above show that in the winter period, the concentrations of secondary inorganic aerosols (sulfate, nitrate, and ammonium) has increased dramatically. Thus it is important to explore how sensitive secondary inorganic aerosol is to perturbations in precursor emissions. Three sensitivity simulations (change SO_2, NH_3 and NH_3 emissions separately) were conducted to examine how changes in emissions of each species affect aerosol concentrations. The predicted changes of $PM_{2.5}$ and major $PM_{2.5}$ components at the ground-level are shown in Figure 4 and monthly domain mean aerosol changes are summarized in Table 3.

3.2.1 Changes in SO_2 emissions

Due to changes in SO_2 emissions from 1960 to 2010, domain averaged sulfate increase by 3.4μg/m^3 (178.3%), nitrate decreases by -0.3μg/m^3 (-32.3%), and ammonium increases by 0.2μg/m^3 (29.4%). NH_3 reacts preferentially with SO_2 (Seinfeld and Pandis, 2006) and free NH_3 reacts with enhanced H_2SO_4 due to increasing SO_2. As a result, ammonium increases and less HNO_3 gas is transferred to the aerosol phase, which is consistent with the responses to increasing SO_2 emissions in Kharol et al. (2013).

3.2.2 Changes in NH_3 emissions

As shown in Figure 4 and Table 3, changes in NH_3 emissions from 1960 to 2010 result in significant increases in nitrate (1.5μg/m^3, +76.0%) and ammonium (0.6μg/m^3, +84.0%). The domain mean changes of sulfate due to increase in NH_3 is close to zero (about 0.1μg/m^3), because sulfate formation is only indirectly associated with NH_3 availability (Tsimpidi et al., 2007). The significant changes in nitrate and ammonium occurred in south Hebei, Shandong, and Henan province, where anthropogenic NO_x emissions are very high (Figure 1). Although NH_3 emissions substantially increased in Inner Mongolia (Figure 1), responses of nitrate and ammonium are not significant there due to trivial NO_x emissions. The substantial increases of nitrate after NH_3 emission increase indicate that NH_3 limits the NH_4NO_3 formation in North China region in this period.

3.2.3 Changes in NO_x emissions

After changing NO_x emissions from 1960 to 2010 levels, domain mean surface $PM_{2.5}$ decreases by about 0.2μg/m^3, but the changes of individual $PM_{2.5}$ inorganic components vary. The increase of NO_x emissions cause 0.7μg/m^3 (-39.1%) decrease in monthly domain mean sulfate and the domain peak sulfate reduction is about 2.9μg/m^3. The OH radical is critical in the sulfate formation in the regions where SO_2 concentrations are high and there is a competition between NO_x and VOCs to react with OH (Tsimpidi et al. 2012b). When the VOCs/NO_x concentration ratio is close to 5.5:1, the OH reacts with NO_x and VOCs at an equal rate (Seinfeld and Pandis, 2006). When the concentration ratio is lower than 5.5:1, the OH primarily reacts with NO_x, and the region with this concentration ratio is called VOC-limited region. In VOC-limited regions, an increase of NO_x will cause a decrease of OH and ozone concentration. When the VOCs/NO_x concentration ratio is higher than 5.5:1, the OH will preferentially react with VOCs, and the region with this high ratio is called NO_x-limited region. In the NO_x-limited region, an increase of NO_x will increase OH and ozone concentrations. In the simulated winter month, biogenic emissions are low and NO_x emissions in North China are very high, leading to lower VOCs to NO_x ratios, and it can be considered as VOC-limited region. Fu et al. (2012) pointed out that north East Asia is VOC-limited in January and urban areas of Beijing are VOC-limited in both January and July. The model averaged VOCs/NOx ratio changes from 4.2 to 1.2 due to emission perturbations from 1960 to 2010 (shown in Figure S3). As a result, the large increases in NO_x emissions from 1960 to 2010 result in a 47.9% decrease in daytime surface ozone concentration and 55.6% decrease in daytime surface OH concentration, which are shown in Figure 5. Over the entire domain, ozone and OH decrease due to NO_x emission increases (Figure 5). Consequently, sulfate aerosol decrease over the entire domain, as shown in Figure 4(i). Decreases in sulfate might also be related to changes in thermodynamics of the ammonium-sulfate-nitrate system. Although OH decreases, nitrate still rises (0.6μg/m^3, +76.0%) due to the increase in NO_x emissions. The domain mean ammonium decreases by about 5.1% (-0.04μg/m^3). The net effects of NO_x emission increases bring about 0.2μg/m^3 decrease in monthly domain mean $PM_{2.5}$ concentration and the domain peak decrease is about 1.1μg/m^3 (Table 3).

3.2.4 Coupled changes in SO_2, NH_3 and NO_x emissions

As shown above, increasing SO_2 emissions significantly increases $PM_{2.5}$ concentrations in North China region, increasing NH_3 emissions also increases $PM_{2.5}$ concentrations but to a lesser extent, and increasing NO_x emissions slightly decreases $PM_{2.5}$ concentrations. The effects of coupled changes in SO_2, NH_3 and NO_x emissions are not a simple addition of the effect of changing them separately. As listed in Table 3, the monthly domain mean sulfate, nitrate, ammonium, and $PM_{2.5}$ increases more than the effects of changing emissions separately. Domain mean sulfate increases by 5.0μg/m^3 (+264.0%), nitrate increases by 2.6 μg/m^3 (+322.5%), ammonium increases by 2.3μg/m^3 (295.2%) and $PM_{2.5}$ increases by 9.9μg/m^3 due to coupled changes in SO_2, NH_3 and NO_x emissions from 1960 to 2010. The simultaneous increases in SO_2, NH_3 and NO_x

emissions promote dramatic increases of secondary inorganic aerosols in North China, which is partially due to perturbations in VOCs and other species at the same time

3.2.5 Changes in BC and OC emissions

Since BC and OC are treated as primary aerosols in the chosen CBMZ/MOSAIC mechanism, changes in their emissions do not show any impact on other aerosol components. As listed in Table 3, monthly domain mean $PM_{2.5}$ increases by 2.3μg/m^3 and 2.5μg/m^3 due to changes in their emissions from 1960 to 2010, respectively.

3.3 Effects of temperature increases

The model used in this study is a fully online-coupled model, which simulates meteorological variables and chemical variables together. Therefore, it is not possible to increase temperature uniformly, as was done in previous studies using offline models (Dawson et al., 2007; Megartis et al., 2013; Megartis et al., 2014). To examine the sensitivity of $PM_{2.5}$ to temperature change (reflecting the winter warming trends), we decrease temperature by 2 ℃ in the initial and boundary conditions to reflect conditions more like 1960. As a result, the monthly domain mean surface temperature increases 2.0 ℃ (CTL-CTL_T2), but in a non-uniform manner. These responses in domain temperature are partially due to aerosol feedbacks. The spatial distributions of monthly mean surface temperature and temperature changes are shown in Figure 6(a). The monthly mean surface temperature increases more along top left domain boundaries and less over the Bohai sea. The influence of increasing temperature on biogenic emissions is included using temperature-sensitive biogenic emission model MEGAN (Guenther et al., 2006).

Due to the perturbation in temperature as mentioned above, sulfate, nitrate, ammonium and $PM_{2.5}$ are predicted to increase in most areas of the domain (Figure 7). Predicted monthly mean sulfate increases by $0.06\mu g/m^3$ (+3.1%), nitrate increases by $0.03\mu g/m^3$ (+4.2%), and ammonium increases by $0.02\mu g/m^3$ (+2.8%). The increases of sulfate, nitrate and ammonium are mostly attributed to the increasing OH radicals, as shown in Figure 6(b). After the temperature perturbation, daytime OH increases by about 3.6% on domain average. It was found that higher temperature increased volatilization of ammonium nitrate and partitioned it to the gas phase (Megaritis et al., 2014), but it is not significant here due to the low temperature in winter. In addition, the increase of sulfate, nitrate, and ammonium could be partially due to accelerated gas-phase reaction rate at higher temperature (Dawson et al., 2007; Megaritis et al., 2014). It may also due to enhanced photolysis caused by decreases in cloudiness after temperature perturbations (as shown in Figure S4: changes in liquid water path).

As shown in Figure 7 (d-e), the concentrations of primary aerosols (BC and OC) also increase after the temperature perturbations. This is due to changes in other physical parameter, such as wind direction, wind speed, and PBLHs, which are key factors in the diffusion of air pollutants. Figure 6(c) shows that monthly PBLHs in most North China areas decrease after the temperature perturbation, and PBLHs over the Bohai sea decrease the most, with monthly mean decrease over 50 meters. The monthly domain average daytime PBLHs decrease about 2.3% due to changes in temperature vertical profiles. Surface horizontal winds also change (Figure 6(d)), which directly affect the distributions and magnitudes of $PM_{2.5}$ concentrations in North China along with PBLH changes.

The responses of $PM_{2.5}$ concentrations to temperature perturbation are different from the responses of sulfate, nitrate, ammonium, BC and OC (Figure 7), with decreases in northwestern regions and increases in most areas of the North China Plain. This is because natural dust is dominant in northwestern regions (as shown in Figure 2(f)), and the concentrations of natural dust decrease under lower horizontal wind speeds (Figure 6(d)). The monthly $PM_{2.5}$ concentration decreases by $0.01\mu g/m^3$ on domain average due to temperature perturbation. Because of temperature increase, the numbers of haze days (defined using the daily mean threshold 35 and $75\mu g/m^3$) in urban Beijing do not change.

The discussions shown above are based on emission levels in 1960. The responses to temperature perturbations were also investigated based on emission levels in 2010, and the results are shown in Figure S5, S6 and Table 3. The spatial distributions of the changes are similar to the results shown above, but with larger magnitudes. The domain mean PBL heights decreases slightly more (-8.6 compared to -8.3 meters). The domain mean $PM_{2.5}$ concentrations and $PM_{2.5}$ components exhibit larger increases in North China, although daytime OH concentrations increases less (2.6×10^{-9} compared to 3.3×10^{-9} ppmv), suggesting that the responses of $PM_{2.5}$ concentrations are mostly due to changes in PBL heights and wind fields.

3.4 Effects of RH decreases

The RH was enhanced by 10% in model initial and boundary conditions to represent RH for the previous decades. As a result, the simulated monthly mean RH decreases by 9.3% on domain average. Due to RH perturbation, domain mean $PM_{2.5}$ concentration decreases by $0.7\mu g/m^3$. As shown Figure 8(a), $PM_{2.5}$ concentrations decrease in the Jing-Jin-Ji region but increase in southern areas of the domain. The ammonium nitrate formation equilibrium depends on RH (Tai et al., 2010), so HNO_3 may be shifted to the gas phase under lower RH. In addition, the changes in RH can also affect the wet deposition rate. The increases in southern areas of the domain are mainly due to suppressed in-cloud scavenging, as the decreases in RH inhibit the formation of clouds. As shown in Figure 8(b), liquid water path (LWP) decreases by 75.0%. As a result, the in-cloud scavenging loss rate decreases. The changes of predicted aerosol optical depth at 600nm are shown in Figure 8(c). In most regions, visibility decreases due to lower RH. Because of RH decreases, the numbers of haze days (defined using the daily mean threshold 35 and 75μg/m^3) in urban Beijing do not change. The responses to RH perturbations were also investigated based on emission levels in 2010, and the results are shown in Figure S7 and Table 3. The responses are also similar to changes based on emission levels in 1960, but with larger magnitudes.

3.5 Effects of wind speed decreases

Simulations were also carried out when wind speeds in initial and boundary conditions were increased to estimate the wind speeds for the previous decades. The predicted domain averaged monthly mean wind speed decreases by about 0.7 m/s. As shown in Figure 9(a), the monthly mean near surface horizontal winds are pronounced in mountainous areas (northwest areas of the domain) and relatively smaller in other areas. Figure 9(b) shows the changes of wind speeds (CTL-WS20) due to model perturbations. The predicted monthly mean $PM_{2.5}$ concentrations decrease by 2.3μg/m^3 on domain average, but the responses of $PM_{2.5}$ vary within the domain. As shown in Figure 9(c), $PM_{2.5}$ concentrations decrease in the northwestern areas because of lower production of natural dust under lower horizontal wind speeds. However, in most areas of the North China Plain, $PM_{2.5}$ concentrations increase under lower wind speeds (Figure 9(c)). The domain peak increase is about 2.4 μg/m^3, which is based on low predicted $PM_{2.5}$ concentrations using emissions for year 1960. If the concentration in base case is higher, the responses will be enhanced. As shown in Figure 9(d), the domain maximum increases in $PM_{2.5}$ increases from 2.4 to 9.4 μg/m^3. Because of wind speed decreases, number of haze days that daily mean $PM_{2.5}$ concentrations are above 35μg/m^3 increases by 1.

3.6 Effects of changes in aerosol feedbacks

As mentioned in Gao et al. (2016), high concentrations of aerosol enhance stability of boundary layer and increase $PM_{2.5}$ concentrations. Due to dramatic changes in emissions from 1960 to 2010, the strength of aerosol feedbacks may also have changed. To quantify these changes, we simulated four cases (i.e., CTL, CTL_NF, EMI2010, and EMI2010_NF). CTL-CLT_NF and EMI2010-EMI2010_NF are used to represent the contributions of aerosol radiative effects in 1960 and 2010. The changes in monthly mean daytime PBL heights and $PM_{2.5}$ concentrations are shown in Figure 10. In 1960, the domain averaged PBL height decreases by 6.7 meters due to aerosol radiative effects, and the domain maximum decrease is 25.4 meters. Correspondingly, the domain averaged $PM_{2.5}$ increases by 0.1 μg/m^3 and the domain maximum increase is 0.9 μg/m^3. In 2010, the domain averaged PBL height decreases by 13.8 meters and the domain maximum decrease is 55.2 meters (more than two times compared to 1960). Correspondingly, the domain averaged $PM_{2.5}$ increases by 0.7 μg/m^3 and the domain maximum increase is 5.1 μg/m^3. The enhanced strength of aerosol feedbacks is another important cause of degraded aerosol pollution. Thus, controlling emissions will have a co-benefit of reducing strength of aerosol feedbacks.

3.7 Implications for the effects of emission and meteorology changes on $PM_{2.5}$ concentrations

The simulated responses of $PM_{2.5}$ concentrations to emission changes and meteorology changes presented here, along with the previous presented effects of aerosol feedbacks (Gao et al. 2016), provide important implications for the causes of the dramatic increases in winter $PM_{2.5}$ concentrations.

We calculated domain maximum changes in $PM_{2.5}$ concentration averaged over four stagnant days (January 16-19) owing to emission changes from 1960-2010 (EMI2010-CTL), temperature increases (CTL-CTL_T2), RH decreases (CTL-CTL_RH10), wind speed decreases (CTL-CTL_RH20), and aerosol feedbacks (CTL-CTL_NF). The values are 137.7, 2.0, 2.6, 7.5 and 4.0μg/m^3, respectively. When the perturbations are based on emission levels in 2010, domain maximum changes in $PM_{2.5}$ concentration due to temperature increases (EMI2010-EMI2010_T2), RH decreases (EMI2010-EMI2010_RH10), wind speed decreases (EMI2010-EMI2010_WS20), and aerosol feedbacks (EMI2010-EMI2010_NF) are 4.8, 4.7, 26.4 and 25.5μg/m^3. The effects of emission changes on haze formation are dominant and the effects of aerosol feedbacks are comparable to the effects of wind speed decreases.

The comprehensive comparisons of these factors are also summarized in Table 3. Based on the monthly domain mean responses of $PM_{2.5}$ concentrations to these factors, dramatic emission changes due to urbanization and industrialization are the main causes of degraded air quality and frequent haze occurrences in in North China. $PM_{2.5}$ shows significant responses to changes in SO_2, NH_3, NO_x emissions than BC and OC (about 106.3% higher). In addition, $PM_{2.5}$ shows significant increases in response to changes in SO_2 and NH_3 emissions, as compared to increases in response to changes in NO_x emissions. This region is relatively ammonia-poor in winter, so reducing NH_3 emissions might be effective, which is consistent with previous findings in Europe (Megaritis et al., 2013). SO_2 is the precursor of sulfate, which accounts for a large fraction of PM in this region. Thus, they should be preferentially controlled in order to reduce $PM_{2.5}$ levels. To control SO_2 emissions, the usage of natural gas or other clean energy should be promoted to reduce the usage of coal. NH_3 emissions in China are mainly from agriculture sources (about 90%), including livestock, fertilizer, and agricultural soil (Huang et al., 2012). Lelieveld et al. (2015) found that agricultural emissions make the largest relative role in $PM_{2.5}$ concentration in eastern USA, Europe, Russia and East Asia. To control NH_3 emissions from agriculture sources, some animal feeding and animal housing strategies should be taken. In addition, controlling emissions will also have a co-benefit of reducing strength of aerosol feedbacks.

According to the ECLIPSE_GAINS_4a emission dataset, SO_2 emissions in China will decrease by -26%, NO_x emissions in China will increase by 19%, and NH_3 emissions in China will increase by 14% from 2010 to 2030. We predicted (by perturbating SO_2, NO_x and NH_3 emissions by -26%, 19% and 14%) that these changes will lead to large decreases in winter sulfate (-2.3μg/m^3 on domain average). Nitrate will increase by 1.5μg/m^3 and ammonium will slightly decrease (-0.05μg/m^3) on domain average. The net change of domain averaged $PM_{2.5}$ concentration is not significant (-0.8μg/m^3), so more efforts are needed to control these important gaseous precursors.

From the information listed in Table 3, the responses of $PM_{2.5}$ concentrations to temperature and RH perturbations are not as significant as to wind speed perturbations. From Sect. 3.3, we also found that the effects of temperature perturbation on $PM_{2.5}$ concentration are dominant by changes in PBLH and wind fields. Previous studies have pointed out the occurrences of haze events are highly associated with atmospheric circulation anomalies (Chen and Wang, 2015;

Zhang et al., 2016). Thus, changes in atmospheric circulations may be another important cause of growing haze pollution, in addition to emission changes. Furthermore, aerosol can also change atmospheric circulation, especially in severely polluted East Asia. Thus, controlling emission may have co-benefits of mitigate aerosol effects on atmospheric circulation.

The effects of changing atmospheric circulations on winter haze pollution in China is beyond the scope of this paper, but should be investigated in future studies.

4 Summary

A fully online coupled meteorological and chemical transport model, WRF-Chem was used to study responses of winter $PM_{2.5}$ concentrations to changes in emissions of SO_2, BC, OC, NH_3, and NO_x and to meteorology (temperature, RH, and wind speeds) changes in North China region, where people are suffering from severe winter haze pollution.

The detailed historical emissions dataset MACCity for year 1960 and 2010 were used to evaluate the impacts of changes in emissions of SO_2, BC, and OC. From 1960 to 2010, the dramatic changes in emissions lead to +264.0% increases in sulfate, +322.5% increases in nitrate, +295.2% increases in ammonium, +157.0% increases in BC and 54% increases in OC. The domain mean $PM_{2.5}$ concentrations increase by 14.7$\mu g/m^3$ and the domain maximum increase is about 45$\mu g/m^3$. The responses of $PM_{2.5}$ to individual emission species indicate that the simultaneous increases in SO_2, NH_3 and NO_x emissions dominated the increases in $PM_{2.5}$ concentrations. $PM_{2.5}$ shows significant increases in response to SO_2 and NH_3 emission changes. The increases in NO_x emissions may decrease surface ozone concentration and surface OH radical concentrations, because North China region is VOC-limited in the winter. In addition, OC accounts for a large fraction in $PM_{2.5}$ changes.

The sensitivities of $PM_{2.5}$ to emission changes of its precursors provide some implications for haze pollution control. SO_2, NH_3 and OC should be preferentially controlled. In China, the residential sector, particularly biofuel usage is the primary sources of OC (Lu et al., 2011). The usage of natural gas or other clean energy should be promoted to reduce the usage of coal and biofuel to reduce SO_2 and OC. To control NH_3 emissions from agriculture sources, some animal feeding and animal housing strategies should be taken.

The effects of changes in winter time meteorology conditions were also studied. Emission changes from 1960 to 2010 substantially increase numbers of haze days, but meteorology perturbations do not show any significant impacts. The perturbations in temperature and RH do change $PM_{2.5}$ concentrations, but the strength is not as significant as the effects of wind speed and emission changes. The effects of temperature perturbation are dominated by the changes in surface wind fields and PBLHs. The effect of aerosol feedbacks is comparable to the effect of decreasing wind speeds and the strength of aerosol feedbacks significantly increased from 1960 to 2010.

The above discussions indicate that aerosol concentrations are mainly controlled by atmospheric circulations, except emission changes. Thus, long-term trends in atmospheric circulations maybe another important cause of winter haze events in North China. More studies are necessary to get a better understanding of the aerosol-circulation interactions.

In our previous modeling study of the same period (January 2010), we found that SOA contribution was small, so we did not include SOA in this study. But this indication might be problematic due to current poorly parameterized SOA scheme. In the future, how changes in emissions and meteorology variables affect productions of SOA during winter should be further studied using more advanced SOA schemes. In addition, we did not consider primary PM except BC and OC in the model because there is no information in the MACCity emission inventory, which is another direction for improvements in future studies.

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References

Aksoyoglu, S., Keller, J., Barmpadimos, I., Oderbolz, D., Lanz, V. a., Prévăt, a. S. H. and Baltensperger, U.: Aerosol modelling in Europe with a focus on Switzerland during summer and winter episodes, Atmos. Chem. Phys., 11, 7355–7373, doi:10.5194/acp-11-7355-2011, 2011.

Andreani-Aksoyoglu, S., Keller, J., Prévôt, A. S. H., Baltensperger, U. and Flemming, J.: Secondary aerosols in Switzerland and northern Italy: Modeling and sensitivity studies for summer 2003, J. Geophys. Res. Atmos., 113, 1–12, doi:10.1029/2007JD009053, 2008.

Aw, J.: Evaluating the first-order effect of intraannual temperature variability on urban air pollution, J. Geophys. Res., 108, doi:10.1029/2002JD002688, 2003.

Bowman, F. M. and Karamalegos, A. M.: Estimated effects of composition on secondary organic aerosol mass concentrations., Environ. Sci. Technol., 36(11), 2701–2707, 2002.

Cheung, H. C., Wang, T., Baumann, K. and Guo, H.: Influence of regional pollution outflow on the concentrations of fine particulate matter and visibility in the coastal area of southern China, Atmos. Environ., 39, 6463–6474, doi:10.1016/j.atmosenv.2005.07.033, 2005.

Dawson, J. P., Adams, P. J. and Pandis, S. N.: Sensitivity of PM2.5 to climate in the Eastern U.S.: a modeling case study, Atmos. Chem. Phys., 7, 4295–4309, 2007.

Dawson, J. P., Bloomer, B. J., Winner, D. a. and Weaver, C. P.: Understanding the meteorological drivers of U.S. particulate matter concentrations in a changing climate, Bull. Am. Meteorol. Soc., 95(April), 521–532, doi:10.1175/BAMS-D-12-00181.1, 2014.

Emery, Chris;Tai, Edward;Yarwood, G.: Enhanced Meteorological Modeling and Performance Evaluation for Two Texas Ozone Episodes, in Prepared for the Texas Natural Resource Conservation Commission, ENVIRON International Corporation, Novato, CA., 2001.

Emmons, L. K., Walters, S., Hess, P. G., Lamarque, J.-F., Pfister, G. G., Fillmore, D., Granier, C., Guenther, a., Kinnison, D., Laepple, T., Orlando, J., Tie, X., Tyndall, G., Wiedinmyer, C., Baughcum, S. L. and Kloster, S.: Description and evaluation of the Model for Ozone and Related chemical Tracers, version 4 (MOZART-4), Geosci. Model Dev., 3(1), 43–67, doi:10.5194/gmd-3-43-2010, 2010.

Fu, G. Q., Xu, W. Y., Yang, R. F., Li, J. B. and Zhao, C. S.: The distribution and trends of fog and haze in the North China Plain over the past 30 years, Atmos. Chem. Phys., 14, 11949–11958, doi:10.5194/acp-14-11949-2014, 2014.

Fu, J. S., Dong, X., Gao, Y., Wong, D. C. and Lam, Y. F.: Sensitivity and linearity analysis of ozone in East Asia: The effects of domestic emission and intercontinental transport, J. Air Waste Manage. Assoc., 62(March 2015), 1102–1114, doi:10.1080/10962247.2012.699014, 2012.

Gao, M., Guttikunda, S. K., Carmichael, G. R., Wang, Y., Liu, Z., Stanier, C. O., Saide, P. E. and Yu, M.: Health impacts and economic losses assessment of the 2013 severe haze event in Beijing area, Sci. Total Environ., 511(January 2013), 553–561, doi:10.1016/j.scitotenv.2015.01.005, 2015.

Gao, M., Carmichael, G. R., Wang, Y., Saide, P. E., Yu, M., Xin, J., Liu, Z. and Wang, Z.: Modeling study of the 2010 regional haze event in the North China Plain, Atmos. Chem. Phys., 16(3), 1673–1691, doi:10.5194/acp-16-1673-2016, 2016.

Granier, C., Bessagnet, B., Bond, T., D'Angiola, A., Denier van der Gon, H., Frost, G. J., Heil, A., Kaiser, J. W., Kinne, S., Klimont, Z., Kloster, S., Lamarque, J.-F., Liousse, C., Masui, T., Meleux, F., Mieville, A., Ohara, T., Raut, J.-C., Riahi, K., Schultz, M. G., Smith, S. J., Thompson, A., van Aardenne, J., van der Werf, G. R. and van Vuuren, D. P.: Evolution of anthropogenic and biomass burning emissions of air pollutants at global and regional scales during the 1980–2010 period, Clim. Change, 109(1-2), 163–190, doi:10.1007/s10584-011-0154-1, 2011.

Guenther, A., Karl, T., Harley, P., Wiedinmyer, C., Palmer, P. I. and Geron, C.: Estimates of global terrestrial isoprene emissions using MEGAN (Model of Emissions of Gases and Aerosols from Nature), Atmos. Chem. Phys., 6(1), 107–173, doi:10.5194/acpd-6-107-2006, 2006.

Guo, W.-L., Hong-Bo, S., Jing-Jin, M., Ying-Juan, Z., Ji, W., Wen-Jun, S. and Zi-Yin, Z.: Basic Features of Climate Change in North China during 1961–2010, Adv. Clim. Chang. Res., 4(2), 73–83, doi:10.3724/SP.J.1248.2013.073, 2013.

Heald, C. L., Henze, D. K., Horowitz, L. W., Feddema, J., Lamarque, J. F., Guenther, a., Hess, P. G., Vitt, F., Seinfeld, J. H., Godstein, a. H. and Fung, I.: Predicted change in global secondary organic aerosol concentrations in response to future climate, emissions, and land use change, J. Geophys. Res. Atmos., 113, 1–16, doi:10.1029/2007JD009092, 2008.

Hu, Z.-Z.: Long-term climate variations in China and global warming signals, J. Geophys. Res., 108, 1–13, doi:10.1029/2003JD003651, 2003.

Huang, X., Song, Y., Li, M., Li, J., Huo, Q., Cai, X., Zhu, T., Hu, M. and Zhang, H.: A high-resolution ammonia emission inventory in China, Global Biogeochem. Cycles, 26(3), n/a–n/a, doi:10.1029/2011GB004161, 2012.

Jacob, D. J. and Winner, D. a.: Effect of climate change on air quality, Atmos. Environ., 43(1), 51–63, doi:10.1016/j.atmosenv.2008.09.051, 2009.

Jia, B., Wang, Y., Yao, Y. and Xie, Y.: A new indicator on the impact of large-scale circulation on wintertime particulate matter pollution over China, Atmos. Chem. Phys., 15, 11919–11929, doi:10.5194/acp-15-11919-2015, 2015.

Jiménez-Guerrero, P., Montávez, J. P., Gómez-Navarro, J. J., Jerez, S. and Lorente-Plazas, R.: Impacts of climate change on ground level gas-phase pollutants and aerosols in the Iberian Peninsula for the late XXI century, Atmos. Environ., 55, 483–495, doi:10.1016/j.atmosenv.2012.02.048, 2012.

Kharol, S. K., Martin, R. V., Philip, S., Vogel, S., Henze, D. K., Chen, D., Wang, Y., Zhang, Q. and Heald, C. L.: Persistent sensitivity of Asian aerosol to emissions of nitrogen oxides, Geophys. Res. Lett., 40(February), 1021–1026, doi:10.1002/grl.50234, 2013.

Lelieveld, J., Evans, J. S., Fnais, M., Giannadaki, D. and Pozzer, A.: The contribution of outdoor air pollution sources to premature mortality on a global scale, Nature, doi:10.1038/nature15371, 2015.

Leng, C., Duan, J., Xu, C., Zhang, H., Zhang, Q., Wang, Y., Li, X., Kong, L., Tao, J., Cheng, T., Zhang, R. and Chen, J.: Insights into a historic severe haze weather in Shanghai: synoptic situation, boundary layer and pollutants, Atmos. Chem. Phys. Discuss., 15, 32561–32605, doi:10.5194/acpd-15-32561-2015, 2015.

Li, C., Martin, R. V., Boys, B. L., van Donkelaar, a. and Ruzzante, S.: Evaluation and application of multi-decadal visibility data for trend analysis of atmospheric haze, Atmos. Chem. Phys. Discuss., 15, 33789–33841, doi:10.5194/acpd-15-33789-2015, 2015.

Li, M., Zhang, Q., Kurokawa, J., Woo, J.-H., He, K. B., Lu, Z., Ohara, T., Song, Y., Streets, D. G., Carmichael, G. R., Cheng, Y. F., Hong, C. P., Huo, H., Jiang, X. J., Kang, S. C., Liu, F., Su, H., and Zheng, B.: MIX: a mosaic Asian anthropogenic emission inventory for the MICS-Asia and the HTAP projects, Atmos. Chem. Phys. Discuss., 15, 34813-34869, doi:10.5194/acpd-15-34813-2015, 2015.

Li, K., Liao, H., Mao, Y. and Ridley, D. a.: Source sector and region contributions to concentration and direct radiative forcing of black carbon in China, Atmos. Environ., 124, 351–366, doi:10.1016/j.atmosenv.2015.06.014, 2016.

Liu, Y., Fan, J., Zhang, G. J., Xu, K. and Ghan, S. J.: Air pollution and climate response to aerosol direct radiative effects: A modeling study of decadal trends across the northern hemisphere, J. Geophys. Res. Atmos., 120(23), 12,221–12,236, doi:10.1002/2014JD022145.Received, 2015.

Lu, Z., Zhang, Q. and Streets, D. G.: Sulfur dioxide and primary carbonaceous aerosol emissions in China and India, 1996-2010, Atmos. Chem. Phys., 11, 9839–9864, doi:10.5194/acp-11-9839-2011, 2011.

Megaritis, a G., Fountoukis, C., Charalampidis, P. E., Denier van der Gon, H. a C., Pilinis, C. and Pandis, S. N.: Linking climate and air quality over Europe: effects of meteorology on PM2.5 concentrations, Atmos. Chem. Phys., 14, 10283–10298, doi:10.5194/acpd-14-10345-2014, 2014.

Megaritis, a. G., Fountoukis, C., Charalampidis, P. E., Pilinis, C. and Pandis, S. N.: Response of fine particulate matter concentrations to changes of emissions and temperature in Europe, Atmos. Chem. Phys., 13, 3423–3443, doi:10.5194/acp-13-3423-2013, 2013.

Mu, Q. and Liao, H.: Simulation of the interannual variations of aerosols in China: role of variations in meteorological parameters, Atmos. Chem. Phys., 14, 9597–9612, 2014.

Pay, M. T., Jiménez-Guerrero, P. and Baldasano, J. M.: Assessing sensitivity regimes of secondary inorganic aerosol formation in Europe with the CALIOPE-EU modeling system, Atmos. Environ., 51, 146–164, doi:10.1016/j.atmosenv.2012.01.027, 2012.

Petäjä, T., Järvi, L., Kerminen, V., Ding, a J., Sun, J. N., Nie, W. and Kujansuu, J.: Enhanced air pollution via aerosol- boundary layer feedback in China, Sci. Rep., 6(18998), doi:10.1038/srep18998, 2016.

Pope, C. A., Ezzati, M. and Dockery, D. W.: Fine-Particulate Air Pollution and Life Expectancy in the United States, N. Engl. J. Med., 2009.

Ren, G., Ding, Y., Zhao, Z., Zheng, J. and Wu, T.: Recent progress in studies of climate change in China, Adv. Atmos. …, 29(5), 958–977, doi:10.1007/s00376-012-1200-2.1.Introduction, 2012.

Seinfeld, John H.; Pandis, S. N.: Atmospheric chemistry and physics: from air pollution to climate chnage, John Wiley & Sons., 2012.

Shi, P.-J., Zhang, G.-F., Kong, F. and Ye, Q.: Wind speed change regionalization in China (1961–2012), Adv. Clim. Chang. Res., 6(2), 151–158, doi:10.1016/j.accre.2015.09.006, 2015.

Song, Y., Liu, Y. and Ding, Y.: A Study of Surface Humidity Changes in China During the Recent 50 Years, Acta Meteorol. Sin., (973), 541–553, doi:10.1007/s13351-012-0501-9.1., 2012.

Steiner, A. L., Tonse, S., Cohen, R. C., Goldstein, A. H. and Harley, R. a.: Influence of future climate and emissions on regional air quality in California, J. Geophys. Res. Atmos., 111, 1–22, doi:10.1029/2005JD006935, 2006.

Tagaris, E., Manomaiphiboon, K., Liao, K. J., Leung, L. R., Woo, J. H., He, S., Amar, P. and Russell, A. G.: Impacts of global climate change and emissions on regional ozone and fine particulate matter concentrations over the United States, J. Geophys. Res. Atmos., 112, doi:10.1029/2006JD008262, 2007.

Tai, a. P. K., Mickley, L. J. and Jacob, D. J.: Impact of 2000-2050 climate change on fine particulate matter (PM 2.5) air quality inferred from a multi-model analysis of meteorological modes, Atmos. Chem. Phys., 12, 11329–11337, doi:10.5194/acp-12-11329-2012, 2012a.

Tai, a. P. K., Mickley, L. J., Jacob, D. J., Leibensperger, E. M., Zhang, L., Fisher, J. a. and Pye, H. O. T.: Meteorological modes of variability for fine particulate matter (PM2.5) air quality in the United States: implications for PM2.5 sensitivity to climate change, Atmos. Chem. Phys., 12, 3131–3145, doi:10.5194/acpd-11-31031-2011, 2012b.

Tai, A. P. K., Mickley, L. J. and Jacob, D. J.: Correlations between fine particulate matter (PM2.5) and meteorological variables in the United States: Implications for the sensitivity of PM2.5 to climate change, Atmos. Environ., 44(32), 3976–3984, doi:10.1016/j.atmosenv.2010.06.060, 2010.

Tsigaridis, K. and Kanakidou, M.: Secondary organic aerosol importance in the future atmosphere, Atmos. Environ., 41, 4682–4692, doi:10.1016/j.atmosenv.2007.03.045, 2007.

Tsimpidi, A. P., Karydis, V. a and Pandis, S. N.: Response of inorganic fine particulate matter to emission changes of sulfur dioxide and ammonia: the eastern United States as a case study., J. Air Waste Manag. Assoc., 57(November 2014), 1489–1498, doi:10.3155/1047-3289.57.12.1489, 2007.

Tsimpidi, A. P., Karydis, V. a and Pandis, S. N.: Response of fine particulate matter to emission changes of oxides of nitrogen and anthropogenic volatile organic compounds in the eastern

United States., J. Air Waste Manag. Assoc., 58(November 2014), 1463–1473, doi:10.3155/1047-3289.58.11.1463, 2008.

Wang, H. J. and Chen, H. P.: Understanding the Recent Trend of Haze Pollution in Eastern China: Roles of Climate Change, Atmos. Chem. Phys. Discuss., (January), 1–18, doi:10.5194/acp-2015-1009, 2016.

Wang, J., Wang, S., Jiang, J., Ding, A., Zheng, M., Zhao, B., Wong, D. C., Zhou, W., Zheng, G., Wang, L., Pleim, J. E. and Hao, J.: Impact of aerosol–meteorology interactions on fine particle pollution during China's severe haze episode in January 2013, Environ. Res. Lett., 9(9), 094002, doi:10.1088/1748-9326/9/9/094002, 2014a.

Wang, L. T., Wei, Z., Yang, J., Zhang, Y., Zhang, F. F., Su, J., Meng, C. C. and Zhang, Q.: The 2013 severe haze over southern Hebei, China: model evaluation, source apportionment, and policy implications, Atmos. Chem. Phys., 14(6), 3151–3173, doi:10.5194/acp-14-3151-2014, 2014b.

Wang, Z., Ding, Y., He, J. and Yu, J.: An Updating analysis of the climate change in China in recent 50 years, Acta Meteorol. Sin., 62(228-236), doi:10.1017/CBO9781107415324.004, 2004.

Yang, Y., Wang, J., Gong, S., Zhang, X., Wang, H., Wang, Y., Li, D. and Guo, J.: PLAM – a meteorological pollution index for air quality and its applications in fog-haze forecasts in north China, Atmos. Chem. Phys. Discuss., 16, 13453–1364, doi:10.5194/acpd-15-9077-2015, 2016.

Zaveri, R. a. and Peters, L. K.: A new lumped structure photochemical mechanism for large-scale applications, J. Geophys. Res., 104(D23), 30387, doi:10.1029/1999JD900876, 1999.

Zaveri, R. a., Easter, R. C., Fast, J. D. and Peters, L. K.: Model for Simulating Aerosol Interactions and Chemistry (MOSAIC), J. Geophys. Res., 113(D13), D13204, doi:10.1029/2007JD008782, 2008.

Zhang, B., Wang, Y. and Hao, J.: Simulating aerosol–radiation–cloud feedbacks on meteorology and air quality over eastern China under severe haze conditionsin winter, Atmos. Chem. Phys., 15, 2387–2404, doi:10.5194/acp-15-2387-2015, 2015a.

Zhang, Q., Streets, D. G., He, K., Wang, Y., Richter, A., Burrows, J. P., Uno, I., Jang, C. J., Chen, D., Yao, Z. and Lei, Y.: NOx emission trends for China, 1995 - 2004: The view from the ground and the view from space, J. Geophys. Res. Atmos., 112(x), 1995–2004, doi:10.1029/2007JD008684, 2007.

Zhang, X. Y., Wang, J. Z., Wang, Y. Q., Liu, H. L., Sun, J. Y. and Zhang, Y. M.: Changes in chemical components of aerosol particles in different haze regions in China from 2006 to 2013 and contribution of meteorological factors, Atmos. Chem. Phys., 15, 12935–12952, doi:10.5194/acp-15-12935-2015, 2015b.

Zhang, Z., Zhang, X., Gong, D., Kim, S.-J., Mao, R. and Zhao, X.: Possible influence of atmospheric circulations on winter hazy pollution in Beijing-Tianjin-Hebei region, northern China, Atmos. Chem. Phys., 16, 561–571, doi:10.5194/acpd-15-22493-2015, 2016.

Zhao, X. J., Zhao, P. S., Xu, J., Meng,, W., Pu, W. W., Dong, F., He, D. and Shi, Q. F.: Analysis of a winter regional haze event and its formation mechanism in the North China Plain, Atmos. Chem. Phys., 13(11), 5685–5696, doi:10.5194/acp-13-5685-2013, 2013.

Zheng, B., Zhang, Q., Zhang, Y., He, K. B., Wang, K., Zheng, G. J., Duan, F. K., Ma, Y. L. and Kimoto, T.: Heterogeneous chemistry: a mechanism missing in current models to explain secondary inorganic aerosol formation during the January 2013 haze episode in North China, Atmos. Chem. Phys., 15, 2031–2049, 2015.

Table 1. Simulation cases and descriptions

Cases	Descriptions
CTL	Base case, anthropogenic emissions are from MACCity dataset for year 1960
EMI2010	Anthropogenic emissions are from MACCity dataset for year 2010
SO_2-2010	Same as CTL case except SO_2 emissions are for year 2010
NH_3-2010	Same as CTL case except NH_3 emissions are for year 2010
NO_x-2010	Same as CTL case except NO_x emissions are for year 2010
CTL_T2	Same as CTL case except temperature BCs and ICs are decreased by 2K
CTL_RH10	Same as CTL case except RH BCs and ICs are increased by 10%
CTL_WS20	Same as CTL case except wind speed BCs and ICs are increased by 20%
CTL_NF	Same as CTL case except aerosol-radiation interactions are excluded
EMI2010_T2	Same as EMI2010 case except temperature BCs and ICs are decreased by 2K
EMI2010_RH10	Same as EMI2010 case except RH BCs and ICs are increased by 10%
EMI2010_WS20	Same as EMI2010 case except wind speed BCs and ICs are increased by 20%
EMI2010_NF	Same as EMI2010 case except aerosol-radiation interactions are excluded

Table 2. Monthly domain mean concentrations of $PM_{2.5}$ and its major components for year 1960, and domain maximum and mean concentrations for changes from 1960 to 2010 due to emission changes ($\mu g/m^3$)

Years		SO_4^{2-}	NO_3^-	NH_4^+	BC	OC	$PM_{2.5}$
1960	Domain mean	1.9	0.8	0.8	1.5	4.6	19.2
1960-2010	Domain maximum	18.9	7.8	6.8	9.9	11.1	45.0
	Domain mean	5.0 (264.0%)	2.6 (322.5%)	2.3 (295.2%)	2.3 (156.6%)	2.5 (54.0%)	14.7 (76.4%)

Table 3. Monthly domain mean changes of sulfate, nitrate, ammonium and $PM_{2.5}$ concentrations (μg/m^3) due to emission and meteorology perturbations, and aerosol feedbacks (the two values of $PM_{2.5}$ changes are for meteorology perturbations and aerosol feedbacks based on 1960 and 2010 emission levels, respectively)

	$SO4^{2-}$	NO^{3-}	NH^{4+}	$PM_{2.5}$
Changes in SO_2 emissions	3.4(178.3%)	-0.3 (-32.3%)	0.2 (29.4%)	3.4
Changes in NH_3 emissions	0.1 (5.3%)	1.5 (189.6%)	0.6 (84.0%)	2.3
Changes in NO_x emissions	-0.7 (-39.1%)	0.6 (76.0%)	-0.04 (-5.1%)	-0.2
Changes in SO_2, NH_3, NO_x emissions	5.0 (264.0%)	2.6 (322.5%)	2.3 (295.2%)	9.9
Changes in BC emissions	-	-	-	2.3
Changes in OC emissions	-	-	-	2.5
Temperature perturbations	-	-	-	-0.01/0.3
RH perturbations	-	-	-	-0.7/-1.1
Wind speed perturbations	-	-	-	-2.3/-0.5
Aerosol feedbacks				0.1/0.7

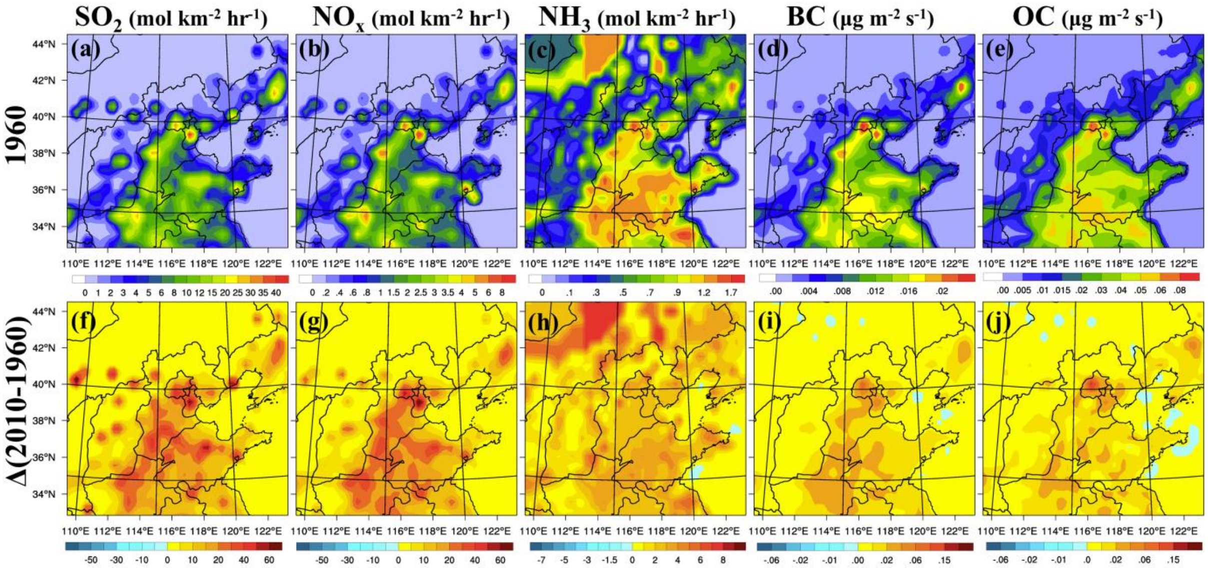

Figure 1. SO_2, NO_x, NH_3, BC and OC emissions for year 1960 (a-e), and the changes of them from 1960 to 2010 (f-j)

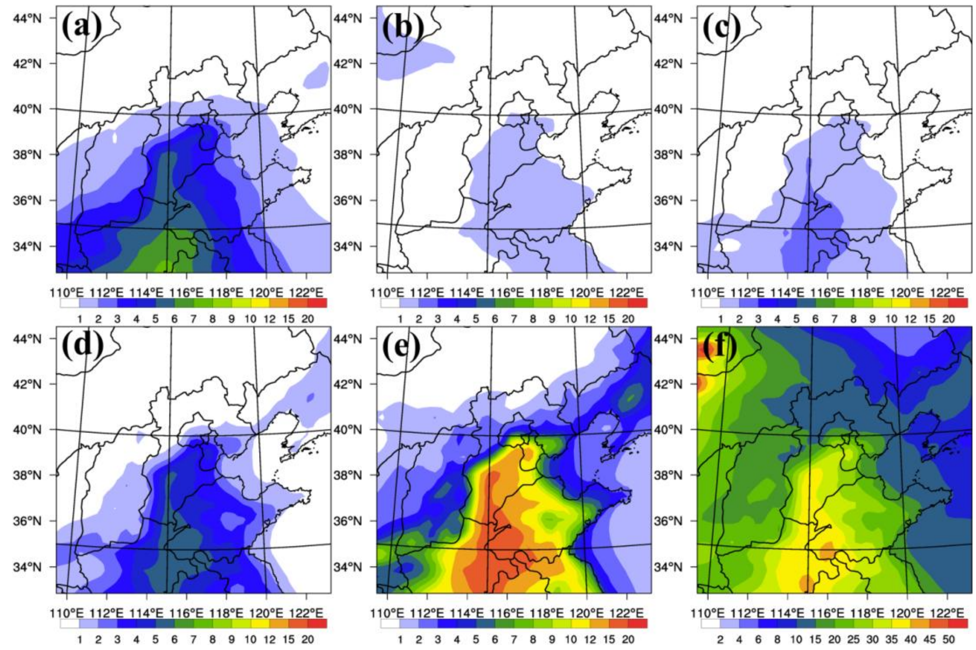

Figure 2. Predicted monthly mean sulfate (a), nitrate (b), ammonium (c), BC (d), OC (e) and $PM_{2.5}$ (f) concentrations based on emissions for year 1960

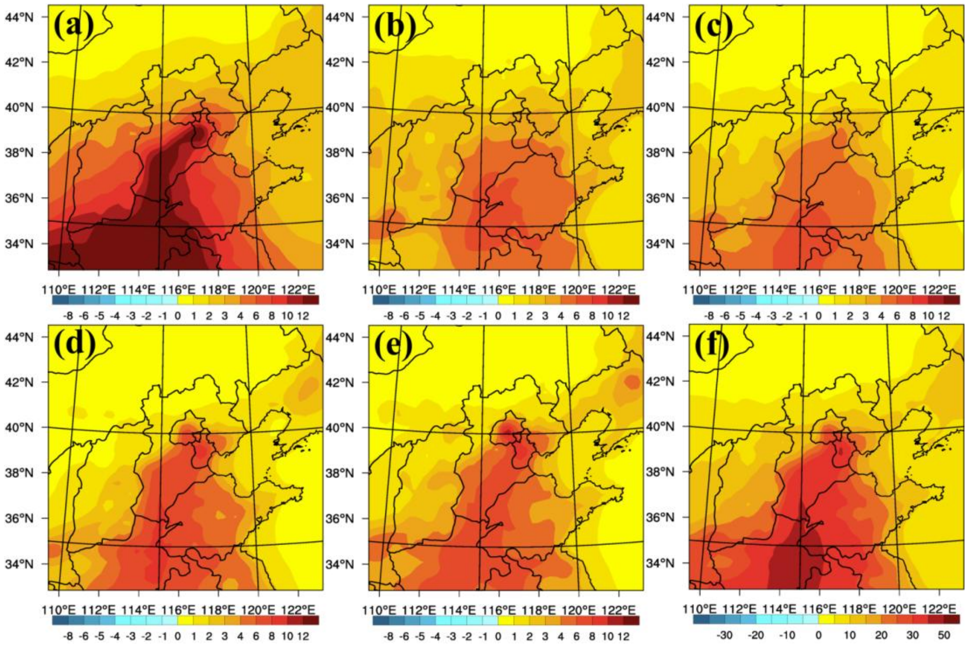

Figure 3. Predicted monthly mean changes of sulfate (a), nitrate (b), ammonium (c), BC (d), OC (e) and $PM_{2.5}$ (f) due to emission changes from 1960 to 2010

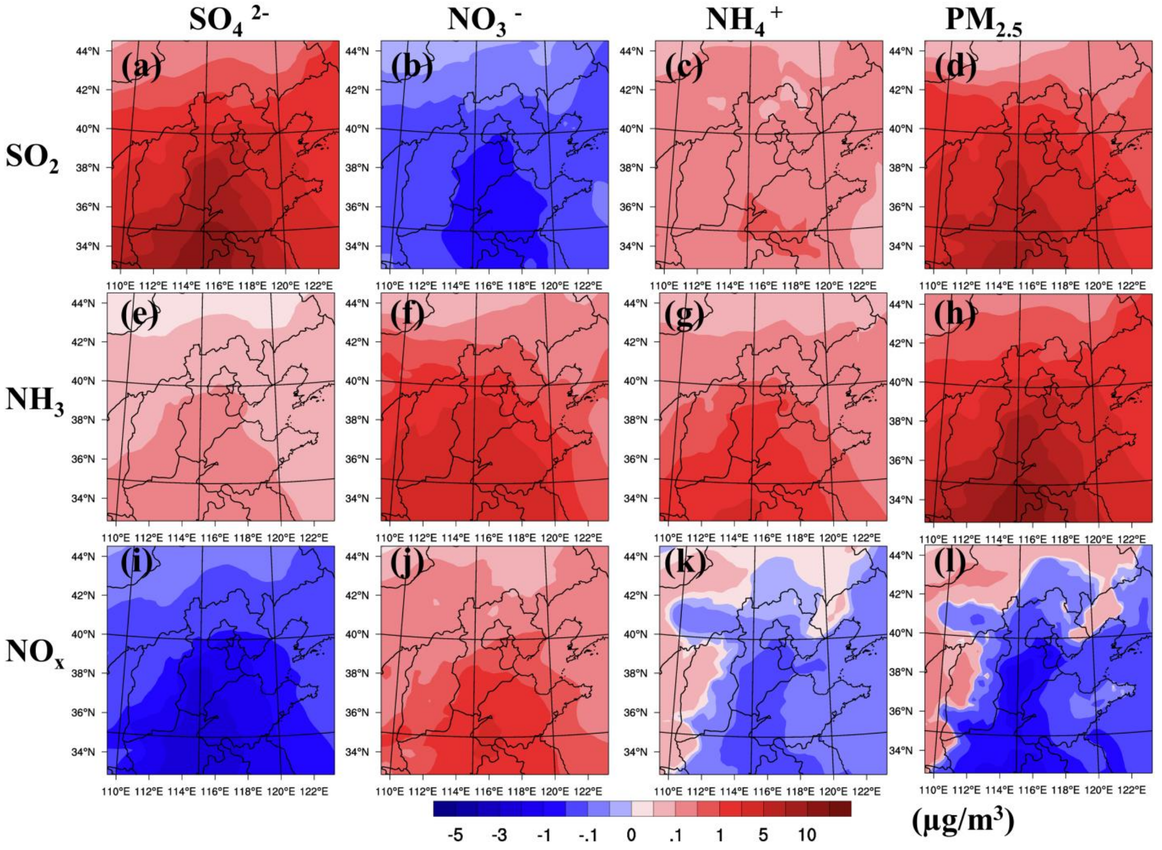

Figure 4. Responses of $PM_{2.5}$ and major $PM_{2.5}$ inorganic species (sulfate, nitrate, and ammonium) to individual changes in SO_2, NH_3 and NO_x emissions from 1960 to 2010

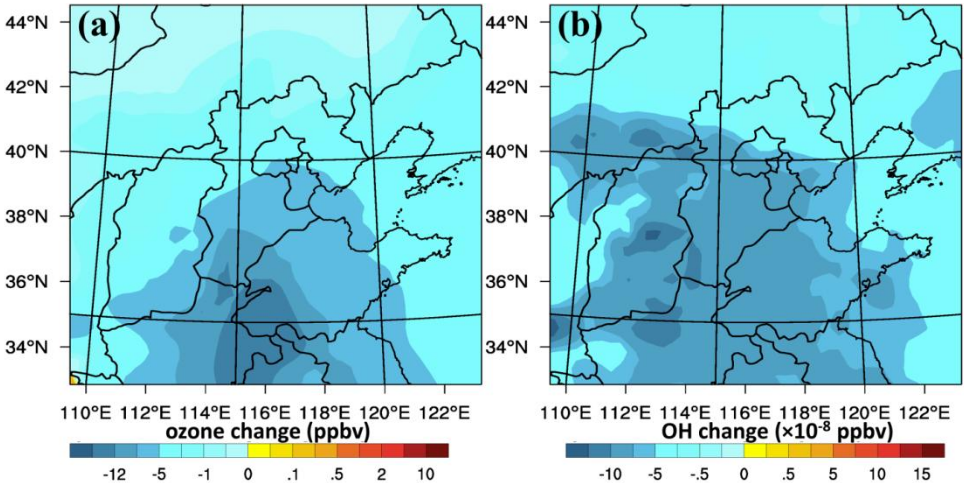

Figure 5. Daytime ozone (a) and daytime OH (b) changes due to NO_x emission increases

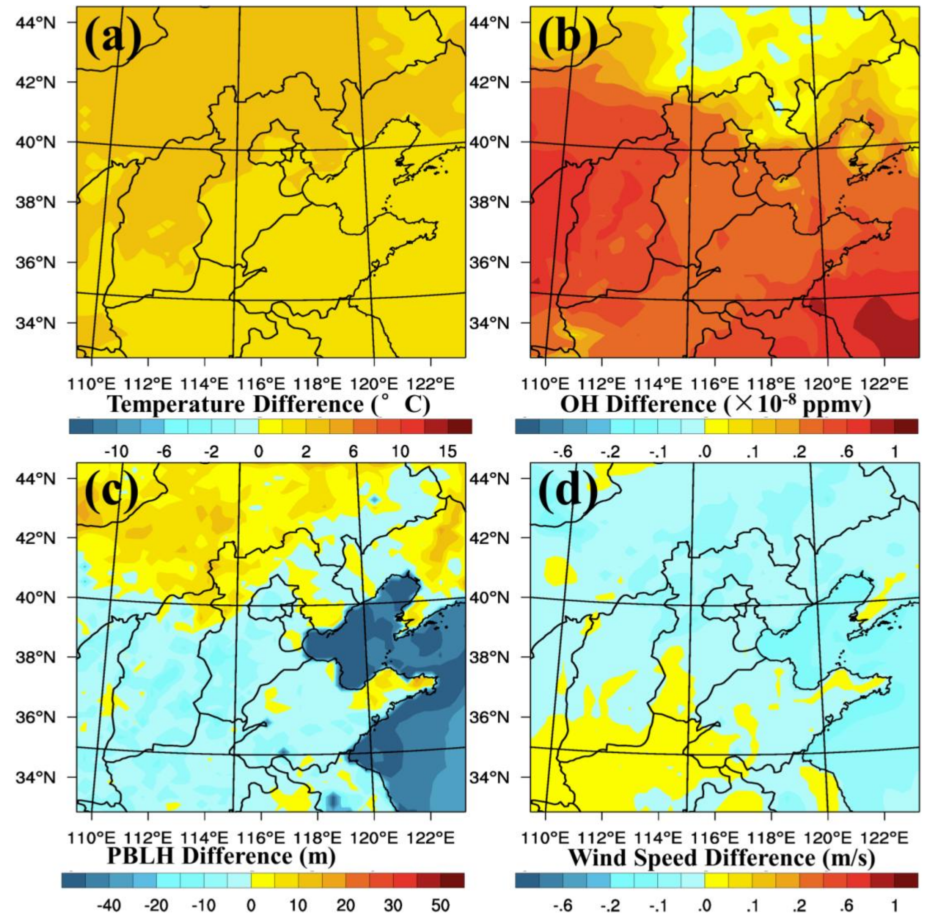

Figure 6. Monthly mean temperature difference due to perturbation in initial and boundary conditions (a), and daily mean OH (b), mean PBLH (c) and mean near surface wind speed changes (d) due to temperature increase

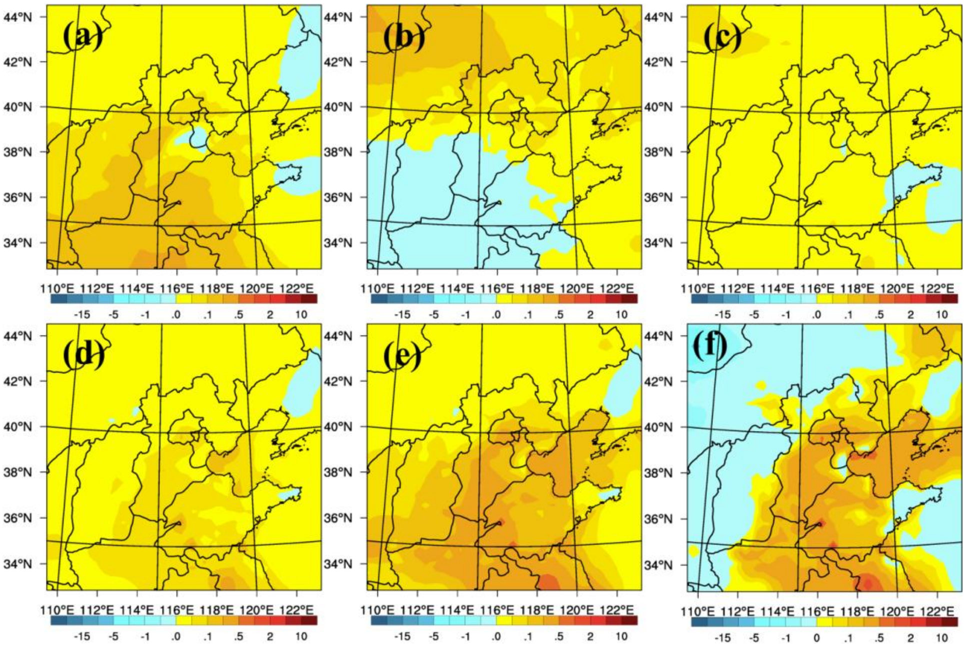

Figure 7. Monthly mean changes of sulfate (a), nitrate (b), ammonium (c), BC (d), OC (e), and $PM_{2.5}$ (f) and due to temperature increase

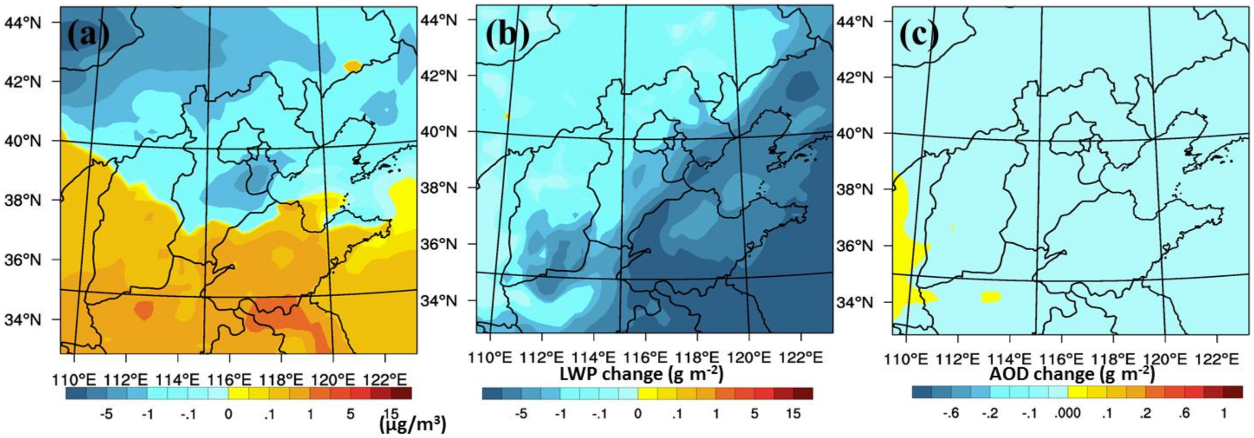

Figure 8. Monthly mean changes of $PM_{2.5}$ (a), LWP (b), and AOD at 600nm (c) due to RH decrease

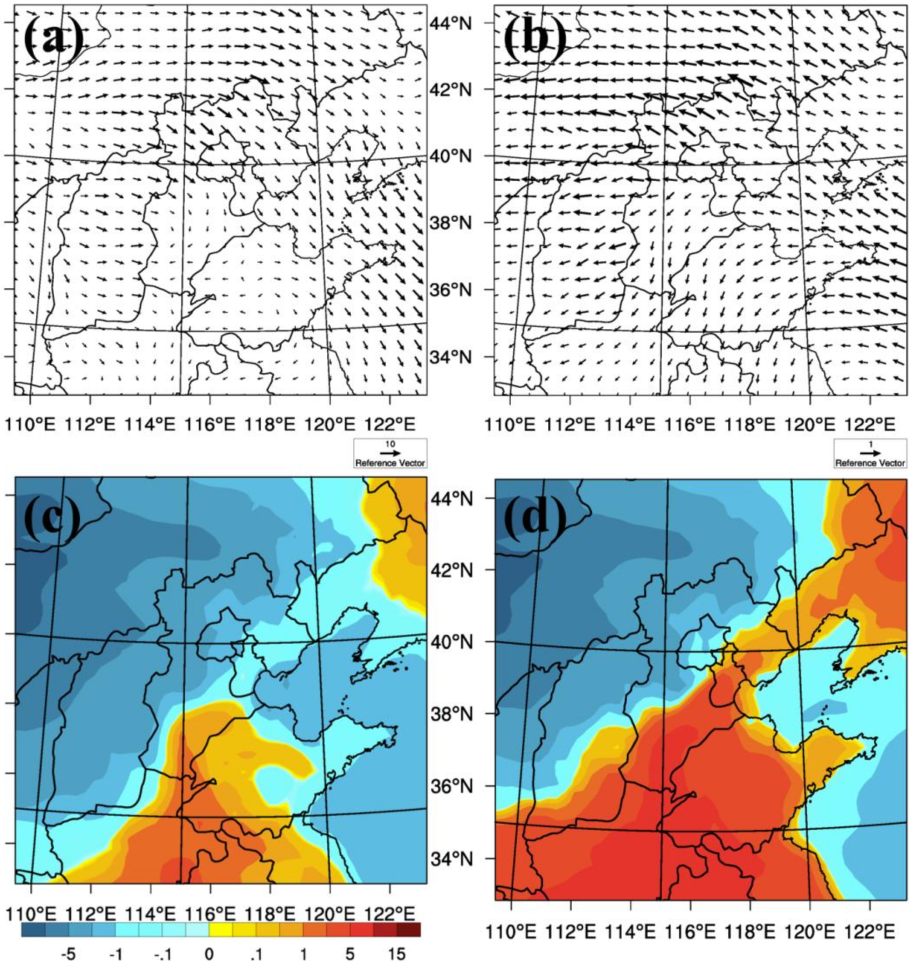

Figure 9. Monthly mean wind fields for WS20 case (a) and changes of wind speeds (CTL-CTL_WS20) (b), and mean changes of $PM_{2.5}$ concentrations based on 1960 emission levels (c) and 2010 emission levels (d)

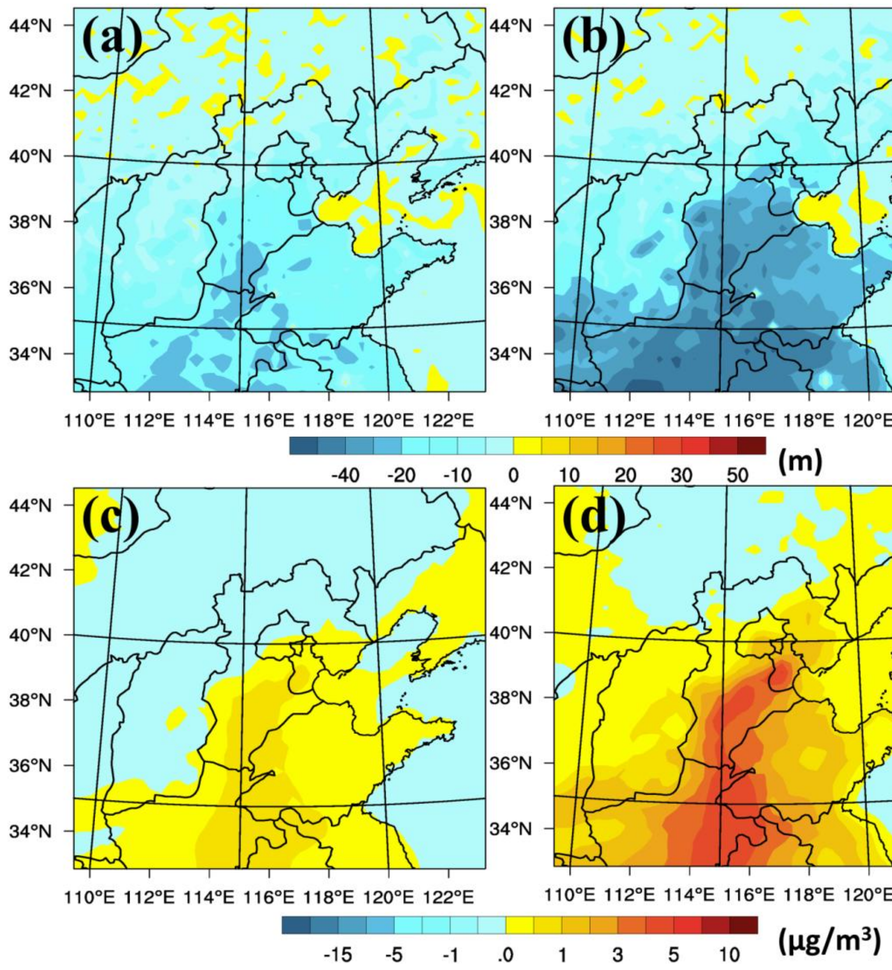

Figure 10. Monthly mean changes of daytime PBL heights for year 1960 (a) and 2010 (b), and of daytime PM$_{2.5}$ concentrations for year 1960 (c) and 2010 (d) due to aerosol-radiation interactions