Global Biogeochemical Cycles

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Key Points:
- We report records of N2O concentration for the last 2,000 yr from the Styx ice core in Antarctica and NEEM ice core in Greenland.
- The data accurately reveal the timing and magnitude of N2O variations on centennial time scales.
- Variations in N2O are associated with changes in tropical and subtropical land hydrology and marine productivity.

Supporting Information:
- Supporting Information S1

Correspondence to:
J. Ahn
jinhoahn@snu.ac.kr

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Y. Ryu, J. Ahn, J.-W. Yang, E. J. Brook, A. Timmermann, T. Blunier, S. Hur, and S.-J. Kim

1School of Earth and Environmental Sciences, Seoul National University, Seoul, South Korea, 2Now at Department of Geosciences, Princeton University, Princeton, NJ, USA, 3Now at Université Paris-Saclay, CNRS, CEA, UVSQ, Laboratoire des Sciences du Climat et de l’Environnement, Institut Pierre-Simon Laplace, Gif-sur-Yvette, France, 4College of Earth, Ocean and Atmospheric Sciences, Oregon State University, Corvallis, OR, USA, 5Center for Climate Physics, Institute for Basic Science, Pusan, South Korea, 6Pusan National University, Pusan, South Korea, 7Niels Bohr Institute, University of Copenhagen, Copenhagen, Denmark, 8Korea Polar Research Institute, Incheon, South Korea

Abstract
The continuous growth of atmospheric nitrous oxide (N2O) is of concern for its potential role in global warming and future stratospheric ozone destruction. Climate feedbacks that enhance N2O emissions in response to global warming are not well understood, and past records of N2O from ice cores are not sufficiently well resolved to examine the underlying climate-N2O feedbacks on societally relevant time scales. Here, we present a new high-resolution and high-precision N2O reconstruction obtained from the Greenland NEEM (North Greenland Eemian Ice Drilling) and the Antarctic Styx Glacier ice cores. Covering the N2O history of the past two millennia, our reconstruction shows a centennial-scale variability of ~10 ppb. A pronounced minimum at ~600 CE coincides with the reorganizations of tropical hydroclimate and ocean productivity changes. Comparisons with proxy records suggest association of centennial- to millennial-scale variations in N2O with changes in tropical and subtropical land hydrology and marine productivity.

Plain Language Summary
Nitrous oxide (N2O) is a greenhouse and ozone-depleting gas. The growing level of N2O in the atmosphere is of global concern, and records of past N2O variations can provide an important context for understanding the links between N2O and climate change. In this study, we report new, high-quality N2O records covering the last two millennia using ice cores obtained from Greenland and Antarctica. Our N2O records show rapid centennial-scale changes in atmospheric N2O and confirm a pronounced minimum near 600 CE. Comparison with climate records suggests that hydroclimate change on land and changes in marine productivity contribute to centennial- to millennial-scale N2O variations.

1. Introduction
Improved knowledge of greenhouse gas-climate feedbacks is required to understand past and future climate changes. Nitrous oxide (N2O) is a particularly important greenhouse gas with a global warming potential ~260 times greater than that of CO2 for a time horizon of 100 yr (Myhre et al., 2013). With the regulation of chlorofluorocarbon emissions, N2O is becoming the most important ozone-destroying substance in the stratosphere (Ravishankara et al., 2009). Nonetheless, the processes governing atmospheric N2O variability still remain elusive.

Microbial production in soils and the ocean are dominant sources of atmospheric N2O (Butterbach-Bahl et al., 2013). The atmospheric N2O concentration is primarily regulated by the balance of these terrestrial and oceanic sources (Freling et al., 2012) with the tropical stratospheric upwelling and photoysis in the stratosphere (Khosrawi et al., 2013; Olsen et al., 2001; Prather et al., 2015). The use of nitrogen fertilizer and other industrial activities in the modern era have added extra reactive nitrogen to the global ecosystems, causing a continuous increase in atmospheric N2O levels since the industrial revolution (Galloway et al., 2008). In addition, natural emissions are thought to be sensitive to future climate change (Battaglia & Joos, 2018; Denman et al., 2007; Martinez-Rey et al., 2015; Voigt et al., 2017).

Nitrification and denitrification are considered to be the two major pathways for natural N2O production (Butterbach-Bahl et al., 2013; Ji et al., 2015). In the presence of molecular oxygen, ammonium (NH4+) can
be oxidized to nitrite (NO$_2^-$) and nitrate (NO$_3^-$) (nitrification), with N$_2$O as a by-product. In anoxic or suboxic conditions, denitrification reduces NO$_3^-$ or NO$_2^-$ into dinitrogen gas (N$_2$) in a stepwise fashion, with N$_2$O as an intermediate product. Because these microbial processes are highly dependent on temperature and oxygen levels in both marine and terrestrial environments, the feedbacks between N$_2$O production and climate conditions are important.

Our understanding of climate-N$_2$O feedbacks relies, in part, on paleoatmospheric records and modeling studies. In this context, air bubbles trapped in polar ice cores provide a unique archive for reconstructing and testing ancient atmospheric composition changes through climate history. Several detailed ice core N$_2$O records have been developed over the past few decades (Fischer et al., 2019; Flückiger et al., 2002, 2004; Schilt et al., 2010; Spahni et al., 2005). These records capture long-term variations, such as glacial-interglacial cycles, millennial-scale variations during the Holocene, and the N$_2$O responses to specific abrupt climatic events, including the preboreal transition at the end of the last glacial period and earlier Dansgaard-Oeschger events (Fischer et al., 2019; Flückiger et al., 1999, 2002; Schilt et al., 2010, 2014; Sowers et al., 2003; Spahni et al., 2005). Although previous studies reported pronounced covariability of N$_2$O with northern hemispheric temperature on glacial-interglacial and millennial scales (Flückiger et al., 2002; Schilt et al., 2010), available ice core N$_2$O records for the Holocene have not been sufficiently consistent to allow for an examination of smaller changes on submillennial time scales. Specifically, the past two millennia are thought to be affected by natural climate variability and the growing human influence. Centennial-scale variations are not well resolved in existing ice core N$_2$O records (MacFarling Meure et al., 2006; Prokopiou et al., 2018), apart from the strong atmospheric N$_2$O increase over the past 200 yr (Flückiger et al., 1999; Machida et al., 1995; Sowers et al., 2002).

To improve our understanding of the key drivers of atmospheric N$_2$O variability on centennial time scales, we present new high-resolution (~15 yr) ice core N$_2$O records from both the Greenland NEEM (North Greenland Eemian Ice Drilling) and the Antarctic Styx Glacier ice cores, which cover the past two millennia, and investigate the underlying control mechanisms.

2. Materials and Methods

2.1. Samples and Gas Chronology

Styx Glacier ice and NEEM ice were used for N$_2$O measurements. The climatic information of the ice coring sites is listed in supporting information Table S1. The 210.5 m long Styx Glacier ice core and firm air samples were obtained during the Korean ice core drilling campaign in 2014–2015 (Han et al., 2015). The gas age scale of the Styx Glacier record was obtained by synchronizing its CH$_4$ record to the West Antarctic Ice Sheet (WAIS) divide CH$_4$ record on the WD2014 age scale (Buizert et al., 2015). The uncertainty of the correlation is ~±20 yr (Yang et al., 2018). The Δage (difference in ice age and gas age) of the Styx Glacier ice was estimated to be ~318 yr (Yang et al., 2018). The gas chronology of the NEEM main ice core was also synchronized to the WD2014 age scale to solve the observed age discrepancy of ~20 yr between the NEEM-2011-S1 and WD2014 age scales (Rhodes et al., 2013) (Figure S1). The NEEM-2011-S1 core is a shallow parallel core to the NEEM main core. The estimated age difference between the main ice core and S1 ice core is less than ~5 yr (Sigl et al., 2015), smaller than the age uncertainty resulted from wiggle matching CH$_4$ records (~±20 yr). No high-resolution CH$_4$ data exist below 408.96 m (291 CE) (Rhodes et al., 2013). We estimated the gas age for the deeper NEEM ice by linear extrapolation of the depth-gas age relationship at depth of 338–408 m.

2.2. Ice Core N$_2$O Measurements

We used a high-precision measurement method (Ryu et al., 2018) to analyze both the NEEM and Styx Glacier ice and generate a composite N$_2$O record. Ice core samples are cut into subsamples (~20 g) in a walk-in freezer at a temperature of −20°C to prevent melting. For each depth, samples are duplicated or quadruplicated to estimate the reproducibility of measurements. Ice pieces are placed in glass flasks sealed to a Conflat Flange with a copper gasket for sealing at a high vacuum level. The flasks are then submerged into a prechilled ethanol bath (< −75°C) and evacuated to remove the ambient air in the flask and any contaminants on the ice core surface, for 50 min. After sufficient evacuation, gases in the ice core bubbles are liberated by melting. The N$_2$O mixing ratio of the liberated air in the headspace is measured by an
Agilent 7890B Gas Chromatograph equipped with a Micro-Electron Capture Detector. Before and after the sample measurement, we analyzed a standard N$_2$O gas with a concentration of 329.9 ppb from the National Oceanic and Atmosphere Administration NOAA-2006A N$_2$O scale. Due to the high solubility of N$_2$O in water, we repeated the freezing-melting cycle to liberate the N$_2$O trapped during the extraction procedure. The additional freezing-melting step results in a ~6% correction to the total N$_2$O concentration.

3. Data Quality

Both ice cores have fairly high accumulation rates (NEEM: 0.22 and Styx Glacier: 0.13 m ice equivalent/yr; supporting information Table S1), resulting in a small smoothing effect on gas records due to gas diffusion and gradual bubble close-off processes in the firn layer (transition zone from snow to ice on top of the ice sheet). The estimated widths of the gas age distribution at half height from firn densification models for both ice cores are smaller than 40 yr (Buizert et al., 2012; Jang et al., 2019). Thus, both ice cores can resolve centennial-scale changes in atmospheric composition. We estimated the uncertainty of the N$_2$O concentration by replicating measurements with series of adjacent eight samples within ~20 cm intervals (corresponding to 1–2 yr in mean age change) and obtained pooled standard deviations of 3.4 ppb for Styx Glacier ice and 2.8 ppb for NEEM ice (supporting information Table S2). These are greater than the analytical uncertainty of 1.5 ppb which is determined from measurements of Styx replicate ice samples within 5 cm depth intervals (Ryu et al., 2018), indicating that N$_2$O concentration in the ice core varies in cm scales. Alteration of N$_2$O concentration by in situ microbial activity in the ice (Miteva et al., 2016; Rohde et al., 2008) could contribute the N$_2$O variations and may explain the greater scatter in longer sample interval. However, we do not believe that in situ production of N$_2$O can explain the N$_2$O variations we describe here, because the variations in N$_2$O measured in the two cores we studied are remarkably coherent (Figure 1a) on centennial time scales. The two cores have different gas age-ice age (Δage) differences (~310 yr for Styx, Jang et al., 2019, and ~188 yr for NEEM, Buizert et al., 2012), which means that dust variations would be offset differently with the gas record in each core. Instead, the agreement indicates that these centennial changes represent the true atmospheric signal. Owing to the higher anthropogenic emissions in the Northern Hemisphere, current atmospheric N$_2$O levels are 0.8 ppb higher in the Northern Hemisphere than in the Southern Hemisphere (Ishijima et al., 2009). However, the uncertainty of our ice core records from Greenland and Antarctica does not allow to detect such a small N$_2$O gradient for the preindustrial period. To focus on the joint variability in both cores, we develop a composite N$_2$O record by averaging the NEEM and Styx Glacier N$_2$O records, which were initially interpolated onto the same time axis. Our new data show subtle centennial-scale changes, which were not well resolved in previous ice core N$_2$O records (Fischer et al., 2019; Flückiger et al., 1999, 2002; MacFarling Meure et al., 2006; Schilt et al., 2010) (supporting information Figure S2).

4. Results

As shown in Figure 1, the composite N$_2$O records exhibit two distinctive features during the late preindustrial Holocene: (1) a minimum near 600 CE and (2) centennial-scale fluctuations of about 10 ppb. The 600 CE minimum was previously observed in the Law Dome ice core N$_2$O record, although the uncertainty of the N$_2$O data (± 6.5 ppb) was not sufficient to resolve the magnitude of the N$_2$O decrease (MacFarling Meure et al., 2006). In our measurements, N$_2$O from both NEEM and Styx Glacier confirms that a local minimum resides near 600 CE, thereby consolidating the finding of a substantial change in the land and/or ocean N$_2$O fluxes (Figure 1a). The N$_2$O concentration decreases from ~265 ppb at 200 CE to ~257 ppb at 600 CE.
This ~8 ppb decrease corresponds to average reduction of ~0.3 Tg N yr\(^{-1}\) in N\(_2\)O emissions. This value is calculated with a simple mass burden to mixing ratio (4.79 Tg N/ppb) (Prather et al., 2015) and the assumption that changes in the stratospheric N\(_2\)O sink are negligible (see supporting information).

Centennial-scale N\(_2\)O fluctuations are also prominent (Figure 1b). To further examine centennial-scale variations in the N\(_2\)O budget, we estimated emissions from the concentration data using a two-box model (Figure 1c). To estimate N\(_2\)O emissions, Monte Carlo simulations incorporating uncertainties in N\(_2\)O concentration and lifetime were run for each time interval (see supporting information for details). We find that emission ranges from 10.2–11.7 Tg N yr\(^{-1}\) on average, similar to the current estimates of the natural N\(_2\)O flux from both top-down and bottom-up approaches (Davidson & Kanter, 2014) (10–12 Tg N yr\(^{-1}\)). As expected, the atmospheric N\(_2\)O concentration shows a delayed response of several decades after the change in the N\(_2\)O emission (Figures 1b and 1c).

5. Discussion: N\(_2\)O and Paleoclimate

N\(_2\)O production relies on the metabolic processes of specific groups of microorganisms. Both terrestrial and marine N\(_2\)O sources are expected to respond to changes in the large-scale climate conditions. Environmental factors such as temperature and oxygen availability are believed to have significant impacts on the regulation of N\(_2\)O production. In the case of terrestrial N\(_2\)O sources, which account for ~70% of the global N\(_2\)O emissions, the water-filled pore space (WFPS) is one of the key factors controlling oxygen availability and N\(_2\)O production in the soil environments (Davidson et al., 2000). An optimum WFPS of 70–80% in the soil increases microbial activity and N\(_2\)O emissions (Bouwman et al., 2013). Tropical forest soils are the main terrestrial N\(_2\)O source (Stehfest & Bouwman, 2006; Xu et al., 2017), and the rainfall variations throughout the monsoon season have been suggested as the key factor influencing the interannual variability of the N\(_2\)O emission (Werner et al., 2007). The soil temperature is another important climate factor that can enhance terrestrial microbial N\(_2\)O production. Model studies suggest that on decadal to multidecadal time scales, temperature is the main climate parameter driving changes in terrestrial N\(_2\)O emission (Xu et al., 2012; Zaehle et al., 2011). Oceanic N\(_2\)O production is tightly associated with high primary productivity regions. Ocean systems are considered to contribute approximately 30% of natural N\(_2\)O emissions (Voss et al., 2013), and high productivity regions contribute a greater portion of the marine N\(_2\)O emissions. The eastern tropical Pacific (ETP) and Arabian Sea (AS), where considerable N\(_2\)O accumulation occurs on the upper boundary of widely developed oxygen minimum zones (OMZs) (Hamersley et al., 2007; Ito & Deutsch, 2013), are well-known major oceanic N\(_2\)O source regions. The thermocline depth and strength of the upwelling activity in these regions may influence the nutrient supply that sustains the primary productivity and, in turn, the N\(_2\)O production rate. The dissolved O\(_2\) in upper ocean can affect N\(_2\)O production and emission variability since both denitrification and nitrification are sensitive to dissolved O\(_2\), closely tied to aerobic remineralization of organic matter (Battaglia & Joos, 2018; Martinez-Rey et al., 2015).

The local N\(_2\)O minimum at 600 CE coincides with an overall weakening of the tropical and subtropical monsoon and lower productivity in major oceanic N\(_2\)O source regions, as indicated by a variety of paleoclimate proxy records (Figure 2 and supporting information Figure S3). The N\(_2\)O level continuously decreased until 600 CE and then increased until 1200 CE. The timing of this N\(_2\)O minimum is coincident with reduced precipitation in the tropical land N\(_2\)O source areas, including subtropical China (Wang et al., 2005), Southeast Asia (Steinke et al., 2014; Wurtzel et al., 2018), India (Anderson et al., 2002, 2010; Gupta et al., 2003), and central America (Bhattacharya et al., 2015; Curtis et al., 1996). Because the optimum conditions for microbial N\(_2\)O production requires a high level of soil moisture (~70% to 80%) to achieve low O\(_2\) availability, a weakening in the monsoon strength in the tropical land source areas may decrease N\(_2\)O production. The hydroclimate proxies in South America are not sufficiently consistent to confirm changes in the rainfall patterns at 600 CE; however, some of them suggest an abnormally drier climate (Bird et al., 2011; Kanner et al., 2013; Novello et al., 2012), illustrating a potential regional contribution of hydroclimate changes in South America to lower N\(_2\)O emissions in this period. There are a few available hydroclimate proxy records in central Europe (Fohlmeister et al., 2012) and central Africa (Shanahan et al., 2009; Wolff et al., 2011), resolving the last 2,000 yr; however, the climate in these areas became wetter or showed no significant changes in this period, possibly contributing to the increase in the regional N\(_2\)O emission. The marine
sediment records from AS and the equatorial eastern Pacific, which account for a significant portion of the oceanic N2O emissions, show an overall reduction in productivity near 600 CE, which would further lead to a reduction in the N2O emission (Figure 2).

To deduce the extent and patterns of the hydrological changes on land, we compared our N2O composite with selected temporally well-resolved hydroclimate proxies (Figure 3). Among the cave speleothem proxy records ($\delta^{18}$O of speleothem calcium carbonate) in China, representing the strength of the Eastern Asian Summer Monsoon (EASM), the lower-latitude cave speleothem records from Dongge cave (Wang et al., 2005) (25.28°N) show a minimum near ~500 CE, while Heshang (Hu et al., 2008) (30.27°N) and Wanxiang (Zhang et al., 2008) caves (33.32°N) records do not share this pattern (supporting information Figure S4). The Wanxiang cave records (Zhang et al., 2008) indicate an even stronger EASM between 400 and 800 CE. This latitudinal discrepancy in the EASM proxies is suggested to result from changes in the width of the tropical rainfall belt with the migration of the Intertropical Convergence Zones (Denniston et al., 2016). Thus, the weakening of the EASM strength was possibly constrained to the tropical and subtropical regions.

The western tropical Pacific and Indian subcontinent also experienced diminishing monsoon activity; the Indonesian-Australian Summer Monsoon (IASM) and Indian Summer Monsoon (ISM) weakened between 400 and 800 CE (Figures 3b and 3c and supporting information Figure S3). The lake sediment titanium to calcium ratio (Ti/Ca) records from Sumba Island (9.22°S, 118.89°E), the southeastern part of Indonesia, documented the past changes in the summer monsoon rainfall and showed a strongly reduced IASM rainfall near 600 CE (Steinke et al., 2014) (Figure 3b). Highly resolved tropical Australian cave records (Denniston...
et al., 2016) and Sumatra Island (Wurtzel et al., 2018) supported the weakened IASM in this period. Principal component analysis on the Flores cave speleothem multiproxy from southeast Indonesia shows a drier hydrology during this period (Griffiths et al., 2016); however, it explains only ~50% of the variation pattern, and the oxygen isotope records (Griffiths et al., 2009) ($\delta^{18}O$ of calcium carbonates) from the Flores speleothem agree with a weakened IASM between 400 and 800 CE.

Like the tropical and subtropical western Pacific regions, whose hydrology is largely affected by the monsoon, the precipitation in the Amazon River basin is also greatly controlled by South American Summer Monsoon (SASM). Speleothem records from Huagapo cave (11.27°S, 75.79°W) in the Eastern Andes show a weakening of the SASM during the $N_2O$ minimum at 600 CE (Kanner et al., 2013) (Figure 3d); however, other SASM proxies are not sufficiently consistent to confirm its weakened strength at 600 CE (Figure 2 and supporting information Figure S3). Although our interpretation of these hydroclimate proxies is limited by the discrepancies among the proxy records, a large portion of the tropical and subtropical monsoon regions appear to indicate a change toward drier conditions near 600 CE.

What caused these globally diminishing monsoon activities at 600 CE is unclear; however, hydroclimate change and the associated reduced soil moisture content in the tropical $N_2O$ source regions likely contributed to the reductions in the terrestrial $N_2O$ emissions. During this period, the northern hemispheric temperature is believed to be lower than average (Ljungqvist, 2010; Mann & Jones, 2003; Moberg et al., 2005) (supporting information Figure S5), which could further subdue the terrestrial $N_2O$ emission by suppressing microbial activity. Previous work suggested that northern hemispheric temperature change caused millennial-scale $N_2O$ changes in the Holocene, showing parallel variation with $N_2O$ (Flückiger et al., 2002; Schilt et al., 2010). However, a quantitative estimation of the temperature change is difficult owing to the spatially dispersed temperature proxy records.

The weakening of the ISM prevents upwelling and reduces the primary productivity in the AS, likely leading to decreased $N_2O$ production owing to the reduction in organic matter input, which is required for denitrification. The abundance of the planktonic foraminifera *Globigerina bulloides* in a sediment core from AS (Anderson et al., 2002, 2010; Gupta et al., 2003) (Figure 3e) shows a minimum at 600 CE, reflecting the reduction in the primary productivity, consistent with the reduction in the marine $N_2O$ flux from AS. Similarly, the nitrogen isotope ratios ($\delta^{15}N$) of organic matter in a sediment core off Peru suggest a reduction in the primary productivity and weakened OMZ development in ETP near 600 CE, followed by a gradual strengthening (Salvatteci et al., 2014) (Figure 3f). OMZs are the regions where active nitrogen loss (both denitrification and anammox) take place, contributing ~4% of the estimated oceanic $N_2O$ production (Battaglia & Joos, 2018). In addition, the combination of aerobic remineralization of organic matter followed by nitrification results in strong regional $N_2O$ production where OMZs are well developed (Yang et al., 2020). Thus, a reduction in the OMZ in ETP and AS could also lead to a decrease in oceanic $N_2O$ emissions. In summary, both terrestrial and marine palaeoclimate records suggest that the reorganization of...
tropical rainfall and wind patterns may have contributed to the marked decrease in N₂O fluxes into the atmosphere near 600 CE.

This period was thought to be affected by changes in the periodical North Atlantic cooling events, known as Bond events (Bond et al., 2001). The North Atlantic cooling can be linked to the changes in Asian southeast monsoon and AS productivity (Gupta et al., 2003). Thus, the North Atlantic cooling can be associated with the subdued productivity of N₂O near 600 CE. Dongge cave records may indicate a possible linkage between Asian monsoon strength and the North Atlantic cooling (Wang et al., 2005). Possible changes in climate during this time period include Pacific sea surface temperature (SST) gradient changes (Kanner et al., 2013) and northern hemispheric temperature changes (Kathayat et al., 2017), both of which could contribute to the reorganization of the tropical and subtropical hydroclimates.

On centennial time scales, regional monsoon proxies and oceanic productivity records are not highly correlated with our N₂O flux estimates. The low correlation among the N₂O flux and climate proxies might be because of age uncertainty of proxies and our N₂O data, which can reach several decades in age difference between the N₂O and the palaeoproxy data. Tropical rainfall change in such a short-term temporal scale may be spatially inconsistent and possibly lead to the lack of predominance in the N₂O emission. The only meaningful correlation for the N₂O flux was achieved with the annually resolved central Pacific El Niño–Southern Oscillation (ENSO) proxies (Figure 3g). ENSO variance records based on Taiwan tree ring δ¹⁸O show in-phase correlation with the calculated N₂O flux (Liu et al., 2017) (r = 0.41). The centennial-scale decrease in the N₂O flux corresponds to a suppressed ENSO variance, whereas the periods of high ENSO variance coincide with enhanced N₂O emission. A correlation coefficient r of 0.41 is a conservative value, given the relative age uncertainty between the records. While we cannot address the physical mechanism between the central Pacific ENSO and N₂O flux changes, it may reflect the relationship between the N₂O production and climate variability in low-latitude regions.

The discussion above focuses on N₂O production changes rather than changes in sinks related to photochemical decomposition in the stratosphere. The fluctuations in our N₂O records appear to match with proxies of solar activity changes. N₂O concentrations are in phase with total solar irradiance reconstruction data (Steinhilber et al., 2009), showing low levels of N₂O during the solar minima events of the past millennium (supporting information Figure S6). A recent study reported that the N₂O lifetime varies with solar activity (Prather et al., 2015); the lifetime of N₂O decreases in solar maxima (while it increases in solar minima), indicating more rapid N₂O loss and decrease of N₂O during the solar maxima. However, our observation is a positive correlation between N₂O and solar activity. This may imply that the changes in N₂O loss due to solar activity are not large enough to counterbalance changes in N₂O sources during the past two millennia.

To better understand the physical linkage between climate and N₂O changes, further numerical modeling experiments are required as well as chemical and isotopic data. The isotopic composition of N₂O molecule (δ¹⁵N, δ¹⁸O, and site preference) has been used as the footprint for distinguishing its sources and production processes (Fischer et al., 2019; Prokopiou et al., 2018; Schilt et al., 2014; Sutka et al., 2006), and it may provide a way to deduce the mechanistic explanation for the N₂O variations. However, published N₂O isotope data do not have sufficient temporal resolution and precision to facilitate the investigation of the relative contribution of marine and terrestrial sources or microbial production mechanisms (nitrification vs. denitrification) on submillenial time scales (Prokopiou et al., 2018). Future studies of N₂O isotopologues and higher-resolution N₂O isotope records will be very useful to improve our understanding of the connection between N₂O emissions and regional and large-scale climate.

6. Conclusion

We present the submillennial-scale N₂O variations during the past two millennia with highly resolved ice core records and evaluate their possible causes. Our data show variability of ~10 ppb on a centennial time scale. A pronounced local minimum at ~600 CE coincides with the changes in tropical hydroclimate and ocean productivity. Both terrestrial and marine N₂O production rate changes in response to climate are likely explanations for the atmospheric N₂O concentration changes. Disentangling the underlying changes in terrestrial and marine N₂O production rate changes requires further analysis of N₂O isotopes and targeted modeling experiments with state-of-the-art Earth system models.
Data Availability Statement

The data are available at NOAA (National Oceanic and Atmospheric Administration) Paleoclimate website (https://www.ncdc.noaa.gov/paleo-search/study/30752).

References

Anderson, D. M., Baulcomb, C. K., Du Vivier, A. K., & Gupta, A. K. (2010). Indian summer monsoon during the last two millennia. *Journal of Quaternary Science, 25*(6), 911–917. https://doi.org/10.1002/jqs.1369

Anderson, D. M., Overpeck, J. T., & Gupta, A. K. (2002). Increase in the Asian southwest monsoon during the past four centuries. *Science, 297*(5581), 596–599. https://doi.org/10.1126/science.1072881

Battaglia, G., & Joos, F. (2018). Marine N2O emissions from nitrification and denitrification constrained by modern observations and projected in multimillennial global warming simulations. *Global Biogeochemical Cycles, 32*, 92–121. http://doi.org/10.1002/2017GB005671

Bhattacharya, T., Byrne, R., Böhnel, H., Wogau, K., Kienel, U., Ingram, B. L., & Zimmerman, S. (2015). Cultural implications of late Holocene climate change in the Cuencita Oriental, Mexico. *Proceedings of the National Academy of Sciences of the United States of America, 112*(6), 1693–1698. https://doi.org/10.1073/pnas.1405653112

Bird, B. W., Abbott, M. B., Vuille, M., Rodbell, D. T., Stansell, N. D., & Rosenmeier, M. F. (2011). A 1,230-year-long annually resolved record of the South American summer monsoon from the Peruvian Andes. *Proceedings of the National Academy of Sciences of the United States of America, 108*(21), 8583–8588. https://doi.org/10.1073/pnas.1003719108

Bond, G., Kromer, B., Beer, J., Muscheler, R., Evans, M. N., Showers, W., et al. (2001). Persistent solar influence on North Atlantic climate during the Holocene. *Science, 294*(5549), 2130–2136. https://doi.org/10.1126/science.1065680

Bouwman, A. F., Beusen, A. H. W., Griffioen, J., Van Groenigen, J. W., Heting, M. M., Oenema, O., et al. (2013). Global trends and uncertainties in terrestrial denitrification and N2O emissions. *Philosophical Transactions of the Royal Society, B: Biological Sciences, 368*, 20130112. https://doi.org/10.1098/rstb.2013.0112

Buizert, C., Cuffey, K. M., Severinghaus, J. P., Baggenstos, D., Fudge, T. J., Steig, E. J., et al. (2015). The WAIS divide deep ice core WD2014 chronology—Part 1: Methane synchronization (68–31 ka BP) and the gas age–ice age difference. *Climate of the Past, 11*(2), 153–173. https://doi.org/10.5194/cp-11-153-2015

Buizert, C., Martinerne, P., Petrenko, V. Y., Severinghaus, J. P., Trudinger, C. M., Witrant, E., et al. (2012). Gas transport in firm: Multiple-tracer characterisation and model intercomparison for NEEM, northern Greenland. *Atmospheric Chemistry and Physics, 12*(9), 4259–4277. https://doi.org/10.5194/acp-12-4259-2012

Butterbach-Bahl, K., Baggs, E. M., Dannenmann, M., Kiese, R., & Zecheimert-Boltenstern, S. (2013). Nitrous oxide emissions from soils: How well do we understand the processes and their controls? *Philosophical Transactions of the Royal Society, B: Biological Sciences, 368*(1621), 20130122. https://doi.org/10.1098/rstb.2013.0122

Curtis, J. H., Hodell, D. A., & Brenner, M. (1996). Climate variability on the Yucatan Peninsula (Mexico) during the past 3500 years, and implications for Maya cultural evolution. *Quaternary Research, 46*(1), 37–47. https://doi.org/10.1006/qres.1996.0042

Davidson, E. A., & Kanter, D. (2014). Inventories and scenarios of nitrous oxide emissions. *Environmental Research Letters, 9*(10), 105012. https://doi.org/10.1088/1748-9326/9/10/105012

Davidson, E. A., Keller, M., Erickson, H. E., Vrachot, L. V., & Veldkamp, E. (2000). Testing a conceptual model of soil emissions of nitrous and nitric oxides. *Biocience, 50*(8), 667. https://doi.org/10.1641/0006-3568(2000)050[0667:tcmod]2.0.co;2

Denman, K. L., Brasseur, G., Chidthaisong, A., Ciais, P., Cox, P. M., Dickinson, R. E., et al. (2007). Couplings between changes in the climate system and biogeochemistry coordinating lead authors: This chapter should be cited as. In *Climate change 2007: The physical science basis*. Contribution of Working Group I to the Fourth Assessment Report of the Intergovernmental Panel on Climate Change (pp. 500–587). Cambridge, United Kingdom and New York, NY, USA: Cambridge University Press. Retrieved from https://www.globalcarbonproject.org/pdf/AR4WG1_Pub_Ch07.pdf

Denniston, R. F., Ummenhofer, C. J., Wanamaker, A. D., Lachniet, M. S., Villarini, G., Asmerom, Y., et al. (2016). Expansion and contraction of the Indo-Pacific tropical rain belt over the last three millennia. *Scientific Reports, 6*(1), 34485. https://doi.org/10.1038/srep34485

Fischer, H., Schmitt, J., Bock, M., Seth, B., Joos, F., Spahni, R., et al. (2019). N2O changes from the Last Glacial Maximum to the preindustrial—Part 1: Quantitative reconstruction of terrestrial and marine emissions using N2O stable isotopes in ice cores. *Biogeochemistry, 166*(20), 3997–4021. https://doi.org/10.1007/s10535-019-00201-2

Flückiger, J., Blunier, T., Stauffer, B., Chappellaz, J., Spahni, R., Kawamura, K., et al. (2004). N2O and CH4 variations during the last glacial epoch: Insight into global processes. *Global Biogeochemical Cycles, 18*, GB1020. https://doi.org/10.1029/2003GB002122

Flückiger, J., Dällenbach, A., Blunier, T., Stauffer, B., Stocker, T. F., Raynaud, D., & Barnola, J. M. (1999). Variations in atmospheric N2O concentration during abrupt climatic changes. *Science, 283*(5425), 227–230. https://doi.org/10.1126/science.283.5425.227

Flückiger, J., Menhinick, E., Stauffer, B., Schwander, J., Stocker, T. F., Chappellaz, J., et al. (2002). High-resolution Holocene N2O ice core record and its relationship with CH4 and CO2. *Global Biogeochemical Cycles, 16*(1), 1010. https://doi.org/10.1029/2001GB000147

Fohlenmeier, J., Schröder-Ritschau, A., Scholz, D., Spötl, C., Riechelmann, D. F. C., Muideslo, M., et al. (2012). Bunker Cave stalagmites: An archive for central European Holocene climate variability. *Climate of the Past, 8*(5), 1751–1764. https://doi.org/10.5194/cp-8-1751-2012

Freng, A., Wallace, D. W. R., & Bange, H. W. (2012). Global oceanic production of nitrous oxide. *Philosophical Transactions of the Royal Society, B: Biological Sciences, 367*(1539), 1245–1255. https://doi.org/10.1098/rstb.2011.0360

Galloway, J. N., Townsend, A. R., Erisman, J. W., Bekunda, M., Cai, Z., Freney, J. R., et al. (2008). Transformation of the nitrogen cycle: Recent trends, questions, and potential solutions. *Science, 320*(5878), 889–892. http://doi.org/10.1126/science.1136674

Griffiths, M. L., Drysdale, R. N., Gagan, M. K., Zhao, J. X., Alyifle, L. K., Hellstrom, J. C., et al. (2009). Increasing Australian-Indonesian monsoon rainfall linked to early Holocene sea-level rise. *Nature Geoscience, 2*(9), 636–639. https://doi.org/10.1038/ngeo605

Griffiths, M. L., Kimbrough, A. K., Gagan, M. K., Drysdale, R. N., Cole, J. E., Johnson, K. R., et al. (2016). Western Pacific hydroclimate linked to global climate variability over the past two millennia. *Nature Communications, 7*, 11719. https://doi.org/10.1038/ncomms11719

Gupta, A. K., Anderson, D. M., & Overpeck, J. T. (2003). Abrupt changes in the Asian southwest monsoon during the Holocene and their links to the North Atlantic Ocean. *Nature, 421*(6921), 354–357. https://doi.org/10.1038/nature01340
