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Spatial and temporal evolution of future atmospheric reactive nitrogen deposition in China under different climate change mitigation strategies

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Abstract

Atmospheric reactive nitrogen (Nr) deposition plays a crucial role in linking air pollution to ecosystem risks. Previous modeling studies have indicated that climate change and pollution controls jointly result in significant changes in Nr deposition in China. However, it remains unclear how future emission reductions will influence Nr deposition under different climate pathways. Here, we investigated the spatiotemporal evolution and driving factors of future Nr deposition under various national clean air and climate policies. We applied WRF-CMAQ and assessed the historical (2010s, 2010-2014) pattern and future changes of Nr deposition till the 2060s (2060-2064) in China, by combining two SSP-RCP global climate pathways and three national emission control scenarios. The results show that the implementation of clean air and carbon neutrality policies would greatly reduce oxidized nitrogen (OXN) deposition, mitigate the adverse perturbations of climate change, and reduce the outflow from Eastern China (EC) to the West Pacific. In North China (NC), the weakened atmospheric oxidation capacity (AOC) would elevate the response of OXN deposition to a 20% abatement of emissions (expressed as the ratio of percentage change of deposition to emissions) from 82.6% in the 2010s to nearly 100% in the 2060s. In contrast, the response of RDN deposition to NH_3 emissions would decline, likely attributable to a more NH_3-rich condition. The outcomes of this work broaden scientific understanding on how anthropogenic actions of air quality improvement and carbon emission reduction would reshape the future Nr deposition and support effective policymaking to reduce associated ecological damages.

Keywords: Nr deposition, SSP-RCP, climate change, outflow pollution, emission abatement

1. Introduction

With vigorous development of industrial and agricultural activities worldwide

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since the industrial revolution, the emissions of reactive nitrogen (Nr, including oxidized and reduced nitrogen species, OXN and RDN, respectively) have increased explosively (Kanakidou et al., 2016), elevating the Nr levels in both atmosphere and deposition. Enriched ambient Nr has led to a series of regional haze and ozone (O_3) pollution issues through participation in atmospheric aerosol formation and photochemical reactions (Chen et al., 2021). Furthermore, excessive atmospheric Nr deposits onto land and water bodies through both dry and wet forms, directly hurting the stability and productivity of the entire ecosystem (Flower et al., 2013). Substantial Nr deposition can result in diverse adverse ecological effects, such as water eutrophication (Zheng et al., 2020), soil acidification (Raza et al., 2020), and biodiversity loss (Liu et al., 2017).

China has undergone rapid industrialization and urbanization, accompanied with explosive growth in the consumption of fossil fuels and fertilizers over the past few decades, triggering significant emissions of NO_x and NH_3 (Zhao et al. 2013; Kang et al. 2016). Enhanced Nr emissions made the country one of the hotpots receiving largest Nr deposition worldwide (Liu et al., 2013; Vet et al., 2014). Observations of background sites from the China Nationwide Nitrogen Deposition Monitoring Network (NNDMN) during 2011-2018 revealed that the annual averaged Nr deposition fluxes reached 23.6 kg N ha^{-1} yr^{-1}, vastly surpassing the monitoring results in the United States (8.1 kg N ha^{-1} yr^{-1}), Europe (8.7 kg N ha^{-1} yr^{-1}) and Japan (11.0 kg N ha^{-1} yr^{-1}) (Wen et al., 2020). Employing atmospheric chemistry transport models (CTMs) or advanced statistical models, a series of studies have indicated that Nr deposition fluxes has increased nearly 60% since 1980s, notably in eastern China (Gao et al., 2023, Yu et al. 2019, Zhao et al., 2022, Zhou et al., 2023). Evidently, China is still struggling with serious Nr pollution. The national air pollution control actions over the past decade have resulted in a fast decline in emissions of acidic gaseous pollutants (mainly NO_x and SO_2) but relatively stable NH_3 (Zheng et al., 2018). The imbalance in emission reductions for different species has altered the composition of Nr deposition, e.g., a growth in the proportion of RDN (Liu et al., 2020). More importantly, the increasingly strong capacity of

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atmospheric oxidation, attributed primarily to the persistently high emissions of volatile organic compounds (VOCs), has been weakening the response of OXN deposition to NO_x emissions in eastern part of China and thus preventing effective reduction of Nr deposition. One-unit abatement of NO_x emissions resulted in only less than 80% of OXN deposition, emphasizing the crucial role of active O_3-VOCs-NO_x photochemistry in modulating the Nr deposition (Liu et al., 2022).

Atmospheric Nr deposition are mainly influenced by rainfall, precursor emissions, and long-distance transport (Ellis et al., 2013, Kim et al., 2012, Ma et al., 2023, Zhu et al., 2022). The strengthening climate change and implementation of pollution controls will greatly alter the regional meteorological conditions and air pollutant emissions, resulting in substantial changes in magnitude and spatiotemporal pattern of Nr deposition. The changing deposition will further exert multiple impacts on the biodiversity, carbon sequestration and greenhouse emissions of various ecosystems, and thus influence the climate and ecological environment profoundly (Zhu et al., 2020). There are only a few studies addressing future Nr deposition in China. They employed coupled climate-chemistry global models to conduct simulations under different predefined greenhouse gas (GHG) emission scenarios. Future emissions were primarily referencing the Intergovernmental Panel on Climate Change (IPCC) Representative Concentration Pathways (RCPs) based on the radiative forcing in 2100 (van Vuuren et al., 2011). A pioneering study by Galloway et al. (2004) predicted significant growth in Nr deposition in East Asia, exceeding 50 kg N ha^{-1} $year^{-1}$ based on the IPCC92a emission scenario. The Atmospheric Chemistry and Climate Model Intercomparison Project (ACCMIP) presented a multi-model global datasets of Nr deposition, covering the period from 1850 to 2100 (Lamarque et al. 2013a). The Nr deposition in East Asia was estimated to increase 27% and 39% in the 2030s under the RCP2.6 and RCP8.5 pathways, respectively (Lamarque et al. 2013b). More recently, Zhang et al. (2019) and Sun et al. (2022) reported the possible future changes in OXN and RDN deposition, respectively, based on ACCMIP datasets. The OXN deposition fluxes under both RCP4.5 and RCP8.5 pathways were projected to increase in 2030s but decrease by the

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end of the century, driven primarily by the Nr emission trends. The proportion of RDN in total deposition in eastern China was projected to rise from 38% in 2000 to 56% in 2100 under RCP8.5 pathway, suggesting a transition in the dominant form from oxidized to reduced.

Even previous studies made insightful predictions on the future evolution of Nr deposition in China, they have insufficiently incorporated the potentially profound emission reduction in the context of global climate change. In 2020, China announced the plan to achieve carbon neutrality by 2060, and the effects of a wide range of sharp emission reductions on future environment has become a major research focus (Dong et al., 2021). Researchers have integrated national strategies of emission reduction to assess future air pollution and associated health risks in China under various climate change pathways (Cheng et al., 2021a, Cheng et al., 2023, Shi et al., 2021). For example, the IPCC Sixth Assessment Report (AR6) introduced a scientifically combined set of pathways known as Shared Socioeconomic Pathways (SSPs) and RCPs, denoted as SSP-RCP (IPCC, 2021). New pathways integrate the impact of socioeconomic development into the framework for the evolution of GHG levels, offering more reliable projections of possible outcomes of climate change (Cook et al., 2020; O'Neill et al., 2016, Xin et al., 2020). However, there is a noticeable gap in assessment of China's atmospheric deposition under the SSP-RCP framework. The roles of future emission and climate changes on deposition remain unclear across diverse climate pathways. Moreover, stringent emission controls with diverse progresses for various species and regions will change the atmospheric oxidizing capacity and regional transport of pollution, and thereby alter the response of Nr deposition to emissions of their precursors. Given the crucial role of atmospheric deposition in connecting air pollution to ecosystem risks, it is essential to evaluate these anticipated changes for a comprehensive understanding of the ecological and environmental impacts during the long-term progress of continuous air quality improvement and global warming prevention.

In this study, we applied an air quality model (WRF-CMAQ, see details in methods)

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and assessed the future changes of Nr deposition in China, by combining the SSP-RCP global climate change pathways and the national emission control scenarios. The historical period was chosen as 2010-2014, representing the years with the highest Nr emissions in China, and the future simulation period was determined as 2060-2064. Firstly, we evaluated the model performance of meteorology and Nr deposition for the historical period based on available ground observations. We then quantified the spatial and temporal changes of future Nr deposition and identified the main driving factors under two IPCC pathways, SSP2-4.5 and SSP5-8.5). The SSP5-8.5 pathway represents high GHG emissions characterized by continued reliance on fossil fuels, often viewed as a pessimistic outlook for future climate change. Conversely, the SSP2-4.5 pathway envisions moderate GHG emissions, achieved through the consideration of environmental policies and technological advancements. We further assessed the effects of various emissions abatement scenarios on Nr deposition. Finally, we analyzed the future response of deposition to emission perturbation under different scenarios. The study enhances scientific understanding on the interactions between anthropogenic activities and atmospheric chemistry along with a changing climate, and in turn supports the development of effective environmental policies to alleviate the adverse effects of Nr pollution on ecosystems and human health.

2. Methodology and data

2.1 Model description and driving data

The Community Multiscale Air Quality (CMAQ) model version 5.2 (available at https://epa.gov/cmaq/access-cmaq-source-code) was adopted to conduct atmospheric Nr deposition simulations over mainland China for both historical (2010-2014) and future periods (2060-2064). To avoid the model errors associated with individual years, full-year simulations were conducted for every year of the two five-year intervals, and the five-year averages were used for further analyses. A series of simulation cases were designed by combining individual climate pathways and national emission scenarios to

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separate the roles of multiple factors on future deposition (see details in Section 2.2). Developed by the United States Environmental Protection Agency (USEPA), CMAQ has been demonstrated to possess extensive practicality and sophistication in simulating regional air quality and acid deposition (Appel et al., 2017, Chang et al., 2020, Cheng et al., 2021, Liu et al., 2010). A single domain covering mainland China (186 × 156 grid cells) was adopted for the simulations with a horizontal resolution at 27 × 27 km per grid (Figure S1). Lambert conformal conic projection was applied for the domain centered at (102°E, 37°N) with two true latitudes, 40°N and 25°N. In the vertical direction, 30 eta levels with the pressure of 50hPa at the top level were used. For chemical configuration, the carbon bond 05 (CB05) gas-phase chemical scheme and the AERO 6 aerosol scheme were adopted (Sarwar et al., 2008, Pye et al., 2017, Murphy et al., 2017). The boundary condition of trace gases used in this study was background concentration (default setup in CMAQ model). Simulation of each year included a one-month spin-up time (i.e., 1st-31st December of the previous year) to reduce the impact of the initial conditions on the simulations.

The Multi-resolution Emission Inventory for China version 1.3 developed by Tsinghua University (MEICv1.3, available at http://www.meicmodel.org/?page_id=560; Li et al., 2017; Zheng et al., 2018) provided historical anthropogenic emission data within China in our simulations. Information on future emissions were obtained from the Dynamic Projection model for Emissions in China version 1.1 developed by Tsinghua University (DPECv1.1, available at http://meicmodel.org.cn/?page_id=1917). DPEC links global climate mitigation pathways to local clean air policies and fully incorporates China's strict air pollution control progress since the implementation of the "Action Plan of Air Pollution Prevention and Control" in 2013. It thus corrects the erroneous emission trends of China in the Coupled Model Intercomparison Project (CMIP) scenarios (Cheng et al., 2021b; Tong et al., 2020). Three emission scenarios, named as "Baseline", "Current-goal", and "Neutral-goal", were used in this work (see the simulation case design in Section 2.2). The "Baseline" depicts a high-emission scenario in the absence of climate and pollution

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control policies, equivalent to the SSP5-8.5 climate pathway. The "Current-goal" scenario is a combination that takes into account SSP2-4.5 climate pathway along with existing pollution control policies in China. The "Neutral-goal" scenario integrates China's 2060 carbon neutrality goal with the most stringent pollution control policies. Details of the scenarios were described in Cheng et al. (2021b).

Anthropogenic emissions outside of China were taken from the Asian anthropogenic emission inventory, named MIX, developed by the Model Inter-Comparison Study for Asia (MICS-Asia) project (available at http://meicmodel.org.cn/?page_id=1770; Li et al., 2017). Biogenic emissions were calculated by the Model Emissions of Gases and Aerosols from Nature developed under the Monitoring Atmospheric Composition and Climate project version 2.1 (MEGANv2.1; Guenther et al., 2012). The initial horizontal resolutions of both emission inventories were 0.25° × 0.25°, and they were interpolated into our simulation domain with the resolution of 27 km.

The Weather Research and Forecasting (WRF) model version 3.9.1 (available at https://www2.mmm.ucar.edu/wrf/users/wrf_files/wrfv3.9/updates-3.9.1.html) was applied to provide meteorological fields for CMAQ. Developed and maintained collaboratively by the National Center for Atmospheric Research (NCAR) and the National Oceanic and Atmospheric Administration (NOAA), WRF model has been recognized as a state-of-the-art regional weather model and widely utilized in short-term weather forecasting and regional meteorological research (Huang et al., 2020, Skamarock et al., 2008, Wang et al., 2021). For our historical meteorological simulation, the fifth generation of European Centre for Medium-Range Weather Forecasts (ECMWF) reanalysis dataset, ERA5 (available at https://cds.climate.copernicus.eu/cdsapp#!/dataset/reanalysis-era5-single-levels?tab=form) was adopted as the initial and boundary field (Hersbach et al., 2020). The temporal and spatial resolution was 6 hours and 0.25° × 0.25°, respectively. For simulation of future period, it is commonly practical to employ climate forecast results from global climate models (GCMs) as the initial and boundary conditions. In this study,

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a global bias-corrected multi-model (BCMM) climatological dataset with a horizontal resolution of 1.25°×1.25° at 6-hour intervals (available at https://www.scidb.cn/en/detail?dataSetId=791587189614968832#p2) was adopted to drive WRF model for 2060-2064. The BCMM dataset was reconstructed from 18 GCMs of the CMIP6, with corrections for climatological mean and interannual variance biases based on ERA5 data from 1979-2014, providing more reliable projections of long-term non-linear trends of multiple climate variables compared with original CMIP6 model outputs. Details of BCMM product were described at Xu et al. (2021). We employed Pseudo Global Warming (PGW) method (Kawase et al., 2013, Liu et al., 2021, Lauer et al., 2013, Taniguchi et al., 2020) for statistical downscaling. Specifically, future driving fields were forced with the ERA5 data from reference period (2010-2014) plus a climate perturbation (difference between the years 2060-2064 and 2010-2014) calculated from BCMM results, as shown in Eq. (1) and Eq (2):

$$WRF_{input2060-2064} = ERA5_{2010-2014} + \Delta BCMM_{ssp} \quad (1)$$

$$\Delta BCMM_{ssp} = BCMM_{\overline{2060-2064}} - BCMM_{\overline{2010-2014}} \quad (2)$$

where $\Delta BCMM_{ssp}$ is the CMIP6 multimodel ensemble mean change signal for 2060-2064 relative to 2010-2014 under the SSP2-4.5 or SSP5-8.5 pathway, $BCMM_{\overline{2060-2064}}$ and $BCMM_{\overline{2010-2014}}$ represent the 5-year meteorological averages of BCMM dataset in the future and reference periods, respectively. Nine physical variables were perturbed in this study including zonal wind, meridional wind, air temperature, sea surface temperature, soil temperature, specific humidity, the surface pressure, sea-level pressure and geopotential height. The bilinear interpolation was applied to interpolate BCMM data to the ERA5 grid.

The land-use and land-cover (LULC) data were taken from global data of the U.S. Geological Survey (USGS) (de Meij et al., 2014; Pineda et al., 2004). The physical parameterization schemes used in all simulations are summarized in Table S1 in the Supplement.

The dry deposition (DDEP) of each atmospheric chemical species (i) was calculated as the product of surface concentration ($C^{surface}$) and dry deposition velocity

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(V_d) at the lowest model layer, as shown in Eq. (3):

$$DDEP_i = C_i^{surface} \times V_d \quad (3)$$

According to the classical resistance cascade model (Venkatram and Pleim, 1999; Wesely, 2007), the parameters of V_d are calculated as Eq. (4):

$$V_d = 1/(R_a + R_b + R_c) \quad (4)$$

where R_a is the aerodynamic resistance to the transfer from lowest layer to the roughness height, calculated as a function of surface layer turbulence parameters including friction velocity and the Monin-Obukhov length; R_b is the boundary layer resistance to transfer between the roughness height and surface; Rc is the resistance to surface uptake, which can be further divided into several series and parallel components, representing the resistance to the lower vegetation canopy or ground.

The algorithm module for wet deposition (WDEP) is derived from the regional acid deposition model (RADM; Chang et al., 1987) and depends on the precipitation rate (P_r) and cloud water concentration (C_{cloud}) of specific chemical component:

$$WDEP_i = P_r \cdot \bar{C}_i^{cloud} \quad (5)$$

The wet scavenging is considered in two pathways, depending upon whether the pollutant participates in the cloud water chemistry and on the liquid water content. Details on how CMAQ removes pollutants through wet deposition can be found in the official CMAQ Science Documentation (available at https://www.cmascenter.org/cmaq/science_documentation/pdf/ch11.pdf). In this study, OXN included NO, NO_2, HNO_3, N_2O_5, HONO, and particulates as nitrate (NO_3^-), and RDN included NH_3 and particulates as ammonium (NH_4^+).

2.2 Numerical simulation experiment design

To evaluate future changes in the spatiotemporal pattern of atmospheric Nr deposition under different SSP-RCP climate pathways and emission control scenarios, we performed parallel numerical simulation experiments with WRF-CMAQ, as summarized in Table 1. Base simulated the real situation in historical period (2010-2014). Case 1 and Case 2 were designed to predict the atmospheric Nr deposition in the

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2060s, following SSP2-4.5 climate pathway with "Current-goal" emission scenario in DPEC and SSP5-8.5 climate pathway with "Baseline" emission scenario, respectively. Difference between Case 1and Base and that between Case 2 and Base respectively revealed the changing Nr deposition from 2010s to 2060s in SSP2-4.5 and SSP5-8.5.

Cases 3 and 4 applied future climate pathways (SSP2-4.5 and SSP5-8.5, respectively) but historical emissions, and the difference between each of them and Base revealed how climate change would influence Nr deposition under corresponding climate pathway. Meanwhile, the effect of emission change on future Nr deposition was examined by comparing Case 3 and Case 1 for "Current-goal" scenario in DPEC, and by comparing Case 4 and Case 2 for "Baseline" scenario. Case 5 applied SSP2-4.5 climate pathway and "Neutral-goal" emission scenario in DPEC. Comparison between Case 5 and Case 3 revealed the benefit of national emission controls under China's carbon neutrality policy on Nr deposition.

Cases 6-8 were designed based on Cases 1, 3, and 5, respectively. In these cases, emissions in eastern China (EC) were set at the 2060s level, while those in western China (WC) were maintained at the 2010s level. The aim was to explore the effect of diverse emission control progresses for different regions on the future Nr deposition. WC and EC were divided by longitude 110° east in this study, as shown in Figure S1. In Cases 9-12, the emissions of all species were reduced by 20% from those in Cases 3, 1, 2, and 5, respectively, to explore the response of deposition to emission perturbation at different atmospheric conditions caused by varying pollution control levels.

2.3 Observations and model evaluation

WRF-CMAQ model performance was evaluated against available observation of meteorological variables and Nr deposition at monthly or annual level. Daily near-surface observations of four meteorological parameters including temperature at the height of 2 m (T2), relative humidity (RH), wind speed at the height of 10 m (WS10) and accumulated precipitation (PREC) were derived from the National Meteorological Data Center of China Meteorological Administration (CMA,

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http://data.cma.cn/data/detail/dataCode/A.0012.0001.html). The 839 meteorological surface stations, with continuous five-year observations from 2010 to 2014 were selected, as shown in Figure 1. Meanwhile, the monthly observations of Nr deposition fluxes were taken from the Nationwide Nitrogen Deposition Monitoring Network (NNDMN; Xu et al., 2018; 2019). Following our previous study, we selected 28 sites for dry deposition fluxes and 53 sites for wet deposition fluxes, for which at least two-year continuous measurement data were available, to evaluate model performance. Details of monitoring stations can be found in Ma et al. (2023). Statistical indicators were calculated with Eq. (6-9), including normalized mean bias (NMB), normalized mean error (NME) and the correlation coefficient (R) at temporal and spatial scales (Baker et al., 2004; Ma et al., 2023):

$$NMB = \sum_{i=1}^{n}(S_i - O_i)/\sum_{i=1}^{n} O_i \times 100\% \quad (6)$$

$$NME = \sum_{i=1}^{n}|S_i - O_i|/\sum_{i=1}^{n} O_i \times 100\% \quad (7)$$

$$R(temporal) = \sum_{i=1}^{n}(S_i - \bar{S})\,(O_i - \bar{O})/\sqrt{\sum_{i=1}^{n}(S_i - \bar{S})^2(O_i - \bar{O})^2} \quad (8)$$

$$R(spatial) = \sum_{j=1}^{m}(\bar{S_j} - \bar{S_j})\,(\bar{O_j} - \bar{O_j})/\sqrt{\sum_{j=1}^{m}(S_j - \bar{S_j})^2(O_j - \bar{O_j})^2} \quad (9)$$

where S and O are the monthly meteorological variables or annual Nr deposition from model simulation and observation, respectively; $\bar{S}$ and $\bar{O}$ are the monthly mean meteorological variables or annual deposition from model simulation and observation, respectively; i means the individual month or year and j means the individual site.

3. Results and discussion

3.1 Evaluation of model performance

We compared the simulated near-surface temperature, wind speed, relative humidity and accumulated precipitation with observations at the monthly level, as shown in Figure 1. The model reasonably reproduced the spatial pattern of near-surface temperature with the spatial R reaching 0.95 (Figure 1a). Overestimation was found in the southeast and northwest of the country while underestimation over the Tibetan

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Plateau. At the national scale, T2 was generally underestimated with the NMB and NME calculated at -7.76% and 12.75%, respectively. In addition, the temporal R reached 0.99, indicating the simulation was in good agreement with observation at the monthly level. Unlike T2, due to the modeling biases in the topographic effects and the underestimation of urban land use in USGS (Carvalho et al., 2012; Liao et al., 2015), WS10 was overestimated with NMB calculated at 33.13% at the national scale (Figure 1b). Such overestimation was also reported in other studies (Liu et al., 2020, Shen et al., 2021, Zhu et al., 2022). RH is slightly underestimated with NMB and NME calculated at -1.58% and 8.94%, respectively, while both spatial and temporal R were greater than 0.8 (Figure 1c). PREC was generally underestimated, with NMB and NME at -19.39% and 39.15%, respectively. A clear gradient from northwest to southeast China was well captured, and the temporal and spatial R were 0.83 and 0.76, respectively (Figure 1d).

The comparison between the simulated and observed annual Nr deposition averaged over 2010-2014 at the site level are provided by form (dry and wet) and species (OXN and RDN) in Table 2. Nr deposition was underestimated for all cases. The NMB and NME for the dry deposition of OXN (DDEP_OXN) were calculated at -9.07% and 24.76%, respectively, and the analogous numbers for RDN (DDEP_RDN) were at -15.12% and 43.24%. The uncertainty in NH_3 emission inventories was frequently recognized as an important factor contributing to the underestimation (Ma et al., 2023, Chang et al., 2020, Shen et al., 2023). The limited development of intensive livestock breeding and farming in China poses a considerable challenge in acquiring sufficient activity data and accurate emission factors, leading to underestimation of emissions with the “bottom-up” approach. Utilizing satellite constraints, Zhang et al. (2018) estimated that the total NH_3 emissions in China may be underestimated by nearly 40%. Due to lack of direct observation, additionally, the dry deposition at NNDMN sites was calculated by multiplying the observed surface concentrations with V_d simulated from GEOS-Chem (Bey et al., 2001; Xu et al., 2019). Difference in the parameterization schemes for calculating V_d of given trace gases or aerosols between

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CTMs could also introduce modest uncertainty for assessment of OXN deposition (Wu et al., 2018; Chang et al., 2020). Compared to dry deposition, wet deposition of OXN and RDN (WDEP_OXN and WDEP_RDN) was simulated to be far lower than the observations, with the NMBs calculated at -28.76% and -17.86%, respectively. Part of the reason may be underestimation of precipitation (Figure 1d). More importantly, most of wet deposition measured at NNDMN sites was actually "bulk deposition", which included both wet deposition and a small fraction of dry deposition (Xu et al., 2015). Therefore, the bias from observation also contributed to the inconsistency.

Project of the Model Inter-Comparison Study for Asia (MICS-Asia) phase III reported the performances of Nr deposition simulation with multiple models over China, with the overall NMBs and NMEs ranged -47% – 67% and 48% – 82% for OXN, and -70% – -29% and 44% – 72% for RDN, respectively (Ge et al., 2020). The model performance in our study was comparable to previous studies. In addition, both spatial and temporal R were greater than 0.6 for each deposition form and species. Overall, our simulations reasonably reproduced the observed Nr deposition in both magnitude and spatiotemporal patterns.

3.2 Evolution of Nr deposition and the roles of climate and emission changes

Table 3 summarizes the simulated atmospheric Nr deposition over historical (Base) and future periods under SSP2-4.5 (Case 1) and SSP5-8.5 pathways (Case 2). The annual averaged Nr deposition for 2010-2014 was simulated at 14.7 kg N ha^{-1} yr^{-1} for mainland China (Base). The contribution of RDN to total deposition reached 52%, which was in good agreement with the multiple-model ensemble mean value in the MICS-Asia phase III project (Ge et al., 2020). The ratio of wet deposition to total deposition was 0.54 in our simulation, also close to other CTM and nationwide observation results (Ge et al., 2020, Xu et al., 2015, Zhao et al., 2017).

Under the SSP2-4.5 pathway, total Nr deposition would decrease to 9.0 kg N ha^{-1} yr^{-1} during 2060-2064, primarily attributed to a sharp decline in OXN deposition (Case

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1). Accompanied with an active energy transition and effective control of fossil fuel consumption, the substantial reduction of anthropogenic NO_x emissions led to a 56% decline in OXN deposition compared to the reference period. Meanwhile, RDN deposition would be reduced by only 22%, resulting from a modest abatement of NH_3 emissions. Figure S2 in the supplement shows the changes of NO_x and NH_3 emissions in 2060 relative to the historical period (2010-2014) in various scenarios, and Figure S3 in the supplement provides the annual emissions by sector. Large emission changes would occur mainly in the east of China. By 2060s, the national NO_x emissions would decline 55% (-15.1 Mt) under the "Current-goal" emission scenario. Such reductions would come mainly from power, industry and transportation sectors, driven by the predicted transition of energy structure (Figures S3a-b). Due to less improvement in agriculture management, the NH_3 emissions would decline much slower by 28% (-2.9 Mt). Under the SSP5-8.5 pathway, the global economy would maintain rapid growth without sufficient considerations for climate change. A high dependence on fossil fuels (especially coal) for energy consumption would result in a nationwide growth of annual NO_x emissions by 24% (6.5 Mt) from 2010s to 2060s ("Baseline" scenario in DPEC, Figure S2a), and thereby elevate the total Nr deposition to 15.4 kg N ha^{-1} yr^{-1} (Case 2). The proportions of OXN and RDN in future Nr deposition were anticipated to vary across different SSP-RCP pathways. Under the SSP2-4.5 pathway, RDN was predicted to be the dominant species of Nr deposition in the 2060s, with a proportion to the total estimated at 66%. Under the SSP5-8.5 pathway, the proportion of OXN to total deposition was expected to expand from 48% in the 2010s to 55% in the 2060s.

In terms of spatial pattern, our simulations present clearly larger regional difference in China compared to the global results of ACCMIP, owing to finer simulation resolution and more detailed regional emission information. Figure 2 illustrates the spatial distribution of Nr deposition in historical period and the future changes under different SSP-RCP pathways. For 2010-2014, a clear gradient from west to east was found for all deposition forms and species (Figures 2a-d), driven mainly by the spatial distributions of NH_3 and NO_x emissions. Dry deposition of OXN

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(DDEP_OXN) appeared mainly in eastern China, especially in the Beijing-Tianjin-Hebei (BTH), Yangtze River Delta (YRD) and Pearl River Delta (PRD) regions (see Figure S1 for the locations of these regions), resulting mainly from the large NO_x emissions caused by active industrialization and urbanization. Hotspots of RDN dry deposition (DDEP_RDN) appeared mainly in the North China Plain and the Sichuan Basin (SCB) with intensive agricultural activities. Further influenced by precipitation patterns, the southern areas experienced greater wet deposition compared to the north, consistent with previous studies (Han et al., 2017; Zhao et al., 2017). Influenced jointly by the substantial rainfall and local Nr emissions, in particular, SCB was of the largest wet deposition for both OXN and RDN (WDEP_OXN and WDEP_RDN).

The future OXN deposition would exhibit contrasting trends between the SSP2-4.5 and SSP5-8.5 pathways. Compared to historical periods, both dry and wet forms were predicted to decrease in the 2060s under the SSP2-4.5 pathway, with national average reductions of 2.2 kg N ha^{-1} yr^{-1} and 1.8 kg N ha^{-1} yr^{-1}, respectively. Relative large declines would be found in their respective hotspots (Case 1-Base, Figures 2e-f). In contrast, a growth of OXN deposition would appear under the SSP5-8.5 pathway, contributed mainly by wet deposition. The changes of dry deposition would be limited within 1 kg N ha^{-1} yr^{-1} at the national level (Case 2-Base, Figures 2i-2j). For RDN deposition, there would be a nationwide decline in the 2060s under SSP2-4.5 pathway (Case 1-Base, Figures 2g-h). Large decline would be found for wet deposition in the SCB and the surrounding area, with the maximum exceeding 10 kg N ha^{-1} yr^{-1}. The changes under the SSP5-8.5 pathway would be small, with the national average reduced by 0.1 and 0.5 kg N ha^{-1} yr^{-1} for dry and wet deposition, respectively (Case 2-Base, Figures 2k-l).

With Cases 3 and 4 included in the analyses, we further estimated the impacts of climate and emission change on future total Nr deposition and compared them with the joint impact (Figure 3). Under the SSP2-4.5 pathway, the national average difference in Nr deposition due to changing emissions alone (-5.48 kg N ha^{-1} yr^{-1}, Figure 3b) was closer to that from joint impacts (-5.77 kg N ha^{-1} yr^{-1}, Figure 3c), while the difference

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caused by climate change alone was small (-0.29 kg N ha^{-1} yr^{-1}, Figure 3a). Additionally, the spatial correlation (R) between the difference in deposition due to emission change alone and that due to both factors would be 0.89 (Figure 3b), while it would be clearly smaller at 0.66 between those due to climate change alone and both factors (Figure 3a). This indicates that the future long-term Nr deposition would be primarily dominated by emission change. Under the SSP5-8.5 pathway, the total amount of Nr deposition change at the national level would also be dominated by the varying emissions. The emission change alone would lead to a growth of nationwide deposition at 0.83 kg N ha^{-1} yr^{-1} (Figure 3e), 90% of the total growth (0.92 kg N ha^{-1} yr^{-1}, Figure 3f). However, the spatial pattern of deposition would be largely modulated by climate change, with the spatial R between the deposition differences due to climate change alone and both factors reaching 0.84 (Figure 3d). The value would only be 0.55 between differences due to emission change alone and both factors (Figure 3e). In the southern BTH, for example, future climate change would elevate the deposition by over 4 kg N ha^{-1} yr^{-1}. By comparing the roles of emission and climate changes in Nr deposition under different SSP-RCP pathways, our study emphasizes that the rigorous implementation of emission controls in the future can effectively mitigate the adverse perturbations of climate change.

3.3 Varying effects of different emission changing patterns on Nr deposition

We further quantified the effects of emission controls on the deposition of different Nr components (OXN and RDN) and compared them under various future emission scenarios ("Baseline", "Current-goal", and "Neutral-goal"). As illustrated in Figure 4, with an exception of OXN deposition in "Baseline" scenario which would increase 24% (1.42 kg N ha^{-1} yr^{-1}) from 2010s to 2060s, the national Nr deposition would commonly decline for other cases, ranging from 5% to 85% (0.27-4.93 kg N ha^{-1} yr^{-1}). In the "Neutral-goal" scenario, in particular, the national average OXN deposition was predicted to decline to 0.98 kg N ha^{-1} yr^{-1} by 2060s (Figure S4), accounting for only

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17% of the total Nr deposition. This implies that the continuous and substantial reduction in NO_x emissions, implemented as part of the national strategy to address climate change and to improve air quality, would make RDN become the dominant contributor to future Nr deposition. Spatial correlation between future emission change and the resulting deposition change was estimated and summarized in Table S2 in the supplement for different emission scenarios. Compared with OXN, the spatial change in RDN deposition would be more consistent with that of precursor emissions, indicated by a much higher R for RDN (0.67-0.72) than OXN (0.24-0.35). The discrepancy could result from the stronger regional transport of NO_x, which comes largely from high-stack sources (Ma et al., 2020).

Figure 5 compares the relative changes in future Nr deposition and precursor emissions for WC and EC in different emission scenarios. Under the "Baseline" scenario (Figure 5a), the OXN deposition in WC was predicted to increase 47% from 2010s to 2060s. This growth would be notably smaller than that of NO_x emissions (81%), suggesting that a larger amount of OXN in WC would deposit to the east through atmospheric transport. However, the transport might be weakened from WC to EC in the "Current-goal" (Figure 5b) and "Neutral-goal" scenarios (Figure 5c), in which the OXN deposition in WC would decline (46% and 85%, respectively) greater than that of NO_x emissions (41% and 77%, respectively). Additional experiments were conducted to quantify the impact of changing transport from WC on deposition in EC, by keeping the emissions in WC at the 2010s levels (Cases 6-8). The fluxes crossing 110°E from west to east were calculated within the altitude from the surface to 100 hpa and latitude from 20°N to 50°N. Compared to the cases where emissions in WC were maintained at the 2010 levels, the outflow fluxes of OXN would change by 17.57 (Case 2-Case 7), -20.10 (Case 1-Case 6) and -37.12 kg N s^{-1} (Case 5-Case 8) for "Baseline", "Current-goal" and "Neutral-goal" scenarios, respectively (Table S3). Consequently, the OXN deposition in EC would change by 0.30 (2%), -0.28 (-5%), and -0.51 kg N ha^{-1} yr^{-1} (-27%) from 2010s to 2060s due to the emission variation in WC for different scenarios (Table 4).

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The OXN deposition in EC was predicted to increase 13%, despite a 17% growth in NO_x emissions under the "Baseline" scenario (Figure 5a). The additional deposition loss may have been exported off-land through long-distance transport processes. Zhao et al. (2017) demonstrated that 30% of China's Nr emissions from 2008-2010 were transported to the China Sea Area of the Northwest Pacific. We calculated the outflow fluxes of OXN from EC crossing 123°E within the altitude from the surface to 100hpa and latitude from 20°N to 50°N (Table S4). Under the "Baseline" scenario, the outflow fluxes from EC in 2060s would increase by 34.56 kg N s^{-1} compared to the case with the emissions maintained at the 2010s level (Case 2-Case 4). In contrast, the outflow fluxes under the "Current-goal" and "Neutral-goal" scenarios would respectively decline by 94.45 (Case 1-Case 3) and 172.86 kg N s^{-1} (Case 5-Case 3) attributable to the emission abatement in EC, making the relative changes in NO_x emissions and OXN deposition would be essentially equal. The result implies that effective implementation of China's clean air and carbon neutrality policies would definitely weaken its role of exporting pollution to west Pacific.

For RDN deposition, the relative change in emissions and deposition would be essentially the same under the "Baseline" scenario (Figure 5a). However, the change in RDN deposition would be smaller than that of NH_3 emissions for both EC and WC in the remaining two scenarios (Figure 5b and 5c). Given its short atmospheric lifetime (generally a few hours) and thereby limited long-distance transport capability (Hertel et al., 2006), the lag in RDN deposition reduction could primarily result from chemical transformation processes. As a crucial reduced gas in the atmosphere, NH_3 exhibits high capability of neutralizing acid gases, thereby slowing down the formation of acid rain and actively participating in the production of sulfates (SO_4^{2-}) and NO_3^-. With the substantial reduction in acidic pollutants, the secondary formation of ammonium sulfate and ammonium nitrate aerosols would decline, leading to an enhanced proportion of gaseous NH_3 in RDN. Given much larger V_d of gaseous NH_3 than that of particulate NH_4^+, the enhanced NH_3 would result in a growth in dry deposition of RDN, thus slowing the decline of total RDN deposition.

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3.4 Responses of future Nr deposition to emission perturbation

Figure 6 shows the predicted response of Nr deposition to a 20% emission reduction for 2010s and 2060s under different emission scenarios. The response was obtained by calculating the ratio of the percent change in deposition to that in emissions. For OXN, the nationwide average response of OXN deposition to NO_x emissions was 83% for the 2010s (Figure 6a). There was a clear north-south difference in the response over EC. We defined Northern China (NC, 30°N-45°N, 110°E-125°E) and Southern China (SC, 20°N-30°N, 110°E-125°E, Figure S1) and calculated the response of OXN deposition to NO_x emission change at 83% and 96%, respectively (Table 5). As a comparison, Liu et al. (2022) reported the response of OXN deposition to NO_x emissions ranging 55-76% in North China Plain and neighboring areas during the 2010s. High ratio of NO_x to VOCs emissions in NC resulted in the NO_x-saturated regime for O_3 formation, and reduced NO_x emissions enhanced the atmospheric oxidation capacity (AOC) and in turn promoted the production of atmospheric nitric acid (HNO_3). Additionally, there was insufficient ambient free NH_3 to completely neutralize the gaseous HNO_3, an important component of OXN_DDEP (Liu et al., 2018; Zhai et al., 2021). The relatively large proportion of HNO_3 in OXN restrained fast decline of OXN_DDEP, given the larger V_d of HNO_3 compared to that of NO_2. Overall, the enhanced AOC, coupled with relatively NH_3-poor condition, resulted in a weak response of OXN deposition to emissions reduction. In our simulations, emissions were controlled for all species including VOCs. Compared to Liu et al. (2022) with NO_x emission reduction only, the extra VOCs emissions reduction might lower AOC due to their great contribution to the formation of O_3 and OH radicals in the atmosphere (McDonald et al., 2018). Thus, the moderately large response in our simulation resulted from the simultaneous reduction of VOC and NO_x emissions, which would partially offset the AOC enhancement induced by NO_x emission control alone, and thereby restrain the OXN deposition to some extent.

Similar to the 2010s, the response of OXN deposition to a 20% emission reduction in the 2060s would be 84% over NC under the “Baseline” scenario, in which VOCs and

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NO_x emissions would remain high levels (Table 5). A 20% reduction in emissions would lead to a 17% decline in near-surface annual mean NO_2 concentrations (Figure S5a in the supplement) but a 3.2% growth in O_3 concentration in NC (Figure S5b). In contrast, under the scenarios of "Current-goal" and "Neutral-goal", a 20% emission reduction would result in 0.82% and 2.7% decline in near-surface O_3 concentration, respectively (Figure S5b), indicating a weakening non-linear mechanism between emission reduction and AOC enhancement with long-term control of air pollution. Meanwhile, the annual mean HNO_3 concentrations would decrease by 14% and 19% (Figure S5c), and OXN_DDEP would decrease by 18% and 19% (Figure S5d) in "Current-goal" and "Neutral-goal" scenarios, respectively. The reductions would be greater than those for the historical period and the future "Baseline" scenario (10% and 11% for HNO_3 concentration and 14% and 14% for OXN_DDEP, respectively). Consequently, the response of total OXN deposition to emission controls would reach 92% and 95%, respectively. Compared to NC, greater effectiveness of emission abatement on decreasing OXN deposition was found in SC for both 2010s and all the future scenarios in 2060s. The response was estimated to range 93%-103.00% (Table 5), similar to the results of 80-120% in the United States (Tan et al., 2020).

The response of RDN deposition to a 20% reduction of emissions was estimated at 96% in 2010s, clearly larger than that in the United States (60-80%, Tan et al., 2020). The value would decline to 94% and 92% for "Current-goal" and "Neutral-goal" scenarios in 2060s, respectively, implying that the national air quality and carbon neutrality policies would enhance the nonlinear response of RDN deposition to precursor emission change, towards current US condition. As mentioned in previously, part of the reason could be the transition to a more NH_3-rich condition in the future, resulting from more stringent emission controls of SO_2 and NO_x than NH_3. The proportion of gaseous NH_3 (with larger V_d than particulate NH_4^+) to total RDN would be enhanced, which would in turn delay the reducing RDN deposition. In addition, our simulations did not account for the bidirectional feedback between atmospheric NH_3 and soil. Soil volatilization could weaken the sensitivity of dry deposition of RDN to

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changing NH_3 emissions.

4 Conclusion remarks

Combining two global SSP-RCP climate change pathways and three Chinese emission control scenarios, we assessed the spatiotemporal evolution of future atmospheric Nr deposition in China, its main driving factors, and the changing response of Nr deposition to precursor emission controls. Under the SSP5-8.5 pathway, the total Nr deposition would increase from 14.7 in 2010s to 15.4 kg N ha^{-1} yr^{-1} in 2060s, and the spatial pattern of deposition would largely be modulated by climate change. In contrast, under the SSP2-4.5 pathway, Nr deposition is predicted to decrease to 9.0 kg N ha^{-1} yr^{-1} by the 2060s, strongly driven by emissions changes. Implementation of clean air and carbon neutrality policies would make RDN become the dominant contributor to future Nr deposition. In the "Neutral-goal" scenario, in particular, the national average OXN deposition was predicted to decrease to 0.98 kg N ha^{-1} yr^{-1} by the 2060s, accounting for only 17% of the total Nr deposition. Previous studies at the global scale have also indicated the increasing role of RDN deposition in the future, but the growth of RDN share was commonly predicted to be slower, due to insufficient knowledge on China's actions on NO_x emission controls. For example, ACCMIP, as reported by Sun et al. (2020), expected that the ratio of RDN to total Nr deposition in eastern China will increase to only 56% by the end of the century.

Through experiments with fixed WC emissions, we further revealed that the OXN deposition from WC to EC in the 2060s would increase by 0.30 kg N ha^{-1} yr^{-1} (2%) compared to the 2010s under the "Baseline", but decline by 0.28 kg N ha^{-1} yr^{-1} (5%) and 0.51 kg N ha^{-1} yr^{-1} (27%) under the "Current-goal" and "Neutral-goal" scenarios, respectively. Similarly, the outflow OXN fluxes from EC in 2060s would decline 94.45 kg N s^{-1} (49%) and 172.86 kg N s^{-1} (89%) in the latter two scenarios in 2060s, respectively. The response of OXN deposition to a 20% abatement of emissions in NC was estimated at 84% under the "Baseline" scenario, while it would approach 100% in the "Current-goal" and "Neutral-goal" scenarios with the declining share of gaseous

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HNO_3 in OXN due to weakened AOC. In contrast, the response of RDN deposition to a 20% abatement of emissions would decline in the latter two scenarios, attributed partly to a more NH_3-rich condition and thereby a growing share of gaseous NH_3 in 2060s.

Our study suggests that future rigorous implementation of clean air and carbon neutrality policies can mitigate the adverse effects of climate change on Nr deposition, and weaken the transport of air pollution to West Pacific. It highlights the potential changes in the source-sink relationship for China, and supports scientific analyses on sources and mitigation of Nr pollution, not only for China but also for downwind areas. More attention needs to be paid to NH_3 emission controls due to its increasing importance on Nr deposition. The sharp decline in future Nr deposition driven by profound emission abatement may substantially reduce the ecological damages like acidification and eutrophication. Meanwhile, it might potentially weaken the carbon sink capacity of terrestrial ecosystems. A comprehensive consideration of the balance between Nr control and terrestrial carbon sinks is essential for the future.

Our findings are subject to some limitations. Firstly, given the computationally intensive of numerical simulation, the Nr deposition was simulated with a single model (CMAQ) in this work. As suggested by the MICS-Asia III project, there existed clear difference in Nr deposition simulation among multiple CTMs, and in particular the consistency of dry deposition of OXN was relatively poor, with coefficient of variation (CV) ranging 0.4-0.5 throughout most of China (Ge et al., 2020). Multi-model ensemble methodology is thus recommended in future work to reduce the bias of single-model simulation. Secondly, the role of climate change on future Nr deposition might be underestimated. Climate-driven effects on emissions were not considered in this study, such as the increase of NH_3 volatilization due to global warming (Ren et al., 2023). In addition, we mainly addressed the future evolution of Nr deposition under the mean state of climate, but neglected the potential impact of extreme climatic events. For example, the changing frequency of heavy precipitation was reported as a key factor influencing the variation of Nr deposition (Chen et al., 2023). Therefore, more analyses

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should be conducted on the connection between the changing extreme climate events and atmospheric deposition.

Data availability

All data in this study are available from the authors upon request.

Author contributions

MMa developed the strategy and methodology of the work and wrote the draft. Y Zhao improved the methodology and revised the manuscript. JCao provided useful comments on the paper. BZheng provided the historical emission inventory. DTong provided the future emission inventory.

Competing interests

The authors declare that they have no conflict of interest.

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Figure captions

Figure 1 Evaluations of simulated monthly average temperature at the height of 2 m (T2, a), wind speed at the height of 10 m (WS10, b), relative humidity (RH, c), and accumulated precipitation (PREC, d) in Mainland China. The dots represent the site-level observations. The normalized mean bias (NMB), normalized mean error (NME), root mean squared error (RMSE) and the correlation coefficient (R) for the comparisons are shown in the lower left corner of each panel.

Figure 2 Spatial distribution of annual averaged Nr deposition fluxes (kg N ha^{-1} yr^{-1}) for different forms and species in 2010s and the changes between 2010s and 2060s. Panels (a-d) represent the results of 2010s (Base simulation). Panels (e-h) represent future deposition changes under the SSP2-4.5 pathway (Case 1 – Base). Panels (i-l) represent the changes under the SSP5-8.5 pathway (Case 2 – Base).

Figure 3 Changes in annual total Nr deposition fluxes (kg N ha^{-1} yr^{-1}) from 2010s to 2060s attributed to climate change (a, d), emission change (b, e), and both (c, f). Panels (a-c) represent the changes under the SSP2-4.5 pathway, respectively and Panels (d-f) represent the changes under the SSP5-8.5 pathway. Domain-averaged spatial correlation (R) between the impact of climate or emission change and both is presented in panels (a, d) or (b, e).

Figure 4 Changes in OXN (a-c) and RDN deposition (d-f) from 2010s to 2060s attributed to emission variation in “Baseline”, “Current-goal” and “Neutral-goal” scenarios.

Figure 5 Relative changes in Nr emissions and deposition in WC and EC from 2010s to 2060s under different emission scenarios.

Figure 6 Predicted response (%) of OXN (a-d) and RDN deposition (e-h) to a 20% perturbation of emissions in 2010s and 2060s for different emission scenarios. The response is obtained by calculating the ratio of the percent change in deposition to that in emission.

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Tables

Table 1 Description of the designed simulation cases.

Name	Emissions input	Meteorological input
Base	MEIC, 2010-2014	ERA5 reanalysis, 2010-2014
Case1	DPEC "Current-goal", 2060	SSP2-4.5 BCMM, 2060-2064
Case2	DPEC "Baseline", 2060	SSP5-8.5 BCMM, 2060-2064
Case3	MEIC, 2010-2014	SSP2-4.5 BCMM, 2060-2064
Case4	MEIC, 2010-2014	SSP5-8.5 BCMM, 2060-2064
Case5	DPEC "Neutral-goal", 2060	SSP2-4.5 BCMM, 2060-2064
Case6	Same as Case1, but emissions in WC are maintained at 2010s levels.	SSP2-4.5 BCMM, 2060-2064
Case7	Same as Case2, but emissions in WC are maintained at 2010s levels.	SSP5-8.5 BCMM, 2060-2064
Case8	Same as Case5, but emissions in WC are maintained at 2010s levels.	SSP2-4.5 BCMM, 2060-2064
Case9	Same as Case3, but with 20% reduction in emissions for all species.	SSP2-4.5 BCMM, 2060-2064
Case10	Same as Case1, but with 20% reduction in emissions for all species.	SSP2-4.5 BCMM, 2060-2064
Case11	Same as Case2, but with 20% reduction in emissions for all species.	SSP5-8.5 BCMM, 2060-2064
Case12	Same as Case5, but with 20% reduction in emissions for all species.	SSP2-4.5 BCMM, 2060-2064

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Table 2 The normalized mean bias (NMB), normalized mean error (NME) and the correlation coefficient (R) between the simulated and observed annual Nr deposition. Dry and wet Nr deposition fluxes of oxidized nitrogen (OXN) and reduced nitrogen (RDN) averaged over 2010-2014 were evaluated separately.

	OXN_DDEP	OXN_WDEP	RDN_DDEP	RDN_WDEP
NMB (%)	-9.07	-15.12	-28.76	-17.86
NME (%)	34.76	43.24	47.17	41.72
R(temporal)	0.63	0.65	0.65	0.82
R(spatial)	0.73	0.72	0.83	0.69

Note: OXN_DDEP and OXN_WDEP indicate the dry and wet deposition of oxidized nitrogen, respectively. RDN_DDEP and RDN_WDEP indicate the dry and wet deposition of reduced nitrogen, respectively.

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Table 3 Simulated atmospheric Nr deposition fluxes (kg N ha^{-1} yr^{-1}) in China averaged over 2010-2014 and 2060-2064 under different SSP-RCP pathways.

Periods	Species	Dry	Wet	Total
2010-2014 (Case 1)	OXN	3.7	3.4	7.1
	RDN	3.0	4.6	7.6
	OXN + RDN	6.7	8.0	14.7
2060-2064 under SSP2-4.5 (Case 2)	OXN	1.5	1.6	3.1
	RDN	2.9	3.0	5.9
	OXN + RDN	4.4	4.6	9.0
2060-2064 under SSP5-8.5 (Case 3)	OXN	4.0	4.4	8.4
	RDN	2.9	4.1	7.0
	OXN + RDN	6.9	8.5	15.4

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Table 4 Simulated domain-averaged OXN deposition fluxes (kg N ha^{-1} yr^{-1}) over EC for cases where emissions change to 2060s levels in all regions as well as cases where emissions in WC are maintained at 2010s levels. Relative changes (%) are calculated by comparing cases with 2060s emission levels in all regions to cases with 2010s emission levels in WC, then dividing the difference by the 2010s emission levels in WC.

	Emissions in WC are maintained at 2010s levels	Emissions change to 2060s levels in all regions	Relative change
"Baseline"	13.29 (Case7)	13.59 (Case2)	2%
"Current-goal"	5.36 (Case6)	5.08 (Case1)	-6%
"Neutral-goal"	1.90 (Case8)	1.39 (Case5)	-27%

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Table 5 Regional average responses (%) of OXN or RDN deposition to a 20% emission reduction in 2010s and 2060s under different emission scenarios over NC and SC and the whole of mainland China.

	NC	SC	China
	Responses (%) of OXN deposition to NO_x emissions		
2010s	82.60	96.19	82.71
2060s under "Baseline"	83.95	92.54	88.41
2060s under "Current-goal"	91.86	103.00	81.17
2060s under "Neutral-goal"	94.59	98.07	68.83
	Responses (%) of RDN deposition to NH_3 emissions		
2010s	103.11	97.63	96.30
2060s under "Baseline"	104.67	98.42	98.05
2060s under "Current-goal"	100.70	95.99	94.38
2060s under "Neutral-goal"	97.12	95.47	92.44

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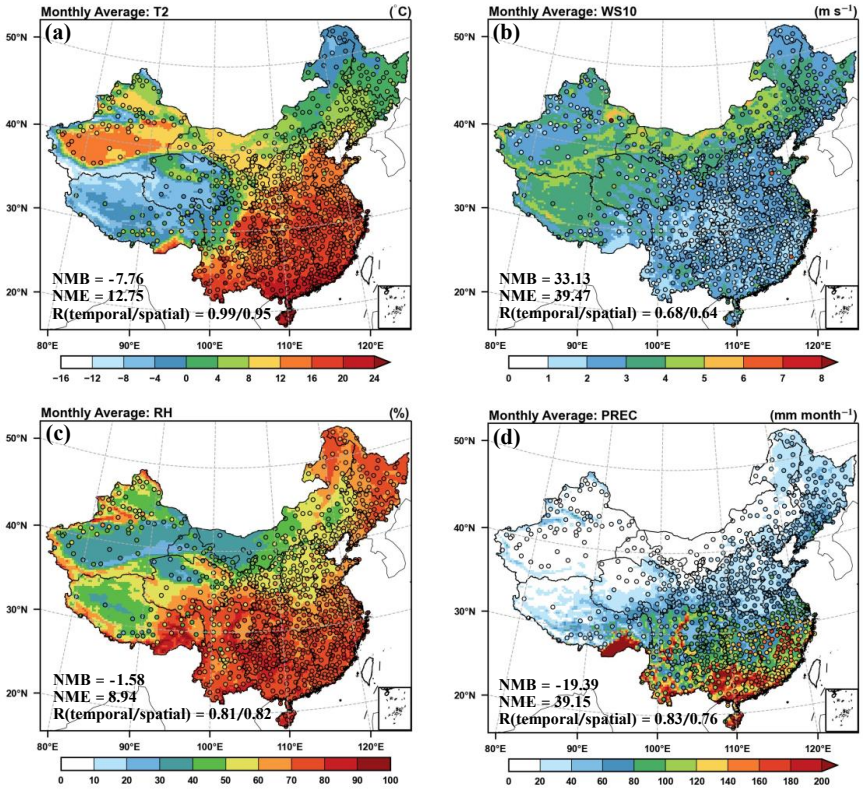
1127 **Figure 1**



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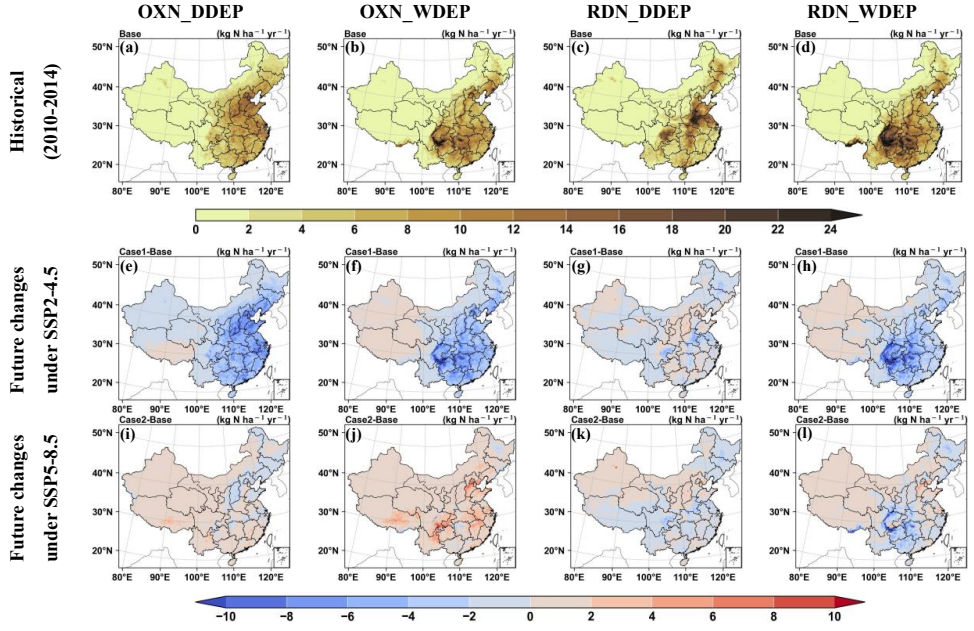
1129 **Figure 2**



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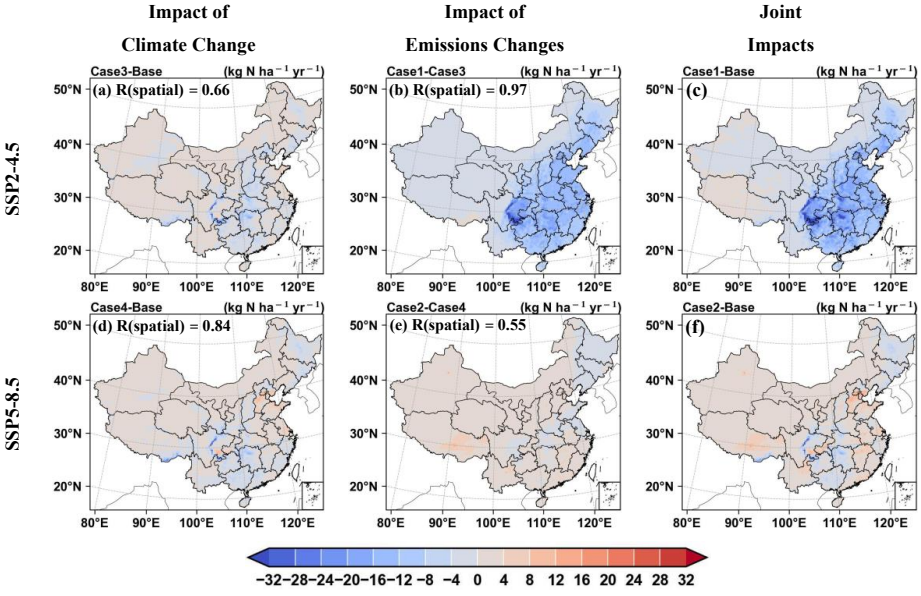
1131 **Figure 3**



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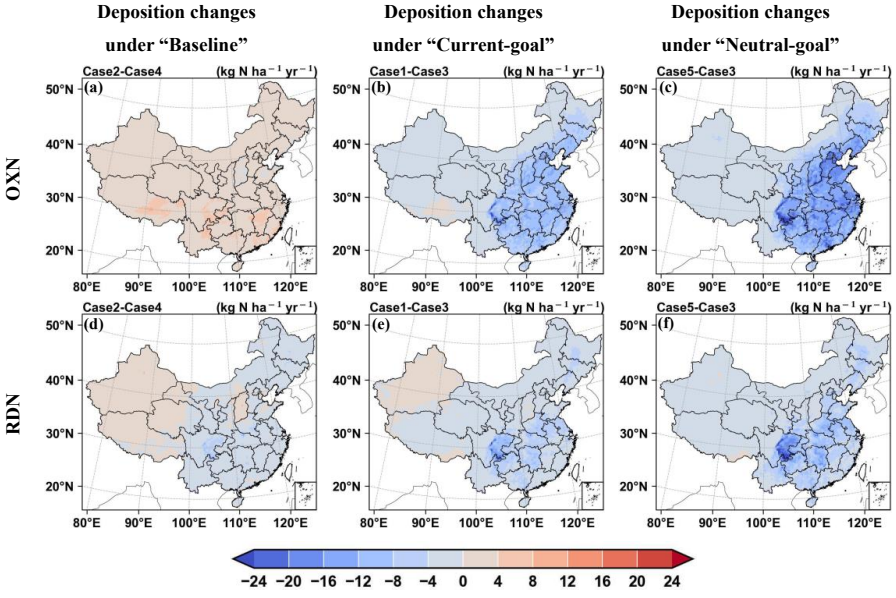
1133 **Figure 4**



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1136 **Figure 5**

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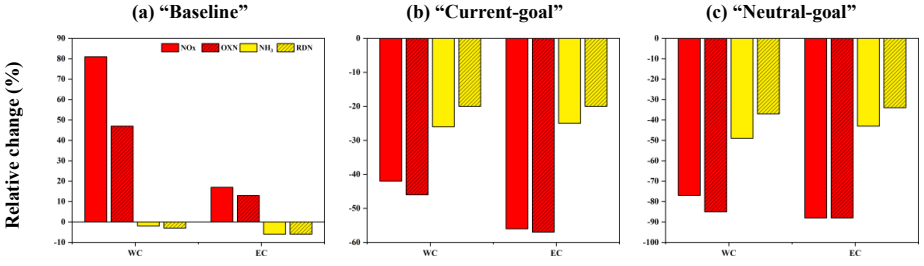

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Figure 6