

Authors' Response to Reviews of

South Asia ammonia emission inversion through assimilating

IASI observations

Ji Xia*, Yi Zhou*, Li Fang, Yingfei Qi, Dehao Li, Hong Liao, and Jianbing Jin

Atmospheric Chemistry and Physics Discussions,

RC: Reviewers' Comment, AR: Authors' Response, □ Manuscript Text

General Comments:

RC: *This paper focuses on optimizing the CEDS NH₃ emission inventory using a top-down inversion system (4D_{En}Var) over South Asia. The authors have previously conducted a similar work in China (published in ERL), but they highlight several novelties in this study:*

- 1. The application of the IASI averaging kernel to derive NH₃ column concentrations;*
- 2. A relatively high spatial resolution NH₃ emission inversion over South Asia (0.5° × 0.625°);*
- 3. The identification of a "double-peak" seasonal pattern difference between prior and posterior emissions.*

Overall, while the study's scope aligns well with the journal ACP and contributes to advancing the field by improving top-down NH₃ emissions over South Asia

Response to Referee #1: We would like to thank the referee for the careful review throughout the paper and the in-depth comments that help to improve our paper.

Specific Comments:

RC: *1) Clarify the use of observations and simulations: The use of IASI and CrIS observations, along with GEOS-Chem simulations, should be more clearly outlined. For example, a table could be included summarizing the exact periods covered by each dataset (i.e., 2015-2023, 4 months in 2019), their respective roles in the study, and the purposes they serve. Additionally, the data filtering process for removing irrational values should be more rigorously justified, particularly regarding skin temperature and cloud fraction considerations. Comparisons between IASI, CrIS, or previous studies could help validate the dataset selection.*

AR: Thanks for comment. We have integrated the relevant information into a table and added it to the manuscript. Additionally, we have added the previously omitted cloud fraction filtering

information in the manuscript, and re-calculated the IASI dataset using the skin temperature condition. Details are below:

Text in manuscript

The assimilated observations for estimating the NH₃ emissions were the monthly IASI column concentration means over the 0.5 ° × 0.625 ° GEOS-Chem grid cell. These values were derived from the latest ANNI-NH₃-v4R-ERA5 product. Despite improvements in NH₃ column retrievals from satellite observations, there remains substantial variability in measurement uncertainty, ranging from 5 % to over 1000 %. (Van Damme et al., 2014b; Whitburn et al., 2016; Van Damme et al., 2017). Data selection was performed by excluding nighttime observations, irrational values (<0), and only using data with a cloud fraction < 0.1 (Van Damme et al., 2018) and skin temperature > 263 K (Van Damme et al., 2014a) during the calculation of the monthly mean. It is important to note that the time coverage of the available version 4 IASI product used was limited: Metop-A provided data for the entire year of 2019, Metop-B provided data from January to July 2019, and Metop-C did not have data for 2019. Therefore, only the data from Metop-A and Metop-B within the 2019 time frame were used in this study.

Data	Period	Use
IASI v3	2015-2023	Annual variation of NH ₃ concentration
IASI v4	entire 2019	Inversion and Validation
Level 2 CrIS	entire 2019	Independent validation
CPCB	entire 2019	Independent validation
GEOS-Chem	entire 2019	Simulation

Table 1. The use of observations and simulations

RC: 2) *Present results in a logical sequence: The results section could benefit from a clearer structure. It might be more intuitive to first present the analysis of observed NH₃ concentrations (Sec. 3.1.1 and 3.2) before discussing the spatial and temporal patterns of NH₃ emissions. Then followed by the validation with the concentration inferred using the inversion.*

AR: Thanks for the in-depth comment. In response to your suggestion, I have revised the logical sequence of the sections in the manuscript. The following is the revised section sequence:

Text in manuscript

- 3 Results and discussion
- 3.1 Observed NH₃ concentrations
- 3.1.1 Spatial NH₃ total column concentration

3.1.2 Seasonal and annual variation of NH ₃ concentration
3.2 Spatial and Seasonal variation of NH ₃ emission
3.3 Validation
3.3.1 NH ₃ total column concentration validation
3.3.2 NH ₃ and PM _{2.5} ground concentration validation

RC: 3) Strengthen the discussion: *The discussion should delve deeper into the reasons behind the changes in the seasonal patterns observed in the posterior emissions compared to the prior. Supporting this analysis with additional figures in either the main text or supplementary materials would be valuable. For instance, including information on:*

- *Seasonal fertilizer application timings for major crops like rice, corn, and wheat;*
- *Meteorological factors affecting NH₃ volatilization and deposition, such as temperature and precipitation;*
- *Biomass burning patterns in 2019, as they may influence spatial and seasonal variations of NH₃ emissions and concentrations.*

AR: Appreciate your comment. We have added the meteorological conditions, such as precipitation and temperature, as well as the seasonal variations in biomass burning, to the supplementary materials. Additionally, the trend of anthropogenic ammonia emissions is mainly attributed to agricultural activities, so the changes in anthropology emissions generally correspond to the planting times of crops. We have identified the planting and harvesting times of crops in the South Asia region from USDA(U.S.DEPARTMENT OF AGRICULTURE, https://ipad.fas.usda.gov/rssiws/al/crop_calendar/sasia.aspx). The detailed emission analysis in revised manuscript is as follows:

Text in manuscript

3.1.2 Seasonal and annual variation of NH₃ concentration

...

The high value in May is attributed to huge amount of biomass burning in South Asia during the spring in Figure S4 (c). We have identified the planting and harvesting times of crops in the South Asia region from USDA(U.S.DEPARTMENT OF AGRICULTURE, https://ipad.fas.usda.gov/rssiws/al/crop_calendar/sasia.aspx). The heavy use of fertilizers in agricultural activities has resulted in the highest emission throughout the year, as will be illustrated in Fig. 4 (b) in Section 3.2. This has lead to the second NH₃ concentration peak in July. The reasons for higher emissions in July but lower concentration levels compared to May could be attributed to meteorological factors. The monsoon season in South Asia results in increased wet deposition, and

notably, 2019 experienced the most intense monsoon since 1994 (NASA, 2020). As shown in the Figure S4 (a) and (b), precipitation and temperature in July are the highest of the year. High temperatures increase ammonia volatilization, and the high precipitation increases the wet deposition of ammonia. These combined factors lead to July having a smaller concentration peak compared to May, despite being another peak month.

...

3.2 Spatial and Seasonal variation of NH_3 emission

...

The substantial emissions in July, as indicated by the posterior inventory, can be attributed to the increased fertilizer application for rice and corn crops during the summer season (Tanvir et al., 2019). Although biomass burning emissions are generally higher in spring in Figure S4 (c), agricultural activities remain the primary contributors to NH_3 emissions (Huang et al., 2016), resulting in July surpassing May in emission intensity. From July to September, as rice and other crops progress through their growth stages, fertilizer application typically decreases, leading to a gradual reduction in NH_3 emissions. Additionally, temperatures decline from August to September in Figure S4 (b), reducing the volatilization rate of NH_3 . This pattern occurs because NH_3 volatilization is strongly influenced by temperature (Fan et al., 2011)

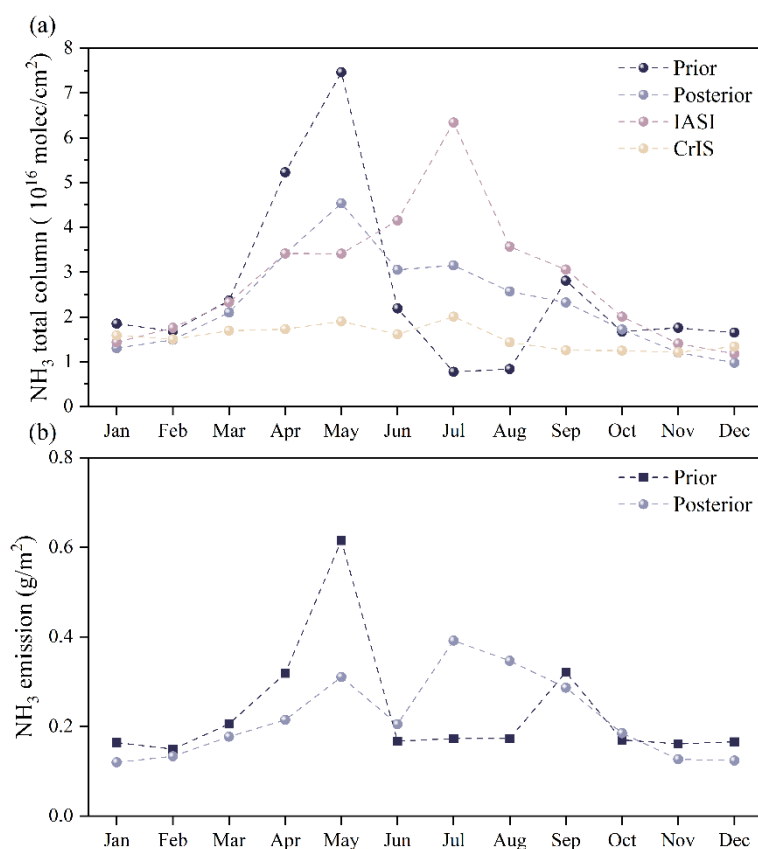


Figure 4. The monthly average total NH_3 column concentrations from the prior and posterior, IASI-observed, and CrIS-observed from January to December (a). The monthly average values of

prior and posterior emissions from January to December (b).

Supplement

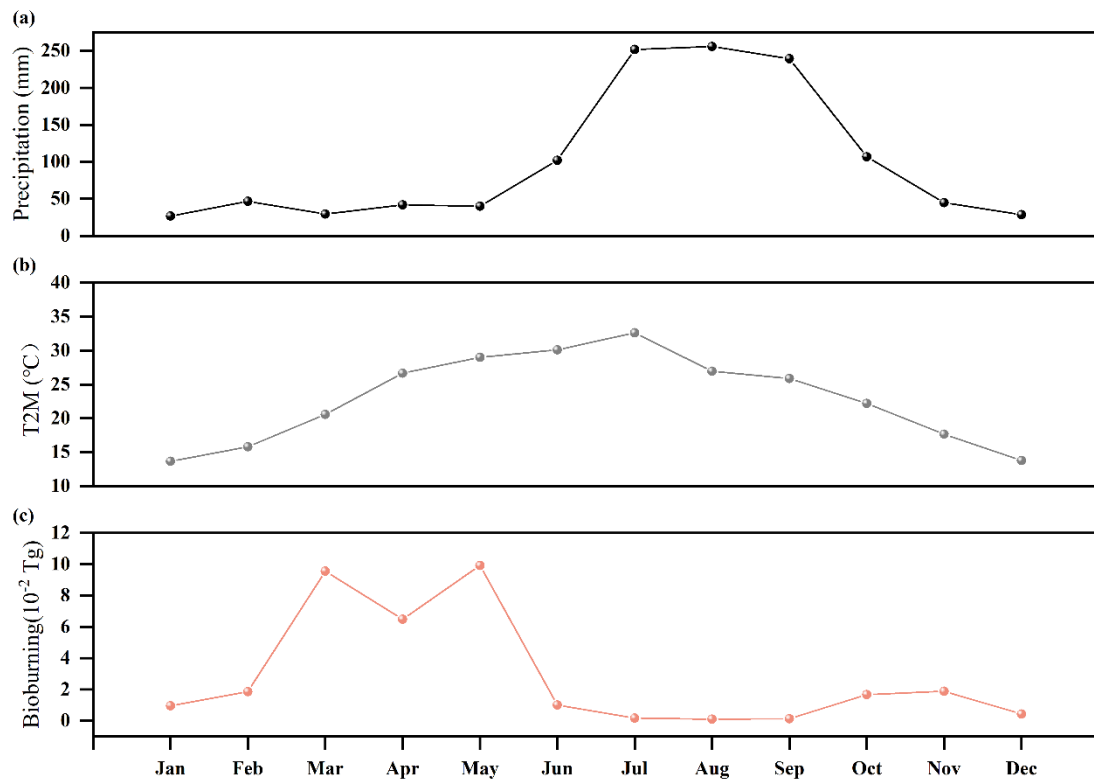


Figure S4. Monthly precipitation (a), temperature (b) from MERRA2 and biomass burning emission (c) from GFED4 in 2019.

RC: 4) Abstract: *One interesting point of this study is the different 'double-peak' seasonal pattern from the CEDS prior. I would expect to highlight this point in the abstract and briefly talk about the potential reasons for this.*

AR: Thanks for the comment. I have revised the abstract to highlight the interesting finding of the "double-peak" seasonal pattern observed in this study, which differs from the CEDS prior. Additionally, I briefly discuss the potential reasons for this discrepancy in the revised abstract. The following is the revised abstract excerpt:

Text in Abstract:

Notably, emissions there exhibit a "double-peak" seasonal profile, with the maximum in July and the secondary peak in May. This differs from the "double-peak" trend suggested by the CEDS prior inventory, which identifies the maximum in May and a second peak in September. The differences may be attributed to a more accurate representation of regional agricultural practices, such as the

timing of fertilizer application and meteorological influences like precipitation and temperature.

RC: 5) *The effect of NH₃ on climate change: inversed cause and effect, Sanderson et al., 2006 studied the effect of climate change on acid deposition. Consider reading more related and updated papers:*

Gong, Cheng, et al. "Global net climate effects of anthropogenic reactive nitrogen." *Nature* 632.8025 (2024): 557-563.

Ma, Ruoya, et al. "Data-driven estimates of fertilizer-induced soil NH₃, NO and N₂O emissions from croplands in China and their climate change impacts." *Global Change Biology* 28.3 (2022): 1008-1022.

AR: Many thanks for your feedback. I have reviewed the recommended papers and other related recent studies, and I have updated the manuscript accordingly blew:

Text in manuscript :

1 Introduction

Ammonia (NH₃), an alkaline compound, has the capacity to react with acidic gases present in the atmosphere, thereby contributing to the formation of secondary aerosols, notably ammonium sulfate and ammonium nitrate (Jimenez et al., 2009). The genesis of fine atmospheric particulate matter poses significant threats to human health (Mukherjee and Agrawal, 2017). Further, ammonia gas, along with its reaction products, plays a pivotal role in soil acidification and the eutrophication of water bodies through both dry and wet deposition (Krupa, 2003), and thereby affecting the balance of ecosystems (Asman et al., 1998) and climate change (Ma et al., 2022; Gong et al., 2024).

RC: 6) *NH₃ level over South Asia, I would expect to see the number of NH₃ concentrations or emissions here, as stated the highest in the world.*

Reply: I have updated the manuscript to include the specific NH₃ concentration data for South Asia. The specific revised excerpt is as follows.

Text in manuscript :

1 Introduction

...

With an enormous livestock population and extensive use of nitrogen fertilizers, South Asia has

experienced the highest level of atmospheric NH_3 globally (Pawar et al., 2021b; Luo et al., 2022). Specifically, the annual average ammonia concentration across India is approximately $1.8\text{--}5.6 \times 10^{16} \text{ mol/cm}^2$, while in the Indo-Gangetic Plain (IGP) of India, the concentration is double that of other regions, reaching a peak of $11.5 \times 10^{16} \text{ mol/cm}^2$ during the high season in July (Kuttippurath et al., 2020).

RC: 7) Use NH_3 instead of ammonia after the definition, which applies to the whole text.

AR: Appreciate your comment. I have made the necessary revisions and replaced "ammonia" with " NH_3 " throughout the manuscript, following the initial definition.

RC: 8) "the environmental impacts can be quantified" -> "enabling the quantification of environmental impacts"

AR: Thanks for your comment. I have revised the sentence as per your recommendation, changing "the environmental impacts can be quantified" to "enabling the quantification of environmental impacts." The specific revised excerpt is as follows:

Text in manuscript:

1 Introduction

...

Over the past decade, scientists have predominantly employed the "bottom-up" approach to estimate NH_3 emissions. When combined with chemical transport models, atmospheric NH_3 dynamics can be simulated, enabling the quantification of environmental impacts.

....

RC: 9) however, these bottom-up estimates of NH_3 emissions are generally considered uncertain (Xu et al., 2019), especially compared to other pollutants mainly derived from fossil fuel combustion" -> However, these bottom-up estimates of NH_3 emissions are generally considered as uncertain (Xu et al., 2019), particularly when compared to pollutants primarily originating from fossil fuel combustion, such as ...

AR: Thanks for comment. I have revised the sentence as recommended, changing it to: "However, these bottom-up estimates of NH_3 emissions are generally considered as uncertain (Xu et al., 2019), particularly when compared to pollutants primarily originating from fossil fuel combustion, such

as...". The specific revised excerpt is as follows:

Text in manuscript :

1 Introduction

...

Over the past decade, scientists have predominantly employed the "bottom-up" approach to estimate NH₃ emissions. When combined with chemical transport models, atmospheric NH₃ dynamics can be simulated, enabling the quantification of environmental impacts. Substantial efforts have been made to quantify the spatiotemporal distribution of NH₃ sources and develop global/regional emission inventories, such as the global NH₃ emission inventory (Bouwman et al., 1997), the anthropogenic emission inventory that includes NH₃ estimates (e.g., Community Emissions Data System, CEDS) (Hoesly et al., 2018), as well as regional NH₃ inventories focusing on South Asia (Yan et al., 2003; Yamaji et al., 2004; Liu et al., 2022). However, these bottom-up estimates of NH₃ emissions are generally considered as uncertain (Xu et al., 2019), particularly when compared to pollutants primarily originating from fossil fuel combustion such as NO₂.

RC: 10) "limitations still remain": unclear what kind of limitations here.

AR: I have clarified the statement regarding the limitations in the manuscript. The next sentence has been revised to explain more specifically that "These primarily arise from the fact that satellite observations can only measure column-integrated NH₃ concentrations, which do not directly reflect emission intensity or the three-dimensional concentration field." The specific revised excerpt is as follows:

Text in manuscript :

1 Introduction

...

The rapid advancement of satellite remote sensing technology has resulted in an expanding array of valuable NH₃ products, such as those from the first satellite NH₃ observations using the Tropospheric Emission Spectrometer (TES) (Beer et al., 2008), as well as higher-resolution retrievals from the Infrared Atmospheric Sounding Interferometer (IASI) (Pawar et al., 2021b) and the Cross-track Infrared Sounder (CrIS) (Beale et al., 2022; Kharol et al., 2022). While these remote sensing measurements play a pivotal role in characterizing atmospheric NH₃ loading, limitations still remain. These primarily arise from the fact that satellite observations can only measure column-integrated NH₃ concentrations, which do not directly reflect emission intensity or the three-dimensional concentration field. In addition to these satellite-based data, very limited ground-based

observations are publicly available over South Asia, and those that do exist are constrained by their inadequate representation of atmospheric NH₃ features (Pawar et al., 2021b).

RC: 11) Add a citation about your emission inversion system.

AR: Thanks for comment. We have added the relevant citation for our emission inversion system in the manuscript (Jin et al., 2023). The updated sentence now reads: "The NH₃ emission inventory will be calculated using our newly developed emission inversion system (Jin et al., 2023)..."

Text in manuscript :

1 Introduction

...

However, there is a paucity of studies focusing on assimilation-based NH₃ emission inversion specific to South Asia, which has some of the highest NH₃ loading hotspots compared to other continents. In this study, we aim to explore the spatial and temporal features of NH₃ emissions over South Asia. The NH₃ emission inventory will be calculated using our newly developed emission inversion system (Jin et al., 2023), by assimilating NH₃ retrievals from the IASI instruments onboard 15 MetOp-A (operational from 2008 to 2018), MetOp-B (operational since 2012), and MetOp-C (operational since 2018) satellites.

RC: 12) The description of emission inversion system and corresponding data could be shortened here and add more details in the data and method part.

AR: Thank you for your valuable comment. The description of the emission inversion system and the corresponding data has been shortened here, and more details have been added in the data and method section. The specific revised excerpt is as follows:

Text in manuscript :

1 Introduction

...

However, there is a paucity of studies focusing on assimilation-based NH₃ emission inversion specific to South Asia, which has some of the highest NH₃ loading hotspots compared to other continents. In this study, we aim to explore the spatial and temporal features of NH₃ emissions over South Asia. The NH₃ emission inventory will be calculated using our newly developed emission inversion system (Jin et al., 2023), by assimilating NH₃ retrievals from the IASI instruments

onboard MetOp-A (operational from 2008 to 2018), MetOp-B (operational since 2012), and MetOp-C (operational since 2018) satellites. Instead of directly assimilating IASI measurements as previous studies have done, we incorporated the averaging kernel information from the latest version of the IASI product. This approach allowed us to update the column concentration observations before assimilation. By doing so, we ensure a fairer comparison between the simulated and observed columnar NH_3 concentrations, a point that has been emphasized in several studies (Eskes and Boersma, 2003; von Clarmann and Glatthor, 2019), but never implemented in the IASI-based emission inversion.

RC: 13) Clearly state the aims of this study, focusing on but not only the NH_3 emission feature.

AR: Many thanks for your feedback. We have revised the relevant section of the manuscript to clearly state the aims of this study. The specific revised excerpt is as follows:

Text in manuscript :

1 Introduction

...

We aim to provide a more accurate estimation of NH_3 emission inventories and to explore their spatial and temporal characteristics across South Asia. Additionally, it serves as a model for effectively calculating atmospheric pollution emissions in regions that have been less studied in the past. The study focuses on NH_3 emissions but also contributes to a broader understanding of atmospheric pollution in under-researched regions.

RC: 14) Which periods are used for MetOp A/B/C, respectively?

AR: The latest v4 version of the IASI data, which includes averaging kernel information, has limited time coverage. Specifically, Metop-A covers the entire year of 2019, Metop-B covers from January to July 2019, and Metop-C has no data for 2019. As a result, we only used the available data from Metop-A and Metop-B within the 2019 time frame. I have made the corresponding changes in the manuscript to reflect this below:

Text in manuscript :

2.1 IASI satellite measurements

...

The assimilated observations for estimating the NH_3 emissions were the monthly IASI column

concentration means over the $0.5^\circ \times 0.625^\circ$ GEOS-Chem grid cell. These values were derived from the latest ANNI-NH₃-v4R-ERA5 product. Despite improvements in NH₃ column retrievals from satellite observations, there remains substantial variability in measurement uncertainty, ranging from 5 % to over 1000 %. (Van Damme et al., 2014; Whitburn et al., 2016; Van Damme et al., 2017). Data selection was performed by excluding nighttime observations, irrational values (< 0.1 (Van Damme et al., 2018) and skin temperature > 263 K (Van Damme et al., 2014) during the calculation of the monthly mean. It is important to note that the time coverage of the available version 4 IASI product used was limited: Metop-A provided data for the entire year of 2019, Metop-B provided data from January to July 2019, and Metop-C did not have data for 2019. Therefore, only the data from Metop-A and Metop-B within the 2019 time frame were used in this study.

RC:15) If you are using the L2 data, how do you aggregate it into the GEOS-Chem grid?

AR: The data is processed by reading daily NH₃ values and adding them to the corresponding GEOS-Chem grid cells. For each valid data point, if its latitude and longitude fall within the specified range, its NH₃ value is added to the appropriate grid cell.

RC: 16) Except for the irrational values, have you detected any large outliers?

AR: Thanks for your comment. We have not detected any significant outliers, aside from the irrational values that were excluded during the data selection process.

RC: 17) It is unclear how the B , A_z^a , and m_z coming from in the Eq. 1.

AR: Thanks for your comment. To clarify, in Eq. 1:

$$m_z = \frac{M_z^m - B_z}{M^m - B}$$

M_z^m represents the modeled concentration of NH₃ at altitude z .

B_z is the background concentration of NH₃ at the same altitude.

M^m represents the total modeled concentration of NH₃ in the atmosphere.

B is the total background concentration.

$$A_z^a = \frac{1}{N} \frac{\hat{X}^a - B}{\hat{X}^l - B}$$

\hat{X}^l represents the a priori (or assumed) concentration of NH₃ at altitude z .

B_z is again the background concentration at that altitude.

\hat{X}^a is the total a priori concentration.

N is a normalization factor, ensuring the matrix A_z^a sums correctly to account for all altitudes.

The specific revised description of the relevant part is as follows:

Text in manuscript :

2.1 IASI satellite measurements

...

Here, \hat{X}^m represents the IASI column concentration retrieved with model profile. \hat{X}^a denotes the initial IASI column concentration, with the background concentration B. The A_z values are AVK for each vertical layer, with the model profile m_z . More detailed information and the corresponding equations are provided in the supplementary materials equation S8 and S9.

Supplement

...

$$m_z = \frac{M_z^m - B_z}{M^m - B}$$

here M_z^m represents the modeled concentration of NH_3 at altitude z. B_z is the background concentration of NH_3 at the same altitude. M^m represents the total modeled concentration of NH_3 in the atmosphere. B is the total background concentration.

$$A_z^a = \frac{1}{N} \frac{\hat{X}^a - B}{\hat{X}^{lz} - B_z}$$

here \hat{X}^{lz} represents the a priori (or assumed) concentration of NH_3 at altitude z. B_z is again the background concentration at that altitude. \hat{X}^a is the total a priori concentration. N is a normalization factor, ensuring the matrix A_z^a sums correctly to account for all altitudes.

RC: 18) I do not see the 'a' in Eq.3, where has it been used?

AR: Appreciate your comment. α is a spatially varying tuning factor. It is used to compensate for the uncertainty in ammonia emissions and to adjust the spatial variability in the emission estimates. α influences the calculation of the background error covariance matrix **B** through its spatial variation, helping to optimize the ammonia emission estimates.

RC: 19) How is the parameter ' $\sigma_{\text{integrated}}$ ' calculated in Eq. 6?

AR: Thanks for comment. $\sigma_{\text{integrated}}$ is the same as σ_{total} in Equation 2. I have modified and unified the parameter names in both equations.

Text in manuscript :

2.1 IASI satellite measurements

...

$$\sigma^{\text{integrated}} = \left\{ (\sigma^{\text{instrument}})^2 + (\sigma^{\text{representing}})^2 \right\}^{0.5} \quad (2)$$

2.3 Emission inversion system

...

$$O_{i,i} = \min \left(1.0 \times 10^{16} \text{ molec cm}^{-2}, \sigma^{\text{integrated}} \right)^2 \quad (6)$$

RC: 20) Add more information such as the boundary layer condition, the spin-up process, model version...

AR: Many thanks for your feedback. I have added more detailed information, including the boundary layer condition, the spin-up process, and the model version. The manuscript fragment with more information about the model is as follows:

Text in manuscript :

GEOS-Chem, a three-dimensional (3-D) global tropospheric chemistry model, is driven by assimilated meteorological data obtained from the Goddard Earth Observing System (GEOS) at the NASA Data Assimilation Office (DAO) (Bey et al., 2001). GEOS-Chem incorporates a fully integrated chemistry system involving aerosol, ozone, NO_x, and hydrocarbons, as described by Park et al. (2004). The wet deposition scheme is explained by Liu et al. (2001) for water-soluble aerosols and by Amos et al. (2012) for gaseous components. Dry deposition is modeled using the resistance-in-series scheme proposed by Wesely and Lesht (1989), as applied by Wang and Jacob (1998). Size-specific aerosol dry deposition follows the approach outlined by Emerson et al. (2020). A nested grid simulation within the GEOS-Chem model v13.4.1 is conducted to simulate the atmospheric environment over South Asia. The nested domain (60°–98°E, 4°–40°N), shown in Fig. 2, has a horizontal resolution of 0.5° latitude by 0.625° longitude, accompanied by 47 vertical layers. The model is driven by meteorological fields from the Modern-Era Retrospective analysis for Research and Applications, Version 2 (MERRA-2) reanalysis dataset provided by the Global Modeling and Assimilation Office (GMAO) at NASA. The model employs a three-months spin-up period to minimize the influence of initial conditions. Lateral boundary conditions for the nested domain are updated every 3 hours using output from the global GEOS-Chem simulation at 2° × 2.5° resolution. Chemical initial conditions are also obtained from the global simulation to ensure consistency.

RC: 21) Sect. 2.3.1 and 2.3.2 can be combined.

AR: Thank you for your kind comment. I have merged Sections 2.3.1 and 2.3.2 into a single section: **GEOS-Chem model and emission inventory**

RC: 22) Do you run the model for the whole year 2019?

AR: Yes, we ran the model for the entire year of 2019.

RC: 23) The step that coarse-grained CEDS into 0.5×0.625 degs included in the GEOS-Chem or seperately?

AR: Thanks for comment. The step of coarse-graining the CEDS data into $0.5^\circ \times 0.625^\circ$ is done separately and is not included in the GEOS-Chem model. We directly made modifications to the original emission inventory.

RC: 24) What is the spatial resolution of the CEDS and is there any regional emission inventory with higher resolution over South Asia that you can use? Does CEDS contain both the anthropogenic and natural sectors?

AR: The spatial resolution of the CEDS emission inventory is $0.5^\circ \times 0.5^\circ$, and it only covers anthropogenic emission data. For the South Asia region, there are higher resolution emission inventories available. For example, MIXv2 is a long-term emission inventory for Asia with a spatial resolution of 0.1° , covering emission data for Southeast and South Asia.

It is important to note, however, that while higher spatial resolution in emission inventories may seem advantageous, it does not necessarily result in better inversion outcomes, as there is no direct correlation between higher resolution and improved inversion accuracy.

RC: 25) Figure 1: Increase the size of the dots and change the color regime. I suggest using different point styles for different measurements.

AR: Appreciate your comment. I have increased the size of the dots and adjusted the color regime as requested. Additionally, I have used different point styles to differentiate the various measurements in Figure 1. Considering the sequence of figures mentioned in the original text, we have also changed Figure 1 to Figure 2. The specific revised content is as follows:

Text in manuscript :

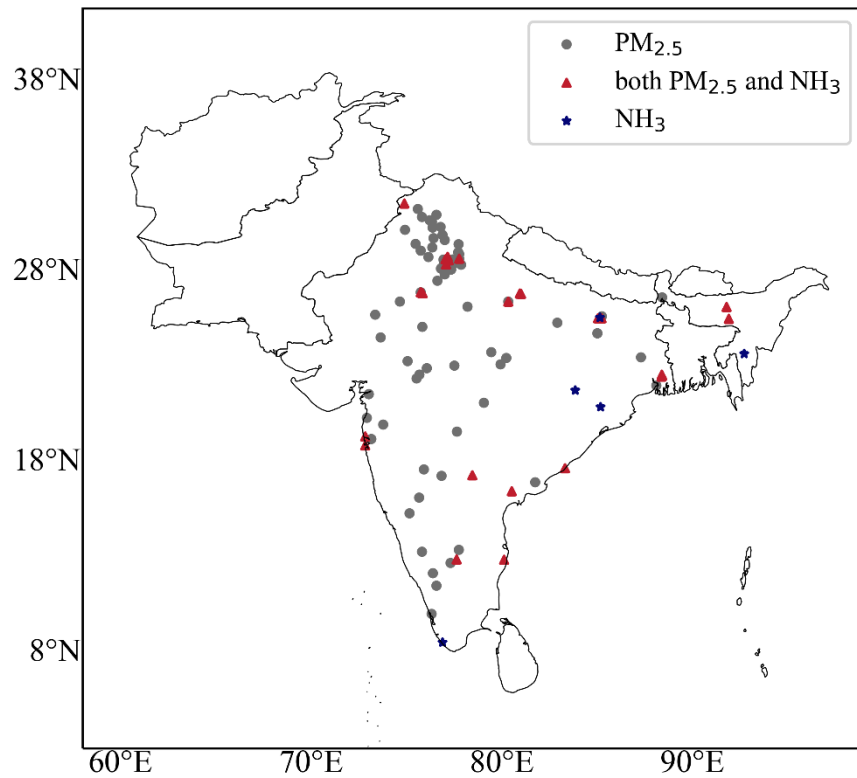


Figure 2. The GEOS-Chem model simulation domain, with dots indicating the locations of ground observation stations from the Central Pollution Control Board (CPCB), India. The three different colored dots represent stations with only $\text{PM}_{2.5}$ observations, stations with both $\text{PM}_{2.5}$ and NH_3 observations, and stations with only NH_3 observations, respectively.

RC: 26) Figure 1: Can you list these stations in the supplementary?

AR: Appreciate your comment. I have added the list of stations in the supplementary material as requested.

Table in supplement :

Station	lon	lat	StationName	City	State
AP001	80.5181667	16.5150833	Secretariat, Amaravati - APPCB	Amaravati	Andhra Pradesh
AP005	83.3	17.72	GVM Corporation, Visakhapatnam - APPCB	Visakhapatnam	Andhra Pradesh
DL001	77.152491	28.815691	Alipur, Delhi - DPCC	Delhi	Delhi
DL002	77.749675	28.725645	Anand Vihar, Delhi - DPCC	Delhi	Delhi
DL019	77.2005	28.6341	Mandir Marg, Delhi - DPCC	Delhi	Delhi
DL020	77.030469	28.68241	Mundka, Delhi - DPCC	Delhi	Delhi
DL024	80.3229863	26.4703136	Nehru Nagar, Delhi - DPCC	Delhi	Delhi
HR012	77.0667	28.4227	Sector-51, Gurugram - HSPCB	Gurugram	Haryana
KA011	77.622813	12.917348	Silk Board, Bengaluru - KSPCB	Bengaluru	Karnataka
MH008	72.82	18.91	Gole Bazar, Katni - MPCCB	Katni	Madhya Pradesh
PB001	74.876512	31.62	Golden Temple, Amritsar - PPCB	Amritsar	Punjab
RJ004	75.836858	26.902909	Adarsh Nagar, Jaipur - RSPCB	Jaipur	Rajasthan
RJ005	75.7994901	26.9164092	Police Commissionerate, Jaipur - RSPCB	Jaipur	Rajasthan
RJ006	75.730943	26.9502929	Shastri Nagar, Jaipur - RSPCB	Jaipur	Rajasthan
TN001	80.1076538	12.9099161	Alandur Bus Depot, Chennai - CPCB	Chennai	Tamil Nadu
TC006	78.451437	17.349694	Zoo Park, Hyderabad - TSPCB	Hyderabad	Telangana
UP012	80.9302753	26.8821003	Central School, Lucknow - CPCB	Lucknow	Uttar Pradesh
UP013	81.005119	26.86812	Gomti Nagar, Lucknow - UPPCB	Lucknow	Uttar Pradesh
WB007	88.3638022	22.5367507	Ballygunge, Kolkata - WBPCB	Kolkata	West Bengal
WB008	88.41002457	22.58157048	Bidhannagar, Kolkata - WBPCB	Kolkata	West Bengal
AS001	91.78063	26.181742	Railway Colony, Guwahati - APCB	Guwahati	Assam
MH012	72.8204	19.3832	Vasai West, Mumbai - MPCB	Mumbai	Maharashtra
WB011	88.380669	22.627847	Rabindra Bharati University, Kolkata - WBPCB	Kolkata	West Bengal
ML001	91.8985	25.5586	Lumpyngngad, Shillong - Meghalaya PCB	Shillong	Meghalaya
BR005	85.043586	25.586562	DRM Office Danapur, Patna - BSPCB	Patna	Bihar
BR006	85.227158	25.592539	Govt. High School Shikarpur, Patna - BSPCB	Patna	Bihar

Table S1. Stations with both PM_{2.5} data and NH₃ data

Station	lon	lat	StationName	City	State
AP002	81.7363176	16.9872867	Anand Kala Kshetram, Rajamahendravaram - APPCB	Rajamahendravaram	Andhra Pradesh
BR001	84.9994	24.7955	Collectorate, Gaya - BSPCB	Gaya	Bihar
BR003	85.2459	25.697189	Industrial Area, Hajipur - BSPCB	Hajipur	Bihar
HR001	76.778328	30.379589	Patti Mehar, Ambala - HSPCB	Ambala	Haryana
HR002	76.9254	28.6701	Arya Nagar, Bahadurgarh - HSPCB	Bahadurgarh	Haryana
HR003	77.196999	28.3419248	Nathu Colony, Ballabgarh - HSPCB	Ballabgarh	Haryana
HR004	76.141105	28.806223	ILB. Colony, Bhiwani - HSPCB	Bhiwani	Haryana
HR005	76.7997	28.2068	Municipal Corporation Office, Dharuhera - HSPCB	Dharuhera	Haryana
HR010	75.467934	29.503664	Huda Sector, Fatehabad - HSPCB	Fatehabad	Haryana
HR015	75.744941	29.14056	Urban Estate-II, Hisar - HSPCB	Hisar	Haryana
HR016	76.337619	29.307814	Police Lines, Jind - HSPCB	Jind	Haryana
HR017	76.4155	29.8006	Rishi Nagar, Kaithal - HSPCB	Kaithal	Haryana
HR018	77.0027	29.6953	Sector-12, Karnal - HSPCB	Karnal	Haryana
HR019	76.875879	29.966942	Sector-7, Kurukshetra - HSPCB	Kurukshetra	Haryana
HR020	76.9938	27.9002	General Hospital, Mandikhera - HSPCB	Mandikhera	Haryana
HR021	76.93609	28.360699	Sector-2 IMT, Manesar - HSPCB	Manesar	Haryana
HR022	75.730943	26.9502929	Shastri Nagar, Narnaul - HSPCB	Narnaul	Haryana
KA001	75.699694	16.172806	Vidayagiri, Bagalkot - KSPCB	Bagalkot	Karnataka
KA012	76.55521	11.55358	Urban, Chamarajanagar - KSPCB	Chamarajanagar	Karnataka
KA013	77.731418	13.428828	Chikkaballapur Rural, Chikkaballapur - KSPCB	Chikkaballapur	Karnataka
KA014	75.797056	13.328028	Kalyana Nagara, Chikkamagaluru - KSPCB	Chikkamagaluru	Karnataka
KL001	76.302765	10.073232	Udyogamandal, Floor - Kerala PCB	Floor	Kerala
MP002	79.446246	23.81748678	Shrivastav Colony, Damoh - MPCCB	Damoh	Madhya Pradesh
MP003	76.064118	22.9682591	Bhopal Chauraha, Dewas - MPCCB	Dewas	Madhya Pradesh
MP004	78.193251	26.203442	City Center, Gwalior - MPCCB	Gwalior	Madhya Pradesh
MP006	75.5213	22.431	Chhoti Gwaltoli, Indore - MPCCB	Indore	Madhya Pradesh
MP007	79.932247	23.168606	Marhatal, Jabalpur - MPCCB	Jabalpur	Madhya Pradesh
MP008	80.23284	23.50016	Gole Bazar, Katni - MPCCB	Katni	Madhya Pradesh
MH002	77.6345232	19.645324	Chandrapur, Chandrapur - MPCB	Chandrapur	Maharashtra
MH004	73.142019	19.25292	Khadakpada, Kalyan - MPCB	Kalyan	Maharashtra
PB002	74.907758	30.233011	Hardev Nagar, Bathinda - PPCB	Bathinda	Punjab
PB003	75.578914	31.321907	Civil Line, Jalandhar - PPCB	Jalandhar	Punjab
PB004	76.209694	30.736056	Kalal Majra, Khanna - PPCB	Khanna	Punjab
PB005	75.8086	30.9028	Punjab Agricultural University, Ludhiana - PPCB	Ludhiana	Punjab
PB006	76.331442	30.649961	RIMT University, Mandi Gobindgarh - PPCB	Gobindgarh	Punjab
PB007	76.366642	30.349388	Model Town, Patiala - PPCB	Patiala	Punjab
PB008	76.5623046	31.0325454	Ratanpura, Rupnagar - Ambuja Cements	Rupnagar	Punjab
RJ001	76.611536	27.554793	Moti Doongri, Alwar - RSPCB	Alwar	Rajasthan
RJ002	74.646594	26.470859	Civil Lines, Ajmer - RSPCB	Ajmer	Rajasthan
RJ003	76.862296	28.194909	RIICO Ind. Area III, Bhiwadi - RSPCB	Bhiwadi	Rajasthan
RJ007	84.9994	24.7955	Collectorate, Jodhpur - RSPCB	Jodhpur	Rajasthan
RJ008	75.821256	25.14389	Shrinath Puram, Kota - RSPCB	Kota	Rajasthan
RJ009	73.340227	25.771061	Indira Colony Vistar, Pali - RSPCB	Pali	Rajasthan
RJ010	73.6321397	24.5886166	Ashok Nagar, Udaipur - RSPCB	Udaipur	Rajasthan
UP003	77.849831	28.406963	Yamunapuram, Bulandshahr - UPPCB	Bulandshahr	Uttar Pradesh
UP006	77.453839	28.685382	Sanjay Nagar, Ghaziabad - UPPCB	Ghaziabad	Uttar Pradesh
UP008	77.482	28.47272	Knowledge Park - III, Greater Noida - UPPCB	Greater Noida	Uttar Pradesh
UP010	77.749675	28.725645	Anand Vihar, Hapur - UPPCB	Hapur	Uttar Pradesh
UP011	80.3229863	26.4703136	Nehru Nagar, Kanpur - UPPCB	Kanpur	Uttar Pradesh
WB002	87.2892225	23.5404352	Sidhu Kanhu Indoor Stadium, Durgapur - WBPCB	Durgapur	West Bengal
WB003	88.109737	22.06047	Haldia, Haldia - WBPCB	Haldia	West Bengal
GJ002	73.010555	21.613267	GIDC, Ankleshwar - GPCB	Ankleshwar	Gujarat
GJ004	73.010555	21.613267	GIDC, Nandesari - Nandesari Ind. Association	Nandesari	Gujarat
GJ005	72.918013	20.362421	Phase-I GIDC, Vapi - GPCB	Vapi	Gujarat
HR023	77.3320667	28.1485564	Shyam Nagar, Palwal - HSPCB	Palwal	Haryana
MH015	79.0517531	21.152875	Opp GPO Civil Lines, Nagpur - MPCB	Nagpur	Maharashtra
MH016	73.7762427	20.0073285	Gangapur Road, Nashik - MPCB	Nashik	Maharashtra
MH021	75.9063906	17.6599188	Solapur, Solapur - MPCB	Solapur	Maharashtra
UP018	77.7622941	28.9535882	Jai Bhim Nagar, Meerut - UPPCB	Meerut	Uttar Pradesh
UP019	77.709723	29.06351	Pallavpuram Phase 2, Meerut - UPPCB	Meerut	Uttar Pradesh
UP021	77.7194031	29.4723508	New Mandi, Muzaffarnagar - UPPCB	Muzaffarnagar	Uttar Pradesh
UP022	77.3231257	28.5447608	Sector - 125, Noida - UPPCB	Noida	Uttar Pradesh
KA015	75.140726	15.351773	Deshpande Nagar, Hubballi - KSPCB	Hubballi	Karnataka
WB014	88.412668	26.6883049	Ward-32 Bapupara, Siliguri - WBPCB	Siliguri	West Bengal
KA017	76.37376	12.21041	Hebbal 1st Stage, Mysuru - KSPCB	Mysuru	Karnataka
KA018	77.298051	12.733409	Vijay Nagar, Ramanagara - KSPCB	Ramanagara	Karnataka
MP010	77.511428	23.10844	Sector-D Industrial Area, Mandideep - MPCCB	Mandideep	Madhya Pradesh
MP011	75.675238	22.624758	Sector-2 Industrial Area, Pithampur - MPCCB	Pithampur	Madhya Pradesh
MP012	75.045981	23.331731	Shastri Nagar, Ratlam - IPCA Lab	Ratlam	Madhya Pradesh
UP025	77.393848	28.56923	Sector-116, Noida - UPPCB	Noida	Uttar Pradesh
UP026	82.9083074	25.3505986	Ardhali Bazar, Varanasi - UPPCB	Varanasi	Uttar Pradesh
KA016	76.822628	17.321993	Lal Bahadur Shastri Nagar, Kalaburagi - KSPCB	Kalaburagi	Karnataka

Table S2. Stations with PM_{2.5} data but without NH₃ data

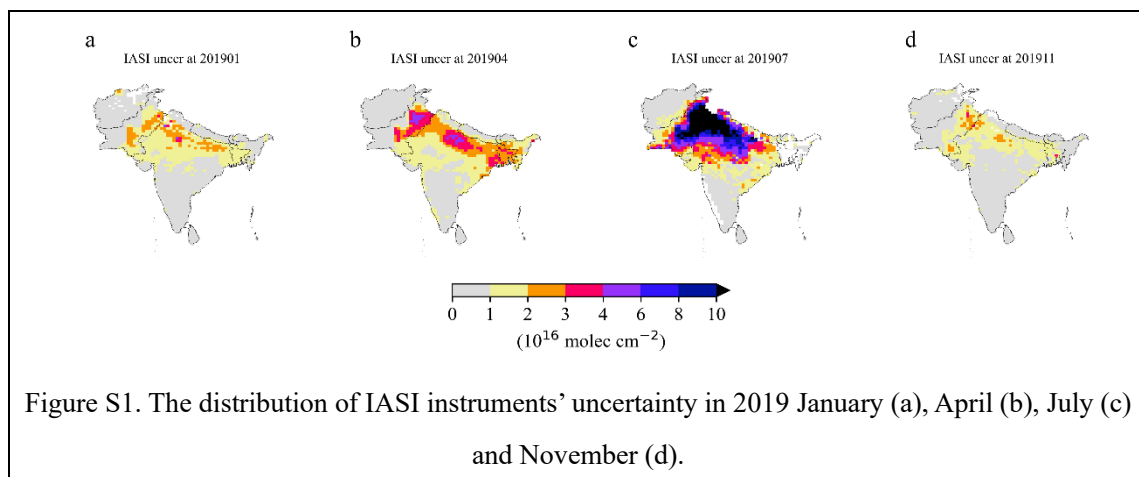
Station	lon	lat	StationName	City	State
BR008	85.147382	25.619651	Muradpur, Patna - BSPCB	Patna	Bihar
KL007	76.8865	8.5637	Kariavattom, Thiruvananthapuram - Kerala PCB	Thiruvananthapuram	Kerala
MZ001	92.7192841	23.7176342	Sikulpuikawn, Aizawl - Mizoram PCB	Aizawl	Mizoram
OD001	83.8396977	21.8004996	GM Office, Brajrajnagar - OSPCB	Brajrajnagar	Odisha
OD002	85.1707021	20.9360711	Talcher Coalfields, Talcher - OSPCB	Talcher	Odisha

Table S3. Stations with NH₃ data but without PM_{2.5} data

RC: 27) Figure 2: Row (b) seems useless and not mentioned in the text, consider moving to the supplementary.

AR: Thanks for comment. I have moved row (b) of Figure 2 to the supplementary material, as it is not discussed in the main text.

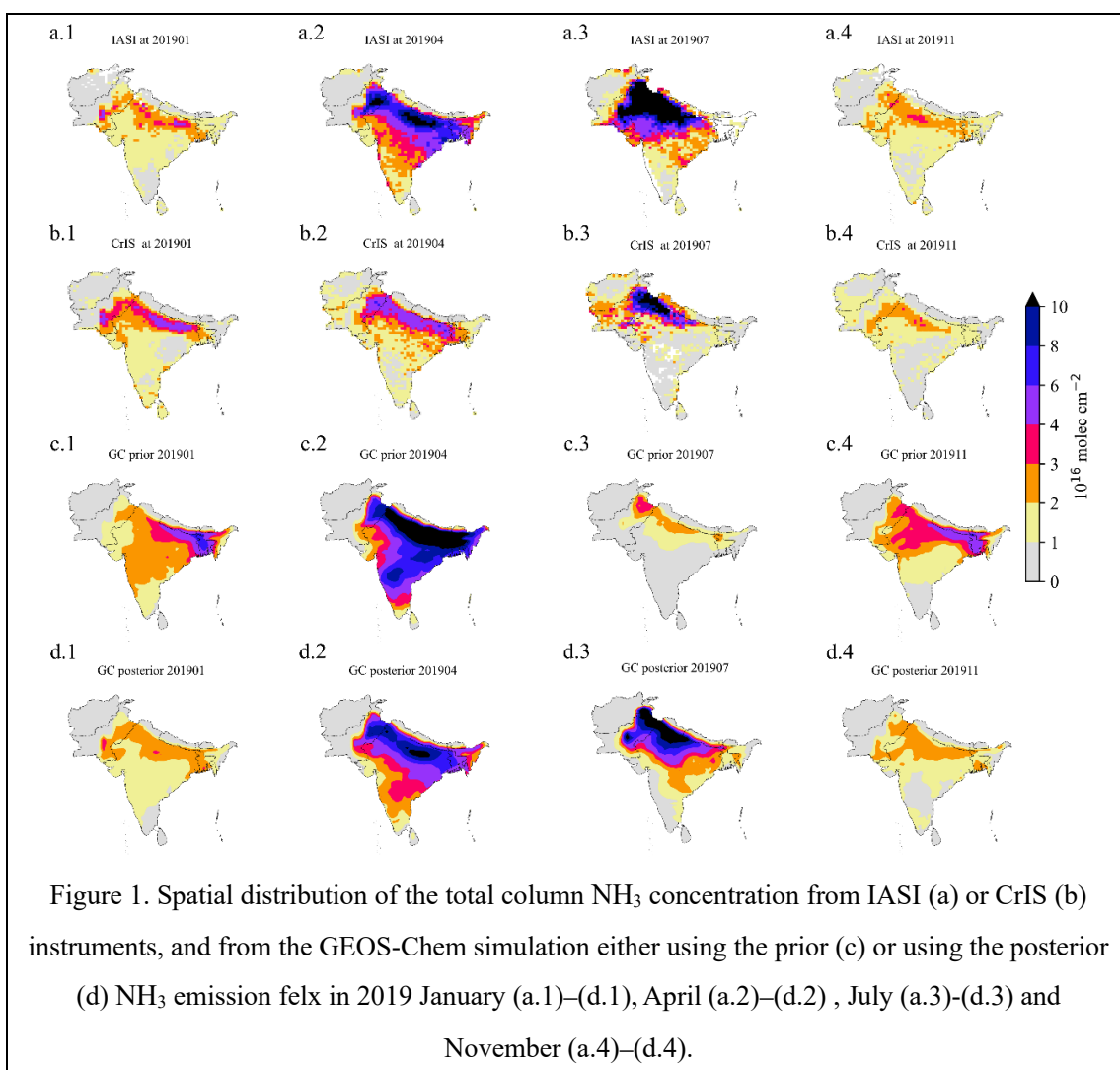
Figure in supplement:



RC: 28) Figure 2: The way you plotted the IASI and GC seems different.

AR: Many thanks for your feedback. Yes, we use scatter plots for the IASI data, while the model simulations are presented using filled contour maps to show their respective spatial distributions.

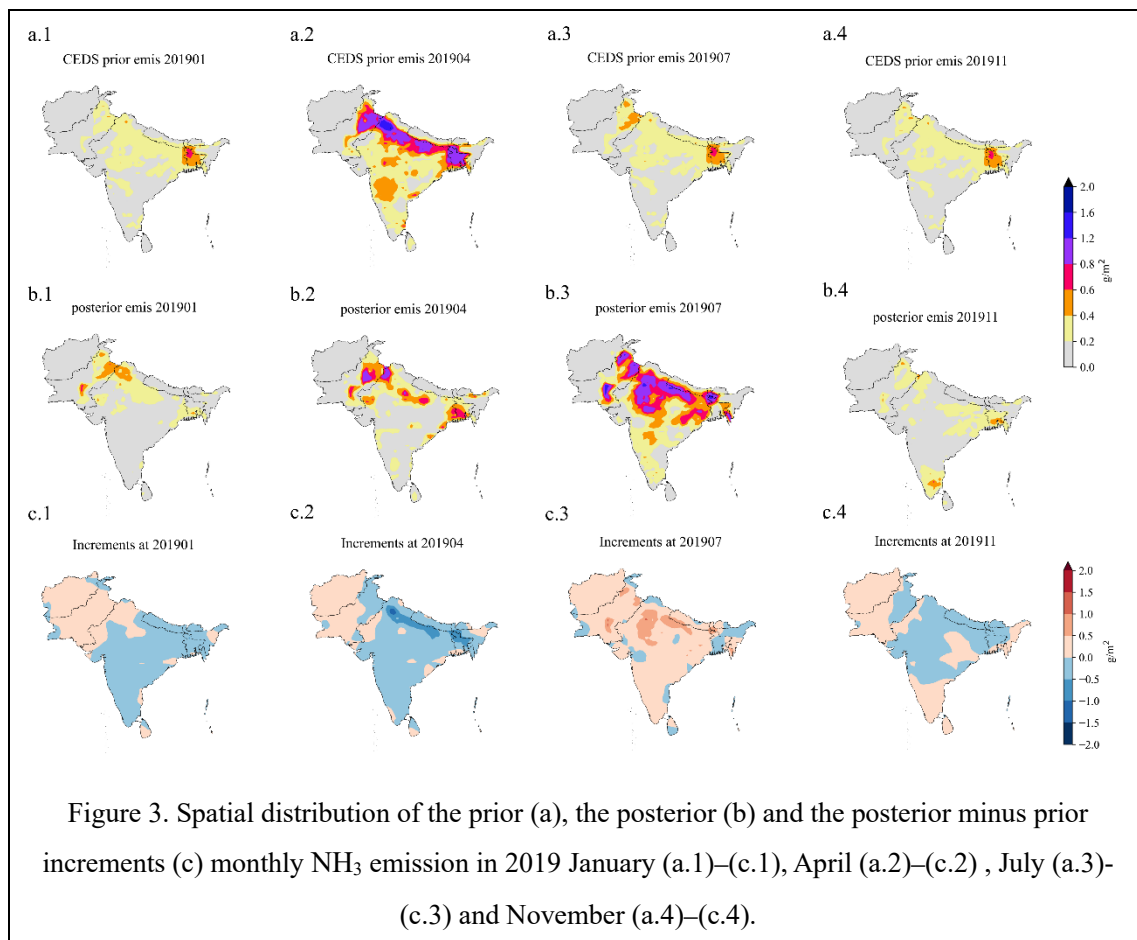
Figure in manuscript:



RC: 29) Figure 3: I am curious why there are 0 values (grey area) in the CEDS prior and posterior, and how you deal with this 0 values or missing gap in the posterior. It may look confused since there are lots of grey area in (a) and (b) but not shown in (c)

AR: Thanks for comment. Apologies for the confusion. The values in the grey area are not zero, but rather fall within the range of 0 to 0.2. They are indeed non-zero, but the values are quite small. We have treated these small values as valid and have not excluded them in the posterior, which may explain why they appear differently in panel (c).

Text in manuscript:



RC: 30) NH_3 total column concentration: "Four months": in 2019. Note all month are presented in Fig.4.

AR: Apologies for the confusion, we have revised this statement in the original text. The specific revised fragment is as follows:

Text in manuscript:

3.1.1 Spatial NH_3 total column concentration

The prior and posterior snapshot of NH_3 column concentration simulations for four months (January, April, July, and November) are presented in Fig. 1 (c)-(d), alongside the IASI measurements shown in panel (a).

RC: 31) "Distributions for the rest months": could you provide timeseries or seasonal pattern of these two satellite observations?

AR: Thanks for comment. The time series for the two satellites are presented in panel (a) of Figure

4, while the spatial distribution is shown in Figure S3.

Figure in manuscript and supplement :

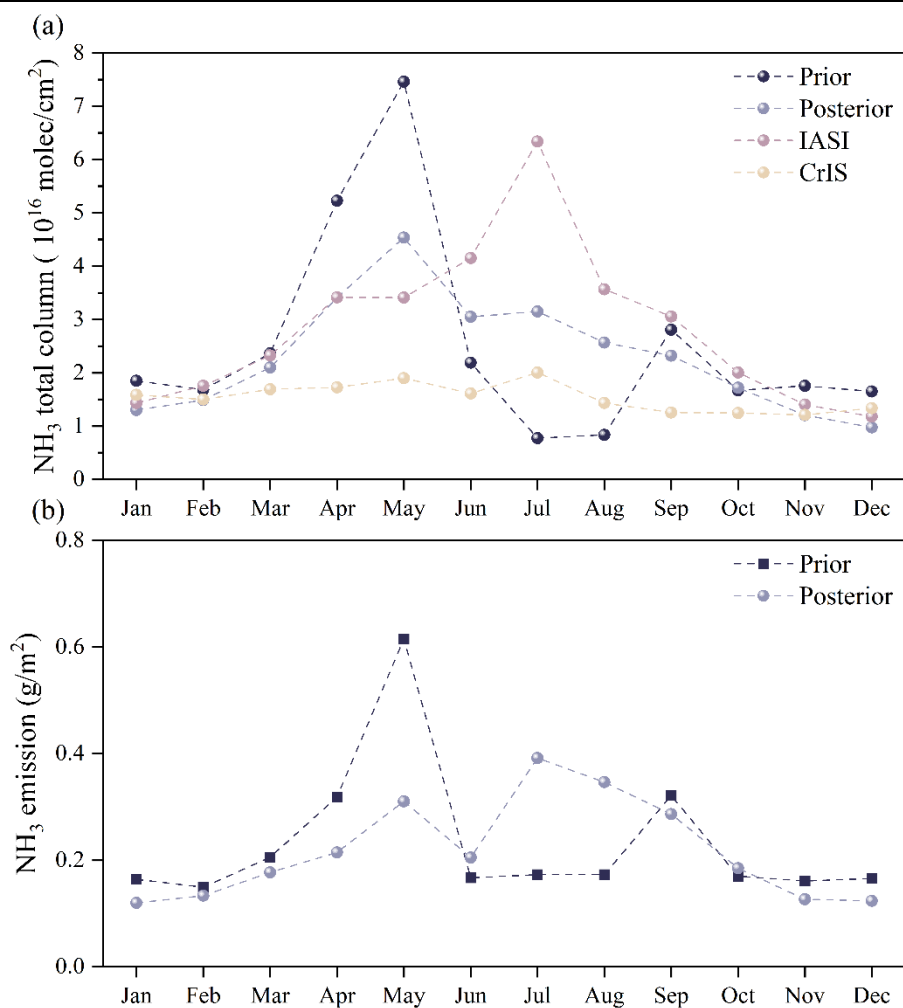
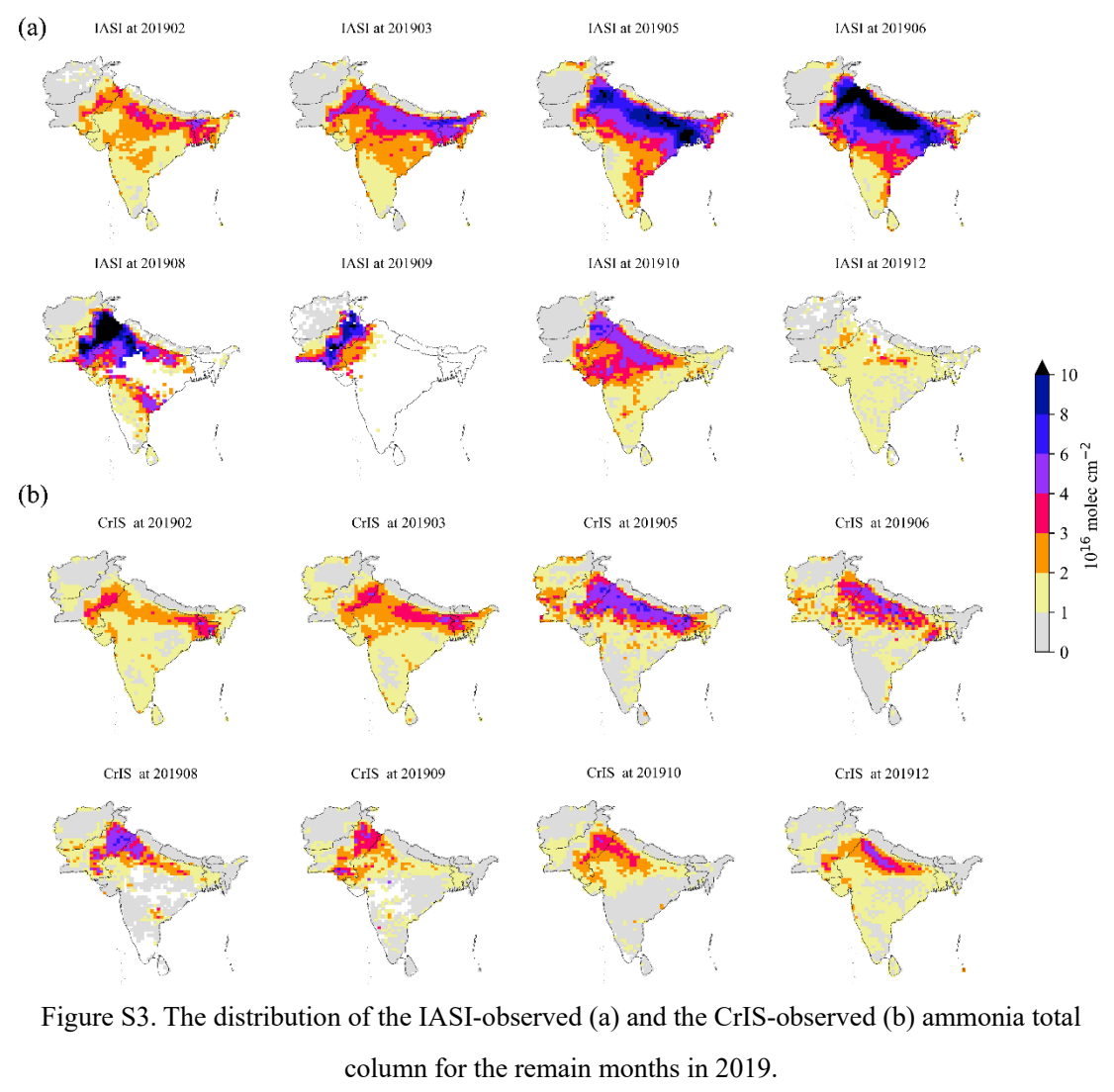


Figure 4. The monthly average total NH_3 column concentrations from the prior and posterior IASI-observed, and CrIS-observed from January to December (a). The monthly average values of prior and posterior emissions from January to December (b).



RC: 32) "background error covariance matrix": does this refer to B in Eq.3 and 5?

AR: Yes, the "background error covariance matrix" refers to B in Equations 3 and 5.

Text in manuscript :

2.3 Emission inversion system

...

$$\mathcal{J}(f) = \frac{1}{2} (f - f_b)^T B^{-1} (f - f_b) + \frac{1}{2} \{y - \mathcal{H}\mathcal{M}(f)\}^T O^{-1} \{y - \mathcal{H}\mathcal{M}(f)\} \quad (3)$$

$$B(i, j) = \sigma^2 \cdot f(i) \cdot f(j) \cdot C(i, j) \quad (5)$$

3.1.1 Spatial NH_3 total column concentration

...

Note that there are still some discrepancies in the posterior simulation vs IASI column measurements. In particular, as shown in panel a.3 vs. d.3 of Fig. 1, the posterior simulation did not fully reproduce the extremely high NH_3 loading observed by IASI in July (with column-integrated concentrations exceeding $10 \times 10^{16} \text{ molec cm}^{-2}$). This occurs because the goal of the assimilation is to achieve the best fit between the posterior, the observations, and the prior emissions, rather than just fitting the observations alone. The extremely high NH_3 concentrations are less likely given the relatively low prior NH_3 emissions and the background error covariance matrix described in Section 2.3.

RC: 33) NH_3 and $\text{PM}_{2.5}$ ground concentration validation: "The mismatch": specify which kind of mismatch, bias mismatch or spatial/temporal

AR: Thanks for comment. The mismatch mentioned refers to both bias and spatial mismatches. The bias mismatch is observed in the differences between the ground observations and simulations, where ground NH_3 concentrations were occasionally higher than the simulations. The spatial mismatch could be due to the locations of the monitoring stations, which are often in urban areas where NH_3 levels are higher due to traffic and human activities.

RC: 34) Can also add the Bias in Fig. 4, 5 and 7.

AR: Many thanks for your feedback. I have added the bias to Figures 4, 5, and 7 as requested.

RC: 35) Fig. 4 and 5 can be combined.

AR: Thanks for comment. I have combined Figures 4 and 5 as per your recommendation. The revised figure along with RC (34) is as follows:

Figure in manuscript:

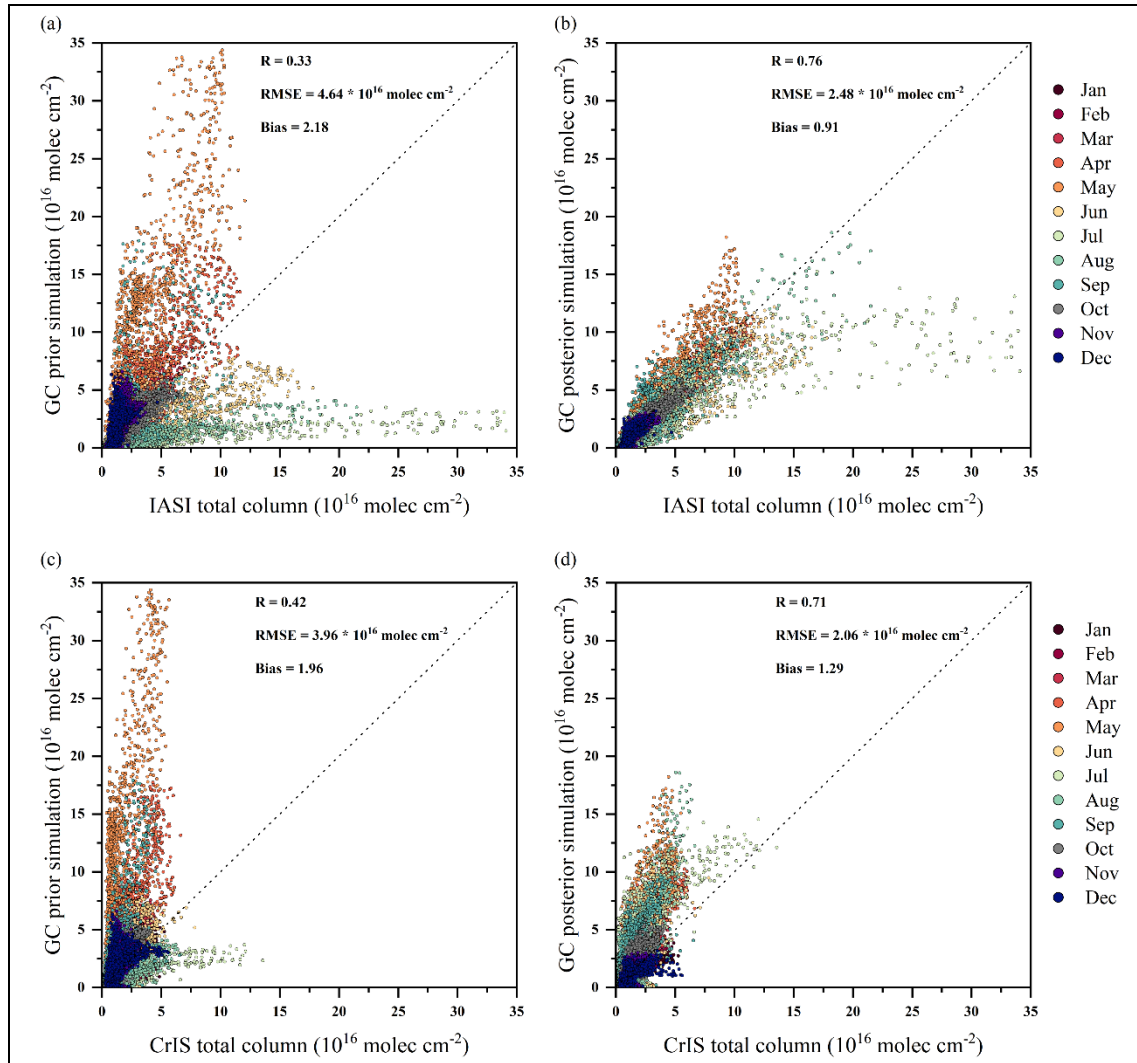


Figure 6. Scatter plot of the IASI (a-b) and CrIS (c-d) observed NH_3 concentrations against the NH_3 simulation over South Asia, either using the prior or the posterior NH_3 emission inventory, from January to December.

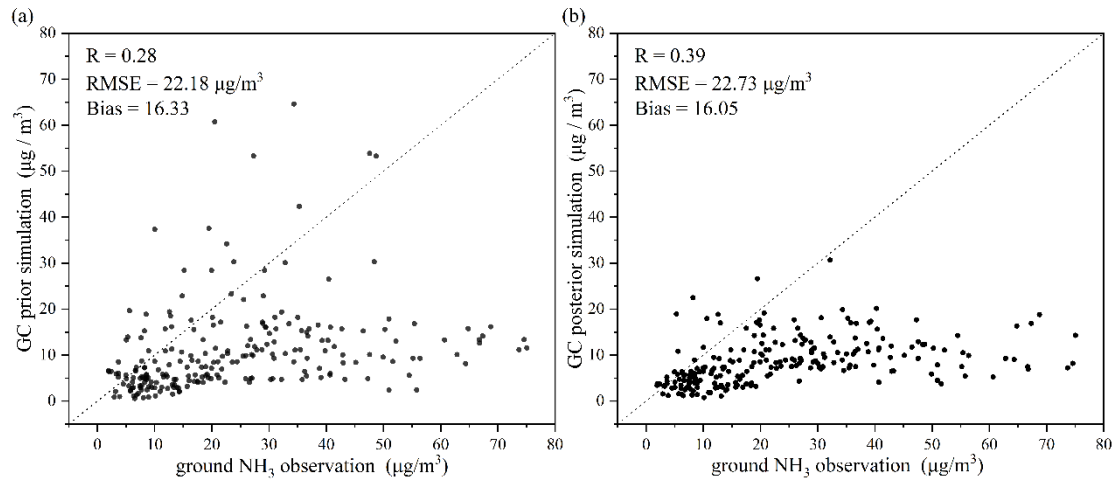
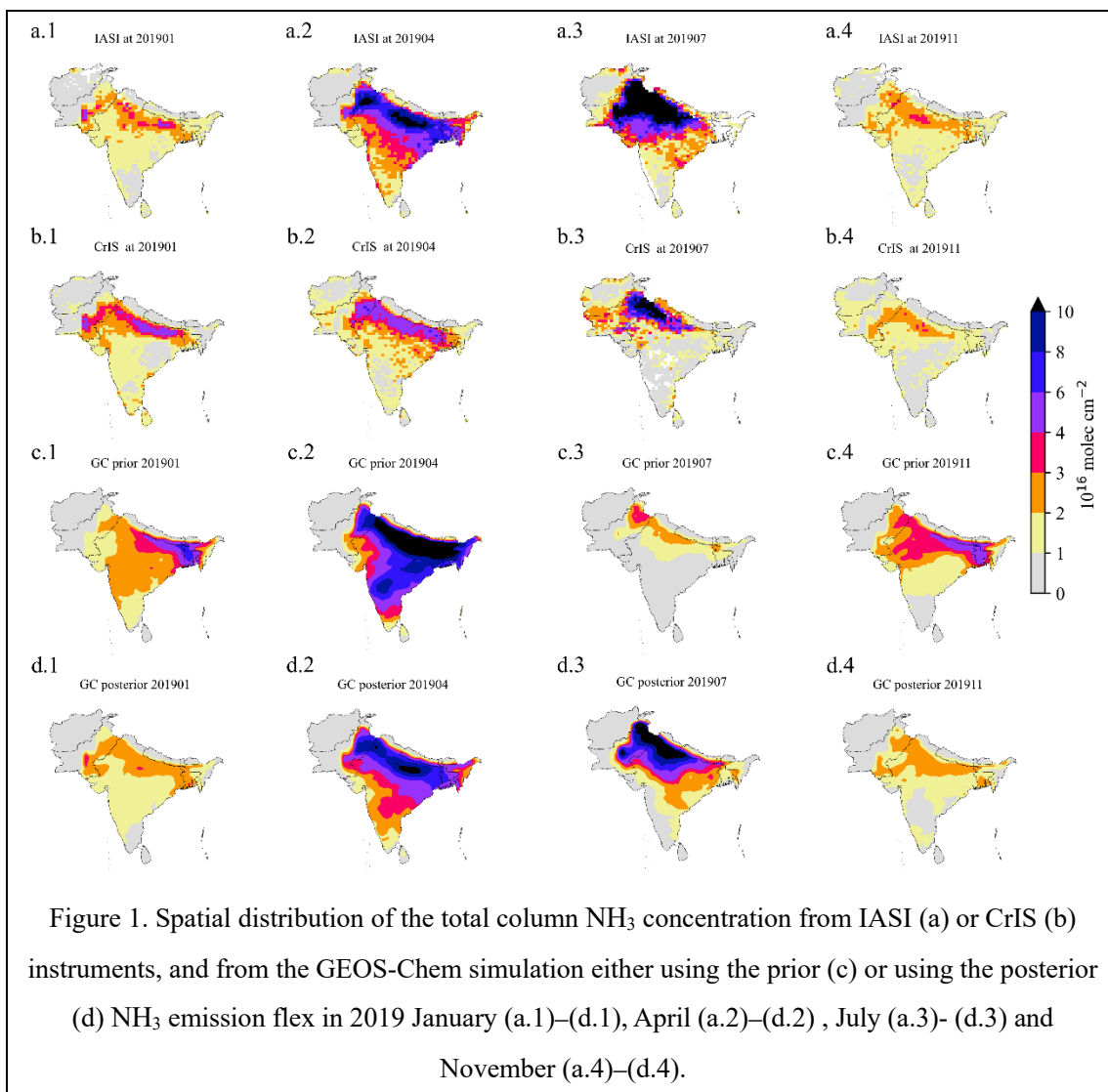


Figure 7. Scatter plot of the ground-observed against the NH_3 simulation over South Asia either using the prior (a) or using the posterior (b) NH_3 emission inventory in 2019.

RC: 36) Fig. 2 and 6 can be combined.

AR: Thanks for comment. I have combined Figures 2 and 6 as per your recommendation. The revised figure is as follows:

Figure in manuscript:



RC: 37) "systematic biases": how do you define the systematic biases, which differentiate from the random error?

AR: Thanks for comment. Systematic bias is a consistent and predictable deviation, usually caused by factors such as biases in input data (e.g., emission inventories). This type of error typically leads to the model consistently overestimating or underestimating the actual values in its predictions, whereas random errors are unpredictable and irregular, occurring due to factors that are not consistent across different measurements or observations.

RC: 38) Seasonal and annual variation of ammonia concentration: The peaks in NH_3 posterior concentration (May) and emission (July) are different and interesting, can you elaborate to explain it?

Reply: Thank you for your kind comment. Although the posterior emission shows that July has the highest emissions, this emission is mainly due to human activities. Specifically, biomass combustion emissions are highest in spring, especially in May in Figure 4 (c), and anthropology emissions are also one of the peaks. Although anthropology emissions are highest in July, meteorological factors must also be considered. As shown in the Figure 4 (a) and (b), precipitation and temperature in July are the highest of the year. High temperatures increase ammonia volatilization, and the high precipitation increases the wet deposition of ammonia. These combined factors lead to July having a smaller concentration peak compared to May, despite being another peak month. The revised related discussion in the text is as follows:

Text in manuscript :

3.1.2 Seasonal and annual variation of NH_3 concentration

...

The high value in May is attributed to huge amount of biomass burning in South Asia during the spring in Figure S4 (c). We have identified the planting and harvesting times of crops in the South Asia region from USDA(U.S.DEPARTMENT OF AGRICULTURE, https://ipad.fas.usda.gov/rssiws/al/crop_calendar/sasia.aspx). The heavy use of fertilizers in agricultural activities has resulted in the highest emission throughout the year, as will be illustrated in Fig. 4 (b) in Section 3.2. This has lead to the second NH_3 concentration peak in July. The reasons for higher emissions in July but lower concentration levels compared to May could be attributed to meteorological factors. The monsoon season in South Asia results in increased wet deposition, and notably, 2019 experienced the most intense monsoon since 1994 (NASA, 2020). As shown in the Figure S4 (a) and (b), precipitation and temperature in July are the highest of the year. High temperatures increase ammonia volatilization, and the high precipitation increases the wet deposition of ammonia. These combined factors lead to July having a smaller concentration peak compared to May, despite being another peak month.

...

3.2 Spatial and Seasonal variation of NH_3 emission

...

The substantial emissions in July, as indicated by the posterior inventory, can be attributed to the increased fertilizer application for rice and corn crops during the summer season (Tanvir et al., 2019). Although biomass burning emissions are generally higher in spring in Figure S4 (c), agricultural activities remain the primary contributors to NH_3 emissions (Huang et al., 2016),

resulting in July surpassing May in emission intensity. From July to September, as rice and other crops progress through their growth stages, fertilizer application typically decreases, leading to a gradual reduction in NH_3 emissions. Additionally, temperatures decline from August to September in Figure S4 (b), reducing the volatilization rate of NH_3 . This pattern occurs because NH_3 volatilization is strongly influenced by temperature (Fan et al., 2011)

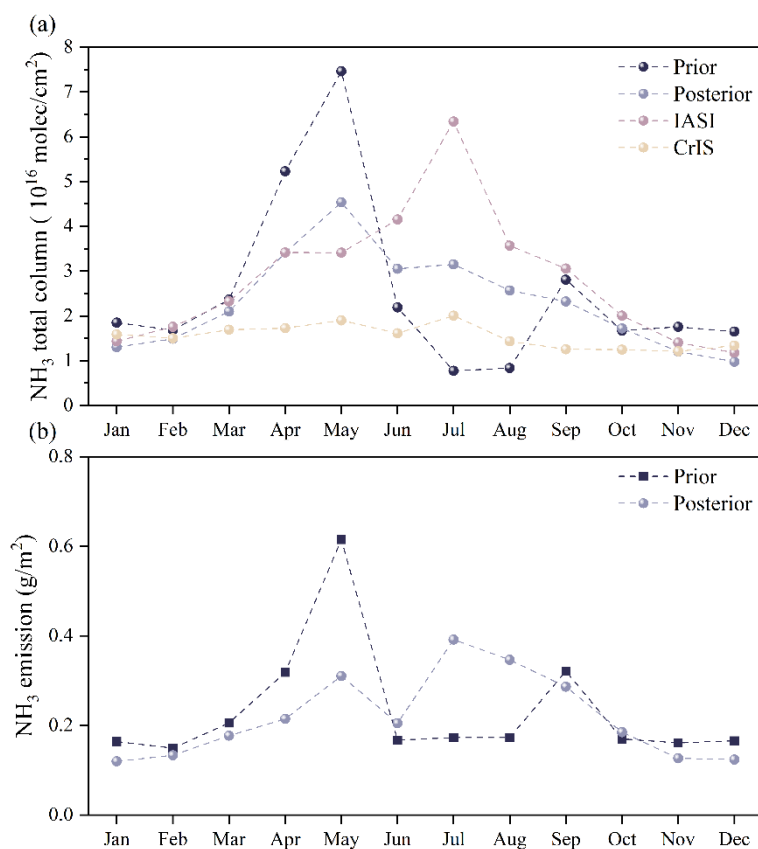


Figure 4. The monthly average total NH_3 column concentrations from the prior and posterior, IASI-observed, and CrIS-observed from January to December (a). The monthly average values of prior and posterior emissions from January to December (b).

Supplement

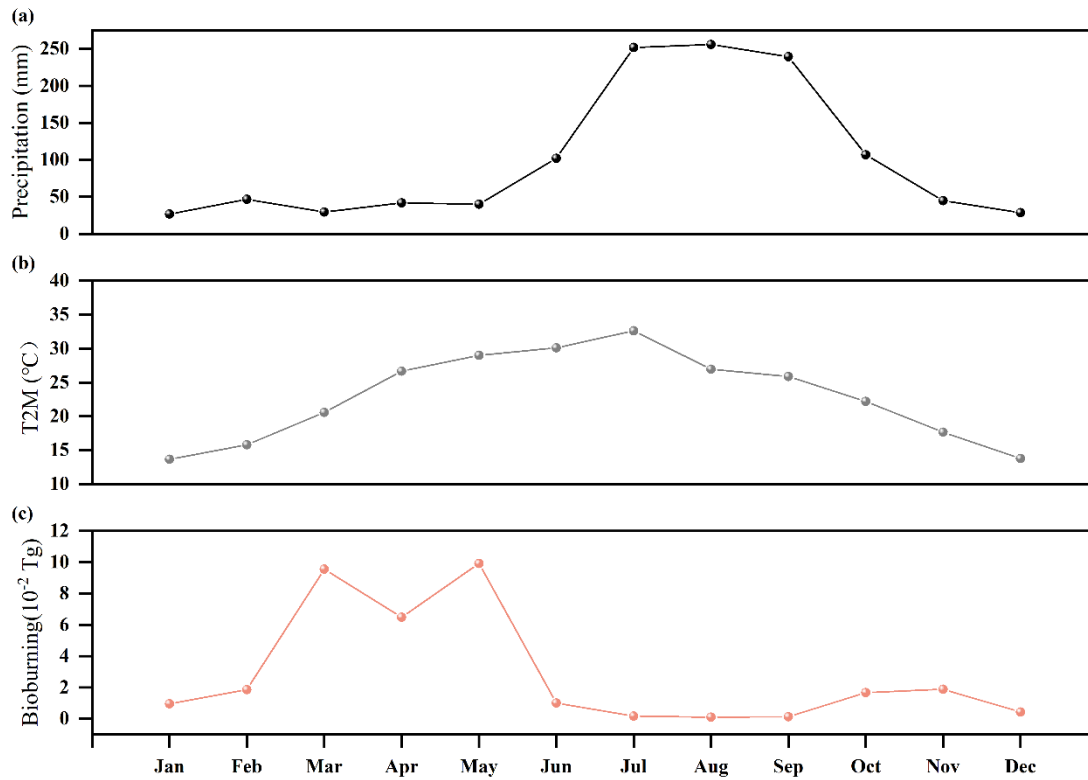


Figure S4. Monthly precipitation (a), temperature (b) from MERRA2 and biomass burning emission (c) from GFED4 in 2019.

RC: 39) *Spatial and Seasonal variation of ammonia emission: No posterior data in Fig.S3.*

AR: Thanks for comment. We have now added the posterior data to Figure S3 as requested.

Figure in manuscript:

