Global estimates of ambient reactive nitrogen components during 2000-2100 based on the multi-stage model

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Abstract

High contents of reactive nitrogen components aggravate air pollution and could also impact ecosystem structure and function across the terrestrial-aquatic-marine continuum. However, the long-term historical trends and future prediction of reactive nitrogen components at the global scale still remains high uncertainties. In our study, the field observations, satellite products, model output, and many other covariates were integrated into the multi-stage machine-learning model to capture the global patterns of reactive nitrogen components during 2000-2019. In order to decrease the estimate uncertainties in the future scenarios, the constructed reactive nitrogen component dataset during the historical period was then utilized as the constraint to calibrate the CMIP6 dataset in four scenarios. The results suggested the cross-validation (CV) R^2 values of four species showed satisfied performance ($R^2 > 0.55$). The concentrations of estimated reactive nitrogen components in China experienced persistent increases during 2000-2013, while they suffered from drastic decreases since 2013 except NH_3. It might be associated with the impact of clean air policy. However, these compounds in Europe and the United States remained relatively stable since 2000. In the future scenarios, SSP3-7.0 (traditional energy scenario) and SSP1-2.6 (carbon neutrality scenario) showed the highest and lowest reactive nitrogen component concentrations, respectively. Although the reactive nitrogen concentrations in some heavy-pollution scenarios (SSP3-7.0) also experienced decreases during 2020-2100, SSP1-2.6 and SSP2-4.5 (middle emission scenario) still kept more rapid decreasing trends. Our results emphasize the need for carbon-neutrality pathway to reduce global atmospheric N pollution.

1. Introduction

Along with the development of global urbanization and industrialization, the anthropogenic emissions of reactive nitrogen (e.g., NO_x, NH_3) experienced drastic increases during the past decades, and caused the higher concentrations of NO_2, NH_3, and many secondary components such as NO_3^- (NO_3-N), NH_4^+ (NH_4-N), and HNO_3 (Chen et al., 2021; Liu et al., 2020b; McDuffie et al., 2020). The reactive nitrogen released from anthropogenic source could significantly alter the global nitrogen cycle throughout the Earth system (Altieri et al., 2021; Zhang et al., 2020). Reactive nitrogen in the atmosphere dominates the chemical formation of tropospheric O_3 and aggravates the particle pollution (Geddes and Martin, 2017), with implications for global air quality and climate change (He et al., 2022; Von Schneidemesser et al., 2015). Moreover, the ambient reactive nitrogen could be deposited into the land surface and could cause lake eutrophication and soil acidification (Bouwman et al., 2002; Chen et al., 2018). Therefore, it is highly necessary to understand the spatial distributions and temporal evolution trends of reactive nitrogen components at the global scale.

Despite the global importance, observational constraints on reactive nitrogen in the atmosphere were still scarce in most parts of the world (Liu et al., 2020b). Furthermore, the majority of monitoring sites focused on China, Europe, and the United States (Du et al., 2014; Li et al., 2020; Li et al., 2019a; Li et al., 2016), and these uneven sites only possessed limited spatial representativeness (Shi et al., 2018), which restricted the accurate assessment of global reactive nitrogen pollution. Fortunately, the satellite observations gave us a unprecedent chance to capture the global variations of atmospheric reactive nitrogen. Geddes et al. (2017) used the satellite products to calibrate the simulated reactive nitrogen oxides (NO_y) and improved the predictive performance (R = 0.83) compared with the chemical transport model (CTM) output alone (Geddes and Martin, 2017). Afterwards, Liu et al. (2022) also used the similar method to estimate the global wet deposition of reduced nitrogen (NH_4^+) and the R value achieved 0.80(Liu et al., 2021). Although the calibration based on satellite products could improve the predictive accuracy compared with CTM output, the simulated values still largely biased from the ground-level observations. Moreover, the method cannot accurately fill the gaps of reactive nitrogen concentrations without satellite coverage. In our previous works, we developed a satellite-based ensemble machine-learning model to predict the wet NH_4^+ deposition across China and the R^2 value reached 0.76 (R = 0.88) (Li et al.,

2020). However, this technique was not expanded to the global scale and the high-accuracy and full-coverage global ambient reactive nitrogen dataset was still lack.

Apart from the historical estimates, the future prediction of reactive nitrogen is also important because these components in the future scenarios could significantly affect the land carbon cycle and greenhouse gas emissions, both of which could aggravate the global climate change and affect the earth system safety (Chen et al., 2015; Zaehle, 2013). To the best of our knowledge, only two studies focused on global aerosol prediction in the future scenarios. Chen et al. (2023) predicted the global $PM_{2.5}$ levels and associated mortalities in 2100 under different climate scenarios and found that SSP3-7.0 scenario was linked with the highest $PM_{2.5}$ exposure. Li et al., (2022) also simulated the global NO_3^- (NO_3-N) and NH_4^+ (NH_4-N) levels in the future four scenarios and demonstrated that both of these components showed marked decreases in most cases except SSP5-8.5 scenario. However, this study predicted the future reactive nitrogen based on historical CTM output alone, which lacks of observation constraints. The result might increase the uncertainty of assessment.

In our study, we developed a multi-stage model to estimate the concentrations of four reactive nitrogen species (NO_3^- (NO_3-N), HNO_3, NH_3, and NH_4^+ (NH_4-N)) during 2000-2019 because these species were most important reactive nitrogen components for human health and ecological ecosystem and also showed abundant ground-level observations. Then, the species over the 2020-2100 period under the SSP1-2.6, SSP2-4.5, SSP3-7.0, and SSP5-8.5 scenarios were also corrected based on the historical estimates. Finally, the long-term dataset of reactive nitrogen during 2000-2100 was constructed. Our results were beneficial to assess the impacts of reactive nitrogen components on air pollution and climate change in the future.

2. Material and methods

2.1 Reactive nitrogen observations

Most of reactive nitrogen observations focused on East Asia, Europe, and the United States. The monthly reactive nitrogen components monitoring data during 2010-2015 in China were downloaded from nationwide nitrogen deposition monitoring network (NNDMN) including 32 sites, and these sites could be classified into three types mixed with urban, rural, and background sites (Xu et al., 2019) (Table S1). The concentrations of reactive nitrogen components were determined using the active DEnuder for Long-Term Atmospheric sampling system (DELTA). The detailed

88   sampling and analysis procedures have been described by Xu et al. (2019). The dataset of reactive

89   nitrogen components in other countries of East Asia during 2000-2019 were download from the

90   Acid Deposition Monitoring Network in East Asia (EANET), which includes 41 sites. The European

91   Monitoring and Evaluation Programme (EMEP) provides records of long-term reactive nitrogen

92   components in 86 sites of most countries across West Europe. Monthly reactive nitrogen

93   components dataset in 84 locations across the United States could be obtained from the Clean Air

94   Status and Trends Network (CASTNET) (Figure S1).

95   2.2 Data preparation

96   The GEOS-Chem (v13.4.0) model driven by MERRA2 meteorological parameters was applied

97   to simulate the historical reactive nitrogen components (daily) during 2000-2019 (Feng et al., 2021).

98   The GEOS-Chem model was composed of detailed ozone-NO_x-VOC-PM-halogen tropospheric

99   chemistry. The grid version of the model with a horizontal resolution of $2° × 2.5°$ was utilized. Wet

100   deposition contained many processes including sub-grid scavenging in convective updrafts, in-

101   cloud rainout, and below-cloud washout (Liu et al., 2001). Dry deposition was estimated based on

102   a resistance-in-series model (Wesely, 2007). The estimates of aerosol optical properties account for

103   the hygroscopic growth (Drury et al., 2010). Vertical mixing in the boundary layer follows a non-

104   local scheme implemented by Lin and McElroy (2010), and convection employs the relaxed

105   Arakawa-Schubert scheme. The anthropogenic emission inventory in 2000-2019 was downloaded

106   from the website of Community Emissions Data System (CEDS) (Hoesly et al., 2018). CEDS

107   emission inventory includes eight sectors such as agriculture, energy, industry, residential, shipping,

108   solvents, transportation, and waste incineration. Then, the daily reactive nitrogen components were

109   averaged to the monthly scale.

110   The IASI instrument aboard on the polar sun-synchronous MetOp platform traverses the

111   equation twice each day (9:30 a.m. and 9:30 p.m. local solar time) (Whitburn et al., 2016a). The

112   measurements in the daytime usually shows the better accuracy than those at night due to the high

113   sensitivity to ambient NH_3 (Van Damme et al., 2017; Whitburn et al., 2016a; Whitburn et al., 2016b).

114   In our study, we used the IASI NH_3 columns in morning during 2008-2019 to estimate the NH_3 and

115   NH_4^+ concentrations globally. Besides, the NH_3 column dataset with a cloud shield higher than 25%

116   and relative error above 100% were eliminated.

The tropospheric vertical column density (VCD) of NO_2 retrieved from OMI aboard on Aura satellite crosses the earth once a day (Kim et al., 2016). OMI-derived tropospheric NO_2 column densities during 2005-2019 was applied to develop the model. The tropospheric NO_2 column density data with cloud radiance fraction > 0.5, terrain reflectivity > 30%, and solar zenith angles > 85° were screened (Cooper et al., 2022). Additionally, the NO_2 columns from GOME (1995-2003), SCIAMACHY (2002-2011) and GOME-2 (2007-) were also collected to simulate the NO_3^- (NO_3-N) and HNO_3 levels. The similar overpass time of these three instruments (from about 09:30 to 10:30 LT, local time) facilitates the simultaneous use to capture consistent long-term coverage. However, the dataset cannot cover the NO_2 columns since 2017. To overcome the inconsistency of these satellite products, we applied the linear regression technique to construct the relationship between OMI-NO_2 columns and GOME/SCIAMACHY NO_2 columns. The results suggested these satellite products showed good relationship (R^2 > 0.6). At last, the long-term (2000-2019) NO_2 columns at the global scale were constructed.

The monthly meteorological parameters derived from ERA-5 comprise of 2 m dewpoint temperature (D_{2m}), 2 m temperature (T_{2m}), surface pressure (S_p), and total precipitation (T_p), 10 m U wind component (U10), and 10 m V wind component (V10). The population density data during 2000-2020 around the world were downloaded from https://hub.worldpop.org/geodata/listing?id=64. The elevation data was extracted from ETOPO at a spatial resolution of 1' (Amante and Eakins, 2009) (https://rda.ucar.edu/datasets/ds759.4/). In addition, the land use types including cropland, forest, grassland, shrubland, tundra, barren land, and snow/ice were obtained from Liu et al. (2020a). Besides, the CMIP6 dataset in four scenarios were also applied to predict the reactive nitrogen concentrations during 2020-2100. The dataset includes 2-m air temperatures, wind speed at 850 and 500 hPa, total cloud cover, precipitation, relative humidity, and short-wave radiation. The modelled meteorological parameters derived from 16 earth system models were incorporated into the machine-learning model. The detailed models are summarized in Table S2.

2.3 Model development

A three-stage model was established to capture the full-coverage reactive nitrogen dataset at the global scale (Figure 1). In the first stage, the ground-level reactive nitrogen species, satellite

146   products (e.g., OMI-NO$_2$ and IASI-NH$_3$ columns), meteorological parameters, land use types,

147   population, and simulated reactive nitrogen components derived from GEOS-Chem model were

148   collected as the independent variables to estimate the gridded reactive nitrogen species at the

149   period/grid with satellite product based on XGBoost algorithm. In the second stage, the

150   meteorological parameters, GEOS-Chem output, land use types, and population were applied to fill

151   the gaps without satellite retrievals. Then, the simulated results based on these models were fused

152   to obtain the full-coverage reactive nitrogen components and the ground-level observations were

153   further used to calibrate the full-coverage dataset and the final reactive nitrogen components at the

154   global scale were simulated. In the last stage, the reactive nitrogen components and meteorological

155   parameters in four scenarios (SSP1-2.6, SSP2-4.5, SSP3-7.0, and SSP5-8.5) during 2020-2100 were

156   collected from CMIP6 dataset including 16 earth system models (Table S2). Then, the data in the

157   future scenarios were integrated into the ensemble model including XGBoost, LightGBM, and

158   convolutional neural networks (CNN) to further calibrate the modeling results based on historical

159   dataset (2000-2019) derived from previous two-stage model. The detailed equations of multiple

160   machine-learning models are summarized as follows:

161   (1) XGBoost model

162

$$F^{(t)} = \sum_{i=1}^{n}[l(y_i, \hat{y}^{(t-1)}) + \partial_{y^{(t-1)}} l(y_i, \hat{y}^{(t-1)}) f_t(x_i) + \frac{1}{2}\partial^2_{y^{(t-1)}} l(y_i, \hat{y}^{(t-1)}) f_t^2(x_i)] + \Omega(f_t) \quad (1)$$

163   where F$^{(t)}$ is the cost function at the t-th period; ∂ is the derivative of the function; $\partial^2_{y^{(t-1)}}$

164   represents the second derivative of the function; l denotes the differentiable convex loss function

165   that reveals the difference of the predicted value ($\hat{y}$) of the i-th instance at the t-th period and the

166   target value (y$_i$); f$_t$(x) represents the increment; $\Omega(f_t)$ reflects the regularizer. Maximum tree

167   depth and learning rate reached 15 and 0.1, respectively.

168   (2) LightGBM model

169

$$\hat{f} = \arg\min_f E_{y,X} Q(y, f(x)) \quad (2)$$

170

$$f_T(X) = \sum_{t=1}^{T} f_t(X) \quad (3)$$

171 where Q(y,f(x)) reflects the a specific loss function; $\sum_{t=1}^{T} f_t(X)$ denotes the regression trees.

172 Maximum tree depth, learning rate, and feature fraction reached 25, 0.2, and 0.7, respectively.

173 (3) CNN model

174 The reactive nitrogen species and meteorological parameters in the future scenarios were applied to

175 CNN model based on the historical (2000-2019) reactive nitrogen species derived from stage 1-2

176 model.

177
$$x \xrightarrow{\ f:U-Net\ } y \qquad (4)$$

178 where x (x_1, x_2,……, x_n) represents the reactive nitrogen species and meteorological parameters

179 derived from CMIP6 dataset; y (y_1, y_2, ……, y_n) denotes the historical (2000-2019) reactive nitrogen

180 species.

181   All of the convolution layers showed the same kernel size of 3×3 and used rectified linear unit

182 (ReLU) as the activation function. Max-pooling layers were employed for adjusting the size of

183 images to capture better bottleneck information. After each block, the image size could be halved

184 by using the max pooling layer with kernel size of 2×2, but the number of channels will be doubled.

185 In our study, the learning rate was set as 0.1 to achieve the best performance.

186   All of the independent variables collected from multiple sources were resampled to 0.25° grids

187 using Kriging interpolation. For example, both of the population density and land use types in each

188 grid were calculated using spatial clipping toolbox. Later on, all of these variables were combined

189 to develop the model. During the development of multi-stage model, it was highly imperative to

190 remove some redundant explanatory variables and then determine the optimal variable group. The

191 redundant variables means that the overall predictive accuracy could degrade after the removal of

192 these variables.

193 **3. Results and discussion**

194 3.1 The modelling performance of historical reactive nitrogen estimates

195   The multi-stage model was applied to capture the spatiotemporal variations of reactive nitrogen

196 concentrations during 2000-2100. In our study, we employed XGBoost model to construct the full-

197 coverage reactive nitrogen dataset during 2000-2020. The cross-validation (CV) R^2 values of the

198 model for NO_3^- (NO_3-N), HNO_3, NH_3, and NH_4^+ (NH_4-N) estimates reached 0.67, 0.62, 0.58, and

199   0.60, respectively (Figure 2). RMSE of NO_3^- (NO_3-N), HNO_3, NH_3, and NH_4^+ (NH_4-N) were 0.55,

200   0.23, 2.32, and 1.71 µg N m^{-3}, respectively. MAE of NO_3^- (NO_3-N), HNO_3, NH_3, and NH_4^+ (NH_4-

201   N) reached 0.19, 0.13, 1.23, and 0.59 µg N m^{-3}. The CV R^2 values of NO_3^- (NO_3-N), HNO_3, and

202   NH_4^+ (NH_4-N) estimates were significantly higher than Jia et al. (2016) (0.22, 0.41, and 0.49), while

203   the CV R^2 value of NO_3^- estimate in our study was comparable to Geddes et al. (2017) (0.68)

204   (Geddes and Martin, 2017). The CV R^2 value of NH_3 estimates were also close to the results

205   obtained by Liu et al. (2019) (0.45-0.71) (Liu et al., 2019). Overall, the predictive performances

206   historical reactive nitrogen was satisfied. Although the CV R^2 values in our study were not

207   significantly higher than those in some previous studies, our study developed the full-coverage (gap-

208   free) ambient reactive nitrogen dataset, which was superior to some previous studies. Based on the

209   constructed full-coverage reactive nitrogen dataset, we also developed the ensemble model to

210   calibrate the CMIP6 dataset in the future scenarios. The CV R^2 values of the model for NO_3^- (NO_3-

211   N), HNO_3, NH_3, and NH_4^+ (NH_4-N) estimates in the future scenarios reached 0.62, 0.67, 0.56, and

212   0.60, respectively (Figure S2). RMSE of NO_3^- (NO_3-N), HNO_3, NH_3, and NH_4^+ (NH_4-N) were 0.58,

213   0.26, 2.12, and 1.91 µg N m^{-3}, respectively. MAE of NO_3^- (NO_3-N), HNO_3, NH_3, and NH_4^+ (NH_4-

214   N) reached 0.22, 0.22, 1.04, and 0.65 µg N m^{-3}. Overall, the ensemble model for these species in the

215   future scenarios still showed satisfied performance, and thus the result could be treated to be robust.

216   3.2 The spatial patterns of nitrogen reactive components

217   The global annual mean concentrations of NO_3^-, HNO_3, NH_3, and NH_4^+ during 2000-2019

218   ranged from 0.03 to 9.08, 0.03 to 1.73, 0.21 to 13.9, and 0.08 to 17.1 µg N m^{-3} with the mean values

219   of 0.43 ± 0.24 (standard deviation over grids), 0.28 ± 0.13, 1.79 ± 0.85, and 0.65 ± 0.36 µg N m^{-3}

220   (Figure S3), respectively. East Asia especially China, West Europe, and the United States obtained

221   widespread attention due to the developed economy and dense anthropogenic activity.

222   In China, the overall mean ambient NO_3^- (NO_3-N), HNO_3, NH_3, and NH_4^+ (NH_4-N)

223   concentrations reached 1.05 ± 0.62, 0.35 ± 0.19, 4.05 ± 1.84, and 2.38 ± 1.26 µg N m^{-3}, ranging

224   from 0.07-9.08, 0.06-1.73, 0.84-11.6, and 0.18-13.1 µg N m^{-3}. At the regional scale, the annual mean

225   NO_3^-, HNO_3, NH_3, and NH_4^+ concentrations followed the order of North China Plain (NCP) (4.38,

226   1.12, 7.22, and 7.69 µg N m^{-3}) > Sichuan Basin (2.40 ± 1.01, 0.52 ± 0.28, 4.92 ± 1.71, and 6.02 ±

227   1.82 µg N m^{-3}) (Figure 3). NCP displayed the higher NO_3^- and HNO_3 concentrations owing to dense

228 human activities and strong industry foundation (Qi et al., 2023; Wen et al., 2018), which could emit

229 a large amount of NO_x to the atmosphere. In both of Yangtze River Delta (YRD) and Pearl River

230 Delta (PRD), the combustion of fossil fuels and traffic emissions might be the major source of NO_x

231 emission, which aggravated nitrate events via gas-particle conversion processes (Huang et al., 2017;

232 Li et al., 2017). For Sichuan Basin, the poor topographical or meteorological conditions were major

233 factors responsible for the severe nitrate pollution (Zhang et al., 2019). It was not surprising that

234 high ambient NH_3 concentrations focused on NCP and Sichuan Basin because many croplands (dry

235 land) are distributed on these regions (Karra et al., 2021; Potapov et al., 2022), which was the major

236 source of NH_3 emissions with frequent N fertilizer applications (Ma et al., 2022). Besides, N manure

237 was another major source of NH_3 emissions in China, and the percentage of N manure to NH_3

238 emissions exceeds 50% (Kang et al., 2016). The spatial pattern of NH_4^+ level was in good agreement

239 with the NH_3 concentration because NH_4^+ was often generated from the reaction of NH_3 with SO_2

240 and NO_2 (Ehrnsperger and Klemm, 2021). Apart from China, many other countries such as South

241 Korea and Japan also showed the higher ambient reactive nitrogen concentrations. As shown in

242 Figure 3, the higher reactive N concentrations occurred on the western coasts of South Korea than

243 on the eastern coasts. The higher reactive N concentrations in Japan mainly focused on the urban

244 areas around Tokyo, which might be linked with the dense anthropogenic emission in this region

245 (Li et al., 2024). In Southeast Asia, Indonesia (NO_3^- (NO_3-N), HNO_3, NH_3, and NH_4^+ (NH_4-N): 0.18,

246 0.47, 5.72, and 0.44 µg N m^{-3}) suffered from the most serious reactive N pollution compared with

247 other surrounding countries.

248   In Europe, the ambient NO_3^- (NO_3-N), HNO_3, NH_3, and NH_4^+ (NH_4-N) concentrations ranged

249 from 0.13 to 2.84, 0.06 to 0.92, 0.35 to 7.81, and 0.22 to 3.77 µg N m^{-3}, respectively. The annual

250 mean NO_3^- (NO_3-N), HNO_3, NH_3, and NH_4^+ (NH_4-N) levels reached 0.57 ± 0.28, 0.25 ± 0.11, 1.58

251 ± 0.68, and 0.89 ± 0.42 µg N m^{-3}, respectively (Figure 4). High concentrations of reactive nitrogen

252 components focused on the northern part of Italy, central and southern part of Germany, North

253 France, Poland, and the western part of Russia, which was in good agreement with the spatial pattern

254 of NO_x and NH_3 emissions (Luo et al., 2022; Qu et al., 2020). Emissions Database for Global

255 Atmospheric Research (EDGAR) suggested that N fertilization and N manure accounted for 43%

256 and 53% of total NH_3 emissions in western Europe (Liu et al., 2019), respectively. Furthermore, Liu

et al. (2019) confirmed that a good relationship between ambient NH_3 level and N fertilization plus N manure (r = 0.62) was observed in Europe. Cooper et al. (2017) employed the inversion model to estimate NO_x emission in Europe and also found that high NO_x emission was also mainly distributed on North France, Germany, the northern part of Italy, and Russia, which partly explained the higher concentrations of reactive nitrogen components in these regions.

In the United States, the ambient NO_3^- (NO_3-N), HNO_3, NH_3, and NH_4^+ (NH_4-N) concentrations reached 0.28 ± 0.12, 0.19 ± 0.08, 2.12 ± 0.66, and 0.49 ± 0.25 μg N m^{-3}, with the range of 0.03-2.35, 0.03-1.31, 0.26-9.96, and 0.10-6.09 μg N m^{-3} (Figure 5), respectively. The hotspots of NO_3^- (NO_3-N), HNO_3, and NH_4^+ (NH_4-N) levels focused on the eastern part of the United States, while the higher NH_3 concentration focused on Central Great Plains and some regions in California such as San Joaquin Valley (6.15 μg N m^{-3}). Both of bottom-up and top-down NO_x and NH_3 emissions suggested that the spatial distributions of reactive nitrogen components were strongly dependent on the precursor emissions (McDuffie et al., 2020; Qu et al., 2020).

Besides, some other regions such as India (1.4, 0.5, 6.6, and 4.4 μg N m^{-3}) especially the northern part of India (3.1, 0.8, 12.6, and 8.4 μg N m^{-3}) also experienced severe reactive N pollution in the atmosphere. Meanwhile, some countries in South America such Brazil and Argentina and in Africa such as West Africa Coast (Nigeria, Ivory Coast, Ghana, Togo, and Benin) (HNO_3 and NH_3: 0.3 and 5.0 μg N m^{-3}) and Democratic Congo (0.4 and 1.6 μg N m^{-3}) also suffered from serious HNO_3 (Brazil and Argentina: 0.3 and 0.2 μg N m^{-3}) and NH_3 (3.6 and 2.8 μg N m^{-3}) pollution. The higher ambient NH_3 concentration focused on the northern part of India might be contributed by two major reasons. First of all, the intensive agricultural activities and high air temperature might be responsible for the higher NH_3 level (Cui, 2023; Wang et al., 2020). Moreover, the relatively low sulfur dioxide (SO_2) and nitrogen oxides (NO_x) emissions coupled with high air temperature restricted the gas-to-particle conversion of NH_3 (Wang et al., 2020). The severe HNO_3 and NH_3 pollution in Brazil, Argentina, and West Africa Coast might be also linked with the dense agricultural activities (Huneeus et al., 2017).

3.3 The seasonal variations of reactive nitrogen components

The ambient NO_3^- (NO_3-N), HNO_3, NH_3, and NH_4^+ (NH_4-N) concentrations exhibited significant seasonal variations (Figure S4-8). NO_3^-, HNO_3, and NH_4^+ displayed the highest and

286   lowest values in winter (December-February) and summer (June-August), respectively. On the one

287   hand, the anthropogenic NO_x emission for domestic heating might be higher in winter compared

288   with other seasons (Lin et al., 2011). On the other hand, the stagnant meteorological conditions

289   limited the pollutant diffusion (Li et al., 2019b; Liu et al., 2020c). Meanwhile, the higher relative

290   humidity in winter facilitated the formation of NH_4NO_3 (Huang et al., 2016; Xu et al., 2012).

291   However, both of ambient NO_3^- and NH_4^+ concentrations showed the lower concentrations in

292   summer, which might be attributable to the decomposition of NH_4NO_3 under the condition of high

293   air temperature. In contrast to the secondary inorganic nitrogen, the ambient NH_3 level showed the

294   highest concentration in summer (1.71 ± 0.45 μg N m^{-3}). China, Europe, and the United States

295   suffered from similar NH_3 peaks in summer (4.20 ± 1.85, 1.77 ± 0.65, and 2.21 ± 1.04 μg N m^{-3}).

296   There are two reasons accounting for the fact. At first, mineral N fertilizer or manure application

297   was mainly performed in spring and early summer (Paulot et al., 2014). Many field observations

298   have obtained similar NH_3 peak in summer (He et al., 2021; Pan et al., 2018). Moreover, summer

299   often showed the higher air temperature, which promotes the volatilization of ammonium and limits

300   the gas-to-particle of gaseous NH_3 (Liu et al., 2019).

301   3.4 The historical trends of reactive nitrogen components during 2000-2019

302   The long-term trends of ambient NO_3^- (NO_3-N), HNO_3, NH_3, and NH_4^+ (NH_4-N) concentrations

303   are shown in Figure 6 and Figure S9-12. The NO_3^- concentration in China displayed rapid increase

304   (9.7%/yr) during 2000-2007, and then it kept the moderate increase (4.2%/yr) during 2007-2013.

305   However, the NO_3^- (NO_3-N) concentration in China experienced the drastic decrease (-2.6%/yr)

306   since 2013. The ambient HNO_3 and NH_4^+ (NH_4-N) concentrations showed similar trends during this

307   period. Due to the impact of Clean Air Action, the concentrations of gaseous precursors (e.g., SO_2

308   and NO_x) suffered from substantial decreases, which could be transformed into nitrate and

309   ammonium via heterogeneous reactions (Huang et al., 2019). However, the decreasing rates of NO_3^-

310   were still much lower than those of gaseous precursors (Li et al., 2023). On the one hand, it might

311   be associated with the increased O_3 level and enhanced atmospheric oxidation capacity (AOC),

312   which led to an increase in the photochemical reaction rate of the secondary components (Wang et

313   al., 2019). On the other hand, strong SO_2 emission control under the Clean Air Action allowed more

314   gaseous NH_3 to form nitrate. The ambient NH_3 level remained relatively stable status during 2000-

2013, while it experienced rapid increases after 2013. The result was in good agreement with Liu et al. (2019). In fact, the ambient NH_3 level in North China Plain still experienced dramatic increase (> 0.2 μg N m^{-3}/yr) during 2000-2013 because enhanced agricultural activities. Zhang et al. (2017) have demonstrated that the livestock manure and fertilizer application generally accounted for 43% and 36% of the agricultural NH_3 emission, respectively. Since 2013, the NH_3 concentration in the entire China suffered from rapid increase, which might be associated with the drastic decrease of sulfate. It was well known that NH_3 could react with HNO_3 and gaseous H_2SO_4 to generate ammonia sulfate and ammonia nitrate (Wang et al., 2022; Wang et al., 2019). Substantial decreases of acidic gases (e.g., SO_2) lead to the reduction of NH_3 conversion to ammonia salts in the atmosphere (Chen et al., 2019), which result in excess NH_3 remaining in the gaseous phase. Different from China, the reactive N concentrations in some other Asia and Africa countries especially India (NO_3^-, HNO_3, NH_3, and NH_4^+: 54%, 46%, 11%, and 94%), South Korea (76%, 42%, 40%, and 9%), Indonesia (21%, 5%, 14%, and 41%), Democratic Congo (9%, 16%, 145%, 41%), and West Africa Coast (4%, 11%, 37%, and 106%) still exhibited stable increases during 2000-2019. The results indicated that no strong reactive N emission control measures were implemented in these countries, which should be further exerted imperatively.

Compared with China, the long-term trends of reactive nitrogen components Europe and the United States were relatively stable. In the Europe, the concentrations of NO_3^- (NO_3-N), HNO_3, and NH_3 exhibited increases during 2000-2007 (0.7%/yr, 2.3%/yr, and 2.1%/yr), while they experienced slight decreases after 2007. The NH_4^+ (NH_4-N) level displayed continuous decrease since 2000. The result was closely linked with the trends of NO_x and NH_3 emissions derived from satellite retrieval (Cooper et al., 2017; Luo et al., 2022). In the United States, both of NO_3^- and NH_4^+ showed persistent decreases during 2000-2019. Zhang et al. (2018) have confirmed that NO_x emission in the eastern US has experienced persistent decrease since 1990, which facilitated the decreases of NO_3^- and NH_4^+ levels. However, the ambient HNO_3 and NH_3 concentrations displayed slight increases during 2000-2007 (2.1%/yr), and then remained relatively stable since 2007. Liu et al. (2019) also found similar characteristic of ambient NH_3 trend in the United States. In fact, the NH_3 concentrations in the Middle Plain and eastern US still showed increases due to the lack of NH_3 emission control policies as well as the decline in acidic gases (Warner et al., 2017). The reactive N concentrations

344　in some countries in South America such as Brazil (9%, 0%, 13%, and 34%) and Argentina (10%,

345　12%, 18%, and 7%) also remained relatively stable because local anthropogenic emission of reactive

346　N did not show dramatic increases in the past two decades (McDuffie et al., 2020).

347　3.5 Projection of future ambient reactive nitrogen components

348　　　For the future reactive nitrogen component estimates, the ensemble model was applied to

349　predict the reactive nitrogen component concentrations under the SSP1-2.6, SSP2-4.5, SSP3-7.0,

350　and SSP5-8.5 scenarios. SSP1-2.6 represents the low emission pathways. In SSP1-2.6, the projected

351　average NO_3^- concentrations in most countries experienced rapid decreases from 2020 to 2100

352　(Figure 7 and Table 1). The mean concentrations of NO_3^- in China, India, Europe, and the United

353　States decreased from 1.16 ± 0.35, 1.23 ± 0.42, 0.41 ± 0.14, and 0.27 ± 0.09 $\mu g\ N\ m^{-3}$ to 0.33 ± 0.10,

354　0.65 ± 0.21, 0.10 ± 0.03, and 0.06 ± 0.02 $\mu g\ N\ m^{-3}$ during 2020-2100 in SSP1-2.6 scenario,

355　respectively. Besides, the NO_3^- concentrations in many other countries of Africa and South America

356　such as Brazil (-127%) and Democratic Congo (-162%) also suffered from drastic decreases in this

357　scenario. SSP2-4.5 scenario represents the middle range of plausible future pathways (Nazarenko

358　et al., 2022). In this scenario, the predicted average NO_3^- concentrations in China, India, Europe,

359　and the United States decreased from 1.19 ± 0.40, 1.43 ± 0.35, 0.44 ± 0.13, and 0.24 ± 0.08 $\mu g\ N$

360　m^{-3} to 0.41 ± 0.14, 0.95 ± 0.32, 0.24 ± 0.08, and 0.05 ± 0.02 $\mu g\ N\ m^{-3}$ during 2020-2100, respectively.

361　SSP3-7.0 and SSP5-8.5 denote the less investment in the environment and heavily relies on

362　traditional energy for rapid economic development, respectively. The ambient NO_3^- in these

363　scenarios generally showed the higher concentrations compared with other scenarios. For instance,

364　the NO_3^- concentrations in China reduced from 1.25 ± 0.40 (SSP3-7.0) and 1.21 ± 0.39 (SSP5-8.5)

365　to 0.75 ± 0.25 (SSP3-7.0) and 0.58 ± 0.18 (SSP5-8.5) $\mu g\ N\ m^{-3}$ during 2020-2100 (Figure 8),

366　respectively. The higher NO_3^- concentrations in SSP3-7.0 and SSP5-8.5 might be associated with

367　the higher anthropogenic NO_x emission. Compard with SSP1-2.6 and SSP2-4.5, the NO_3^-

368　concentrations in some countries during SSP3-7.0 and SSP5-8.5 scenarios displayed slight increases

369　from 2020 to 2040. For instance, the ambient NO_3^- concentrations in Indonesia increased by 12%

370　(SSP3-7.0) and 5% (SSP5-8.5), respectively.

371　　　The temporal variations of ambient HNO_3 were similar to those of NO_3^- concentrations. The

372　mean concentrations of HNO_3 in China, India, Europe, and the United States decreased from 0.25

373 ± 0.09, 0.50 ± 0.16, 0.18 ± 0.06, and 0.08 ± 0.03 µg N m^{-3} to 0.05 ± 0.01, 0.24 ± 0.08, 0.05 ± 0.02,

374 and 0.03 ± 0.01 µg N m^{-3} during 2020-2100 in SSP1-2.6 scenario (Figure S13-S14 and Table S3),

375 respectively. However, the decreasing ratios of ambient HNO_3 levels especially in some developing

376 countries such as Democratic Congo (-13%) and West Africa Coast (-47%) were much less than

377 those of ambient NO_3^- levels. For the SSP3-7.0 and SSP5-8.5 scenarios, the HNO_3 levels in some

378 developing countries such as Democratic Congo (18%), West Africa Coast (16%), Indonesia (13%)

379 even experienced moderate increases. It was assumed that the government gave less investment in

380 environment improvement and the anthropogenic emission did not show marked decrease under the

381 condition of SSP3-7.0 scenario (Chen et al., 2023; Chen et al., 2020).

382 As shown in Figure S15-S18 and Table S4-S5, the higher ambient NH_3 and NH_4^+

383 concentrations also focused on India and North China. In SSP1-2.6, the ambient NH_3 (NH_4^+)

384 concentrations in China, India, Europe, and the United States decreased from 3.51 ± 1.12 (2.00 ±

385 0.62), 6.30 ± 2.12 (4.26 ± 1.42), 1.54 ± 0.51 (0.75 ± 0.24), and 1.79 ± 0.59 (0.53 ± 0.17) µg N m^{-3}

386 to 1.75 ± 0.58 (0.58 ± 0.19), 2.57 ± 0.85 (1.25 ± 0.41), 1.15 ± 0.36 (0.50 ± 0.16), and 1.58 ± 0.52

387 (0.45 ± 0.15) µg N m^{-3} during 2020-2100. Compared with SSP1-2.6, the ambient NH_3 and NH_4^+

388 concentrations in heavy-pollution scenarios (SSP3-7.0, and SSP5-8.5) scenarios did not show

389 marked decreases from 2020-2100. Some developing countries such as Argentina (9%), Democratic

390 Congo (25%), and West Africa Coast (24%) even suffered from persistent increases of ambient NH_3

391 and NH_4^+ levels. It might be associated with ineffective control of NH_3 emission compared with

392 NO_x emission.

393 3.6 Conclusions and limitations

394 The ground-level ambient reactive nitrogen observations, satellite retrievals, GEOS-Chem

395 model output, and many other geographical covariates were integrated into the multi-stage model

396 to reveal the global patterns of ambient reactive nitrogen components during 2000-2019. Then, these

397 high-resolution reactive nitrogen dataset during the historical period was then utilized as the

398 constraint to calibrate the CMIP6 dataset in four scenarios during 2020-2100. The results indicated

399 the cross-validation (CV) R^2 values of four reactive nitrogen species showed satisfied performance

400 ($R^2 > 0.55$). At the spatial scale, four reactive nitrogen components exhibited the higher

401 concentrations in China and India. For the temporal variations, the concentrations of estimated

ambient reactive nitrogen components in China experienced persistent increases during 2000-2013, while they suffered from drastic decreases since 2013 except NH_3, which might be linked with the impact of clean air policy. However, the concentrations of these species in Europe and the United States remained relatively stable since 2000. In the future scenarios, SSP3-7.0 (traditional energy scenario) and SSP1-2.6 (carbon neutrality scenario) displayed the highest and lowest reactive nitrogen component concentrations, respectively.

Global trends of four reactive nitrogen components during 2000-2100 emphasizes the urgent mitigation measures (carbon neutrality pathway) to reduce precursor emissions in order to decrease the concentrations and depositions of reactive nitrogen components especially in China and India. Furthermore, our result could give valuable insights into the impact of reactive nitrogen components on human health and ecological environment. However, this study still shows some limitations. First of all, the observation networks mainly focus on China, Europe, and the United States, and thus the simulations in many other regions might show large uncertainties. Secondly, both of the GEOS-Chem output and CMIP6 future climate scenario data also exhibits large uncertainties, which could impact the reliability of this study. Lastly, our predictions were performed on the basis of the premise that the world was steadily developing, and cannot predict the impacts of uncontrollable factors (e.g., COVID-19, Russia-Ukraine War).

Competing interests

The contact author has declared that none of the authors has any competing interests.

Data availability

The CMIP6 dataset used in this publication is available in https://esgf.nci.org.au/projects/cmip6-nci/.

Author contributions

LR and WGH designed the study; LR developed the model; GYN, ZLJ, and SYB analyzed the observation and model data. LR wrote this manuscript.

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631   **Figure 1** The workflow of global full-coverage reactive nitrogen estimates during 2000-2100.

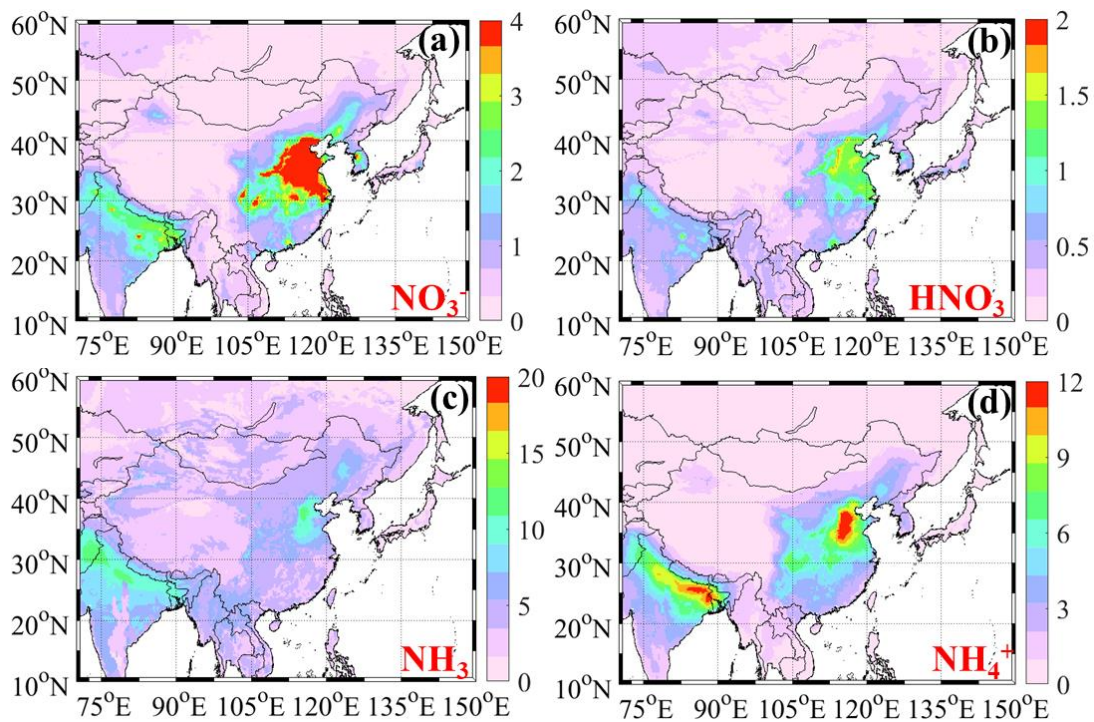632 GC_{output} denotes the GEOS-Chem output.
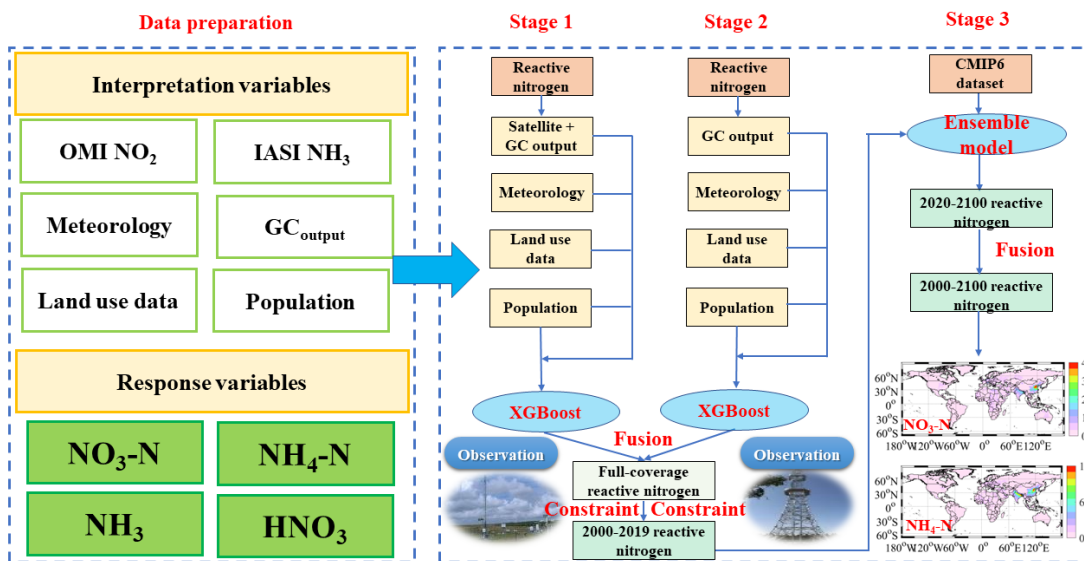
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634 **Figure 2** The predictive performances of four reactive nitrogen components including NO_3^- (a),
635 HNO_3 (b), NH_3 (c), and NH_4^+ (d). The model was constructed with 90% original data and the
636 remained data was applied to validate the model. The black solid line denotes the best-fitting curve
637 for all of the points, while the black dashed line represents the diagonal, which means the same
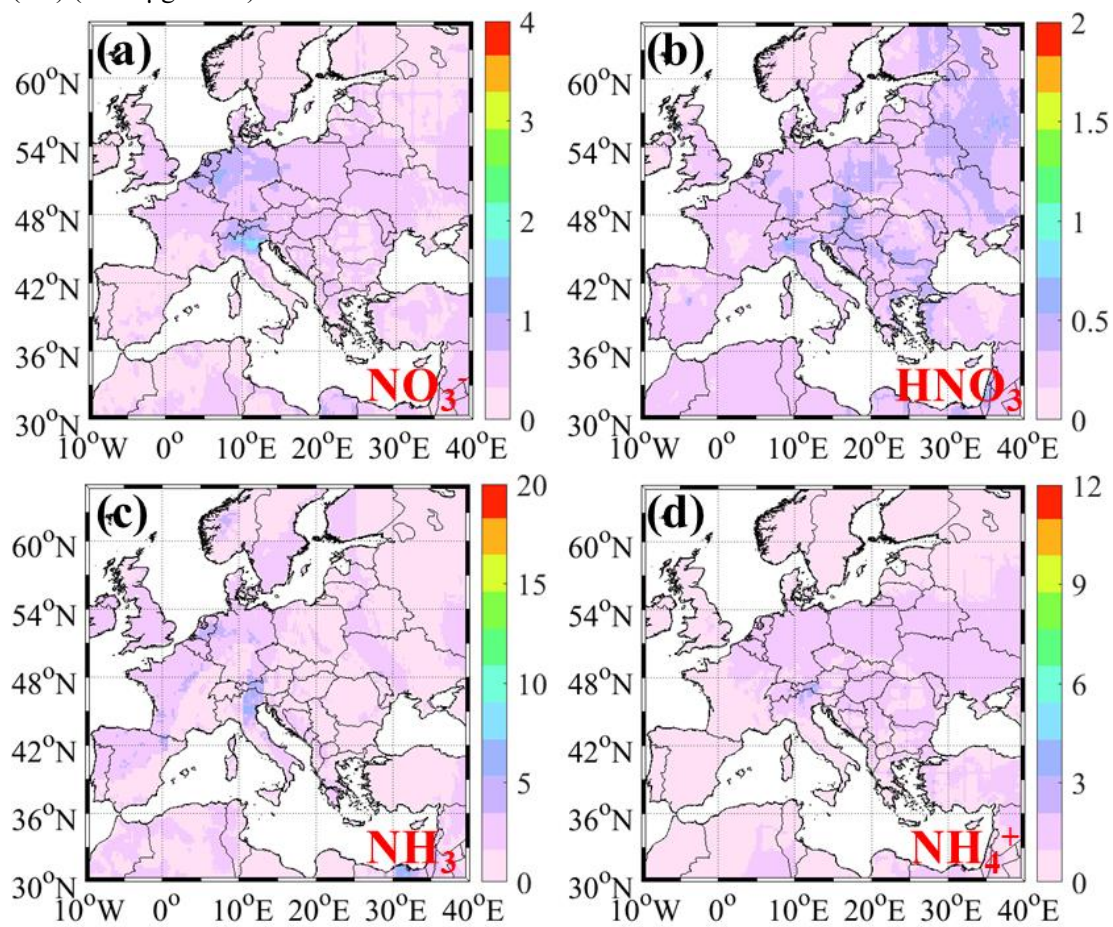638 observed and simulated values. The color scale denotes the sample size.

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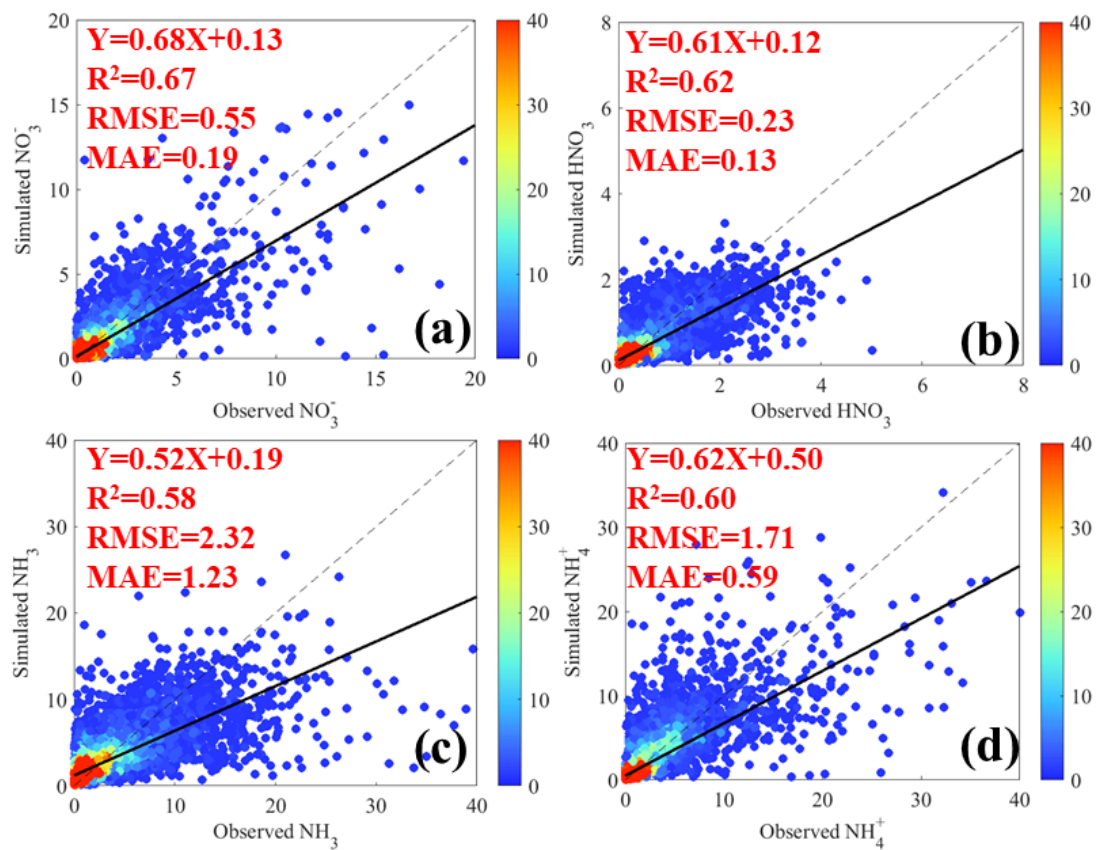
Figure 3 The spatiotemporal variations of NO_3^-, HNO_3, NH_3, and NH_4^+ concentrations in East Asia (a-d) (Unit: μg N m^{-3}).

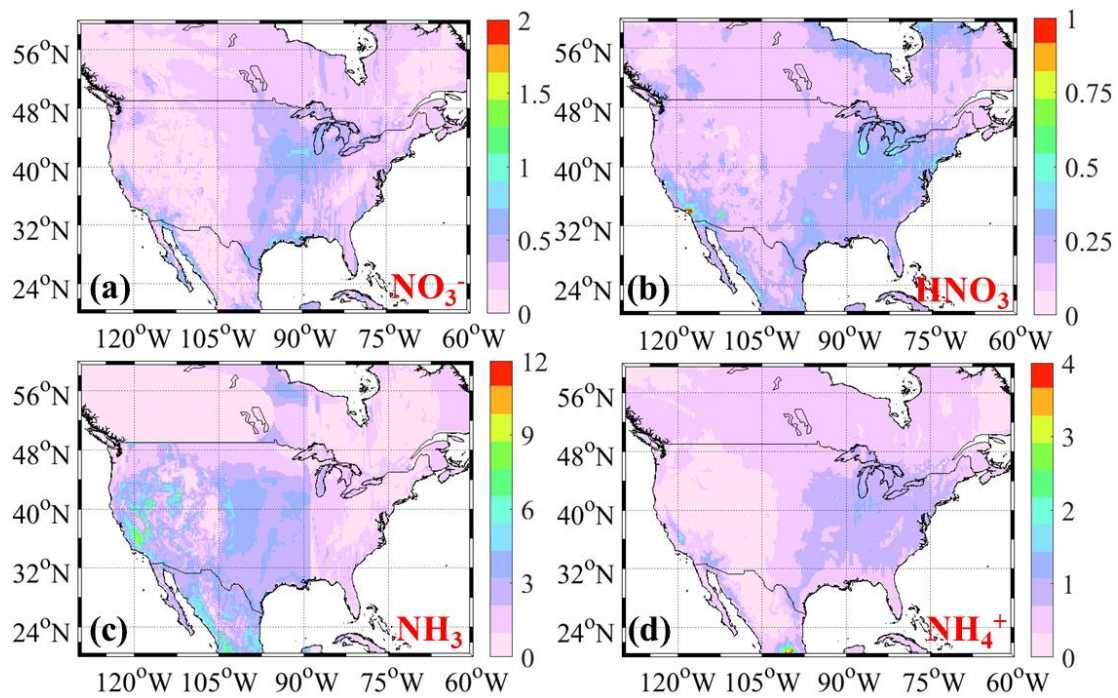644   **Figure 4** The spatiotemporal variations of NO_3^-, HNO_3, NH_3, and NH_4^+ concentrations in Europe
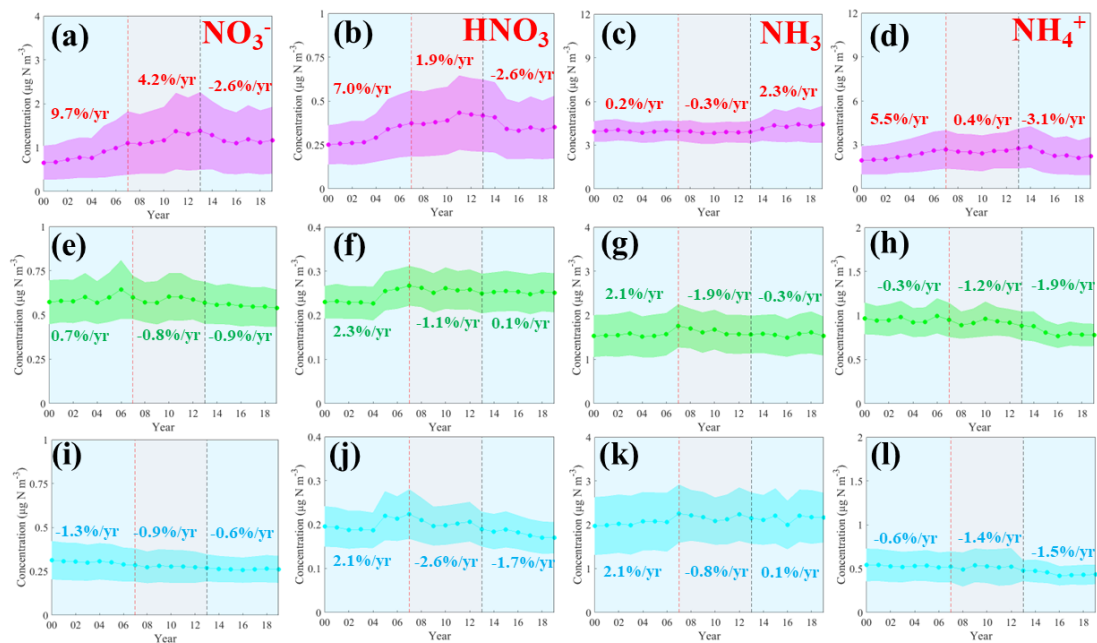
645   (a-d) (Unit: µg N m^{-3}).

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648   **Figure 5** The spatiotemporal variations of NO_3^-, HNO_3, NH_3, and NH_4^+ concentrations in North

649   America (a-d) (Unit: µg N m^{-3}).



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Figure 6 The long-term variations of NO_3^-, HNO_3, NH_3, and NH_4^+ concentrations in China (pink), Europe (green), and the United States (cyan) (Unit: μg N m^{-3}).

654   **Figure 7** Spatial variations of projected global ambient concentrations of reactive nitrogen
655   components under different climate change scenarios (Unit: μg N m^{-3}). Panels (a-b) represent the
656   annual mean concentrations of ambient NO$_3^-$ under SSP1-2.6, SSP2-4.5 during 2021-2100,
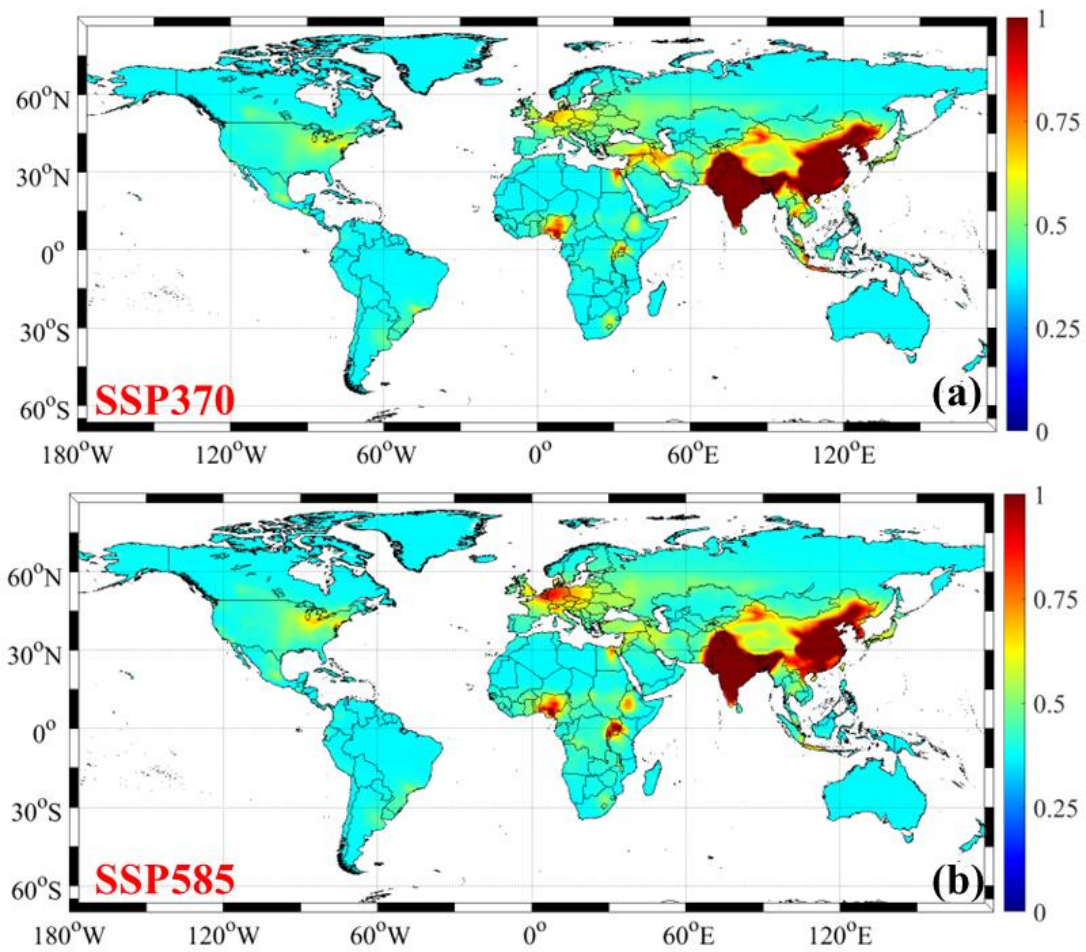657   respectively.

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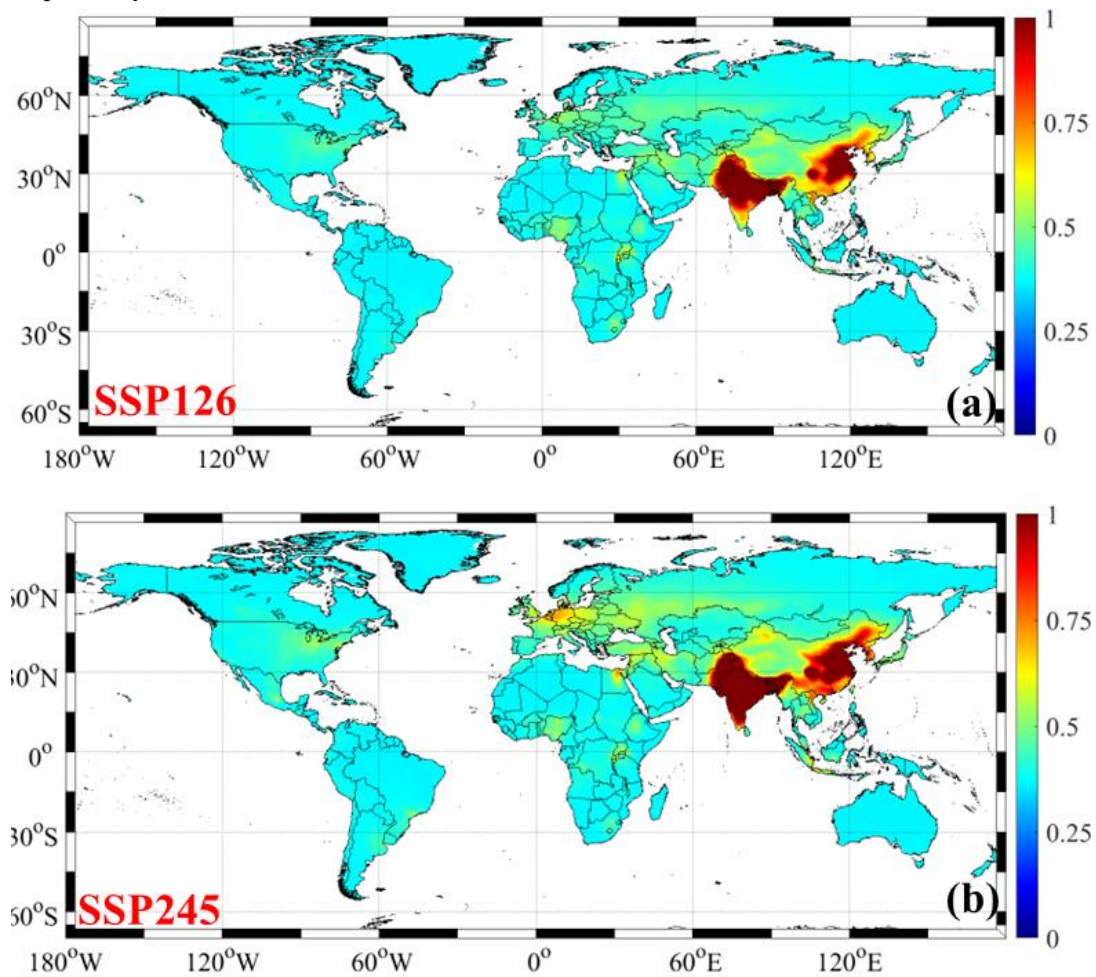
660 **Figure 8** Spatial variations of projected global ambient concentrations of reactive nitrogen
661 components under different climate change scenarios (Unit: μg N m^{-3}). Panels (a-b) represent the
662 annual mean concentrations of ambient NO$_3^-$ under SSP3-7.0, and SSP5-8.5 during 2021-2100,
663 respectively.

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666 **Table 1** The temporal variations of ambient NO_3^- (NO_3-N) concentrations (average concentrations)
667 in many countries during 2000-2100.

Scenario	NO_3^-	China	India	Europe	United States	Brazil	Argentina	Democratic Congo	West Africa Coast	Indonesia	South Korea
Historical	2000	0.66	0.95	0.57	0.31	0.29	0.21	0.56	0.30	0.31	1.06
	2005	0.91	1.26	0.60	0.30	0.31	0.22	0.56	0.30	0.34	1.38
	2010	1.17	1.53	0.60	0.28	0.33	0.23	0.58	0.31	0.32	1.43
	2013	1.39	1.63	0.57	0.27	0.30	0.24	0.58	0.31	0.33	1.57
	2015	1.16	1.34	0.56	0.26	0.32	0.22	0.58	0.32	0.45	1.88
	2019	1.18	1.46	0.54	0.26	0.32	0.23	0.61	0.32	0.37	1.87
SSP1-2.6	2020	1.16	1.23	0.41	0.27	0.25	0.18	0.55	0.28	0.34	1.81
	2040	0.82	1.12	0.28	0.10	0.21	0.14	0.62	0.32	0.28	1.46
	2060	0.69	1.01	0.15	0.05	0.18	0.13	0.43	0.25	0.24	0.69
	2080	0.41	0.89	0.12	0.05	0.17	0.12	0.32	0.17	0.20	0.39
	2100	0.33	0.65	0.10	0.06	0.11	0.08	0.21	0.11	0.16	0.23
SSP2-4.5	2020	1.19	1.43	0.44	0.24	0.26	0.19	0.52	0.29	0.37	1.85
	2040	1.09	1.35	0.43	0.16	0.22	0.16	0.57	0.31	0.35	1.80
	2060	0.89	1.22	0.35	0.11	0.20	0.15	0.51	0.27	0.32	1.25
	2080	0.63	1.06	0.29	0.07	0.17	0.12	0.42	0.23	0.23	0.68
	2100	0.41	0.95	0.24	0.05	0.14	0.10	0.36	0.19	0.20	0.38
SSP3-7.0	2020	1.25	1.59	0.53	0.33	0.31	0.22	0.64	0.34	0.42	1.95
	2040	1.36	1.50	0.47	0.24	0.26	0.19	0.61	0.33	0.47	1.89
	2060	1.18	1.35	0.42	0.19	0.22	0.16	0.56	0.30	0.41	1.56
	2080	0.96	1.15	0.36	0.16	0.18	0.13	0.54	0.29	0.35	1.35
	2100	0.75	1.08	0.33	0.12	0.15	0.11	0.51	0.27	0.30	1.24
SSP5-8.5	2020	1.21	1.50	0.53	0.28	0.28	0.20	0.57	0.31	0.37	1.91
	2040	1.28	1.42	0.49	0.23	0.25	0.18	0.60	0.32	0.39	1.85
	2060	1.05	1.30	0.47	0.24	0.20	0.15	0.55	0.29	0.35	1.44
	2080	0.86	1.10	0.44	0.25	0.15	0.11	0.50	0.27	0.29	1.26
	2100	0.58	1.02	0.31	0.20	0.13	0.09	0.46	0.25	0.25	1.05

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