Measurement report: Surface exchange fluxes of HONO during the growth process of paddy fields in the Huaihe River Basin, China

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Abstract: Significant amounts of nitrous acid (HONO) released from soil affect the chemistry of

- 30 the troposphere, as a major precursor of hydroxyl radical. However, the scarcity of in-situ data on soil– atmosphere HONO exchange flux has constrained the comprehension of emission mechanisms and reactive nitrogen budget. Herein, we performed measurements of HONO and NO_x fluxes over paddy fields in the Huaihe River Basin. The entire experiment experienced various agricultural management activities, including rotary tillage, flood irrigation, fertilization, paddy cultivation and growth, and top-
- 35 dressing. HONO and NO exhibited more upward fluxes, whereas NO² deposited to the ground, with average hourly fluxes of 0.07 ± 0.22 , 0.19 ± 0.53 and -0.42 ± 0.44 nmol m⁻² s⁻¹, respectively. Continuous peaks in HONO and NO fluxes were observed during the rotary tillage, and they exhibited a significant correlation ($R = 0.77$). Moreover, a significant correlation ($R = 0.60$) between HONO flux and the product of $J(NO₂) \times NO₂$ was also observed during the daytime. The results indicate that both soil biological
- 40 emissions and light-driven NO² conversion are likely active, collectively influencing the diurnal pattern of HONO flux. Source analysis revealed that the unknown HONO source $(P_{unknown})$ exhibited a diurnal pattern with higher daytime and lower nighttime values. Sensitivity tests demonstrated that photoenhanced NO_2 conversion on the ground could adequately explain P_{unknown} , while the nocturnal HONO production derived from soil emission fluxes (ranging from 0.32 ppbv h^{-1} to 0.79 ppbv h^{-1}) was sufficient
- 45 to elucidate the nighttime $P_{unknown}$. Our study emphasized the variability of HONO fluxes across various agricultural management activities, as well as the importance of heterogeneous $NO₂$ conversion on the ground surface and soil emissions for HONO production.

1. Introduction

50 Nitrous acid (HONO) and nitrogen oxides ($NO_x = NO + NO₂$) are key components of reactive nitrogen (Nr) cycles and significantly influence the atmospheric oxidation capacity through the hydroxyl radical (OH) and ozone (O_3) atmospheric cycles (Kratz et al., 2022; Monks et al., 2009; Weber et al., 2015). The photolysis of HONO contributes to 20 %–90 % of the OH budget, serving not only as an important source of OH in the early morning but also playing a significant role throughout the entire day

- 55 (Elshorbany et al., 2009; Kim et al., 2014; Kleffmann et al., 2005; Nan et al., 2017; Xue et al., 2020). Despite the significance of HONO in atmospheric chemistry, the formation mechanism of HONO is still not well understood, especially during the daytime. Unexpectedly large discrepancies have been found between HONO measurements and predicted values from known mechanisms, implying the existence of unknown sources of HONO that have not yet been identified (Lee et al., 2016; Liu et al., 2019c; Sörgel
- 60 et al., 2011; Su et al., 2011; Tang et al., 2015). Several potential mechanisms have been proposed to explain atmospheric HONO levels, including direct emissions from combustion processes (Nakashima and Kajii, 2017; Nie et al., 2015); the chemical equilibrium between soil nitrite $(NO₂)$ and hydrogen ions (Su et al., 2011); photosensitized reactions of $NO₂$ on organic substances (George et al., 2005), humic acids (Han et al., 2016; Stemmler et al., 2006), soot (Monge et al., 2010), minerals (Ndour et al., 2008),
- 65 urban grime (Liu et al., 2019a), plant leaves (Marion et al., 2021), etc.; photolysis of adsorbed nitrates or nitric acid (Ye et al., 2017; Zhou et al., 2003; Zhou et al., 2011) and ortho-nitrophenols (Bejan et al., 2006; Guo and Li, 2022); direct emission from ammonia oxidizing bacteria and other microorganisms (Oswald et al., 2013; Scharko et al., 2015); desorption of adsorbed HONO from the surface by acid displacement processes (Vandenboer et al., 2013; Vandenboer et al., 2014; Vandenboer et al., 2015), and 70 chemical reactions of hydroxylamine on the surface of soil particles (Ermel et al., 2018). Furthermore, the NH₃-promoted heterogeneous reaction of NO₂ has recently been proposed based on laboratory and field studies (Ge et al., 2019; Li et al., 2018; Xu et al., 2019), however, this mechanism and its atmospheric influences require further investigation.
- Flux measurement is always considered as a useful tool for quantifying ground-level sources of 75 HONO, providing direct insight into the production and loss processes on the surface. In recent years, micrometeorological methods such as relaxed eddy accumulation (REA) and aerodynamic gradient (AG) have been developed and applied in HONO flux research, with field observations primarily conducted in Europe and North America. Ren et al. (2011), Von Der Heyden et al. (2022) and Zhou et al. (2011)

measured HONO fluxes using the REA method in various environments such as agricultural fields,

- 80 forests, and grasslands. The studies revealed that HONO fluxes were primarily driven by the photosensitized NO² reduction and photolysis of adsorbed HNO3. Laufs et al. (2017), Meng et al. (2022) and Sörgel et al. (2015) performed measurements utilizing the AG method over bare soil, corn canopy, forest canopy, and wheat canopy, obtaining similar conclusions. Additionally, the chamber method provides greater flexibility and is suitable for multipoint observations within agricultural fields. Tang et
- 85 al. (2020) and Xue et al. (2019) investigated HONO emissions from agricultural soil in the Huaihe River Basin and the North China Plain (NCP) by employing the open-top dynamic chamber method, confirming that agricultural soil emission is an important source of atmospheric HONO. However, the limited available HONO flux studies indicated different potential HONO precursors, which demonstrated the necessity for more HONO flux measurements to explore potential HONO formation pathways.
- 90 Moreover, most flux measurements are typically conducted for the short term (less than one month), and cannot cover the entire growing season of crop. Research on HONO fluxes in agriculture has primarily focused on wheat–maize rotations and the effects of fertilization. Paddy fields, as a major crop in southern China with unique growth conditions, have received little attention, resulting in limitations in understanding the Nr budget in paddy field ecosystems.
- 95 Cropland, which covers 50 % of the global habitable areas (FAO, 2022), plays a crucial role in the global nitrogen budget. The application of nitrogen fertilizers has been instrumental in boosting food production. Nevertheless, the overuse of fertilizers has also resulted in soil degradation, declining air quality, and adverse effects on human health. Simultaneously, the extensive application of synthetic nitrogen fertilizers in cropland, coupled with their low nitrogen use efficiency (< 50 % on average) 100 (Mueller et al., 2017; Zhang et al., 2015), has led to the release of excess Nr from the soil through microbial processes. Among these, NO_x mediates the production and destruction of $O₃$, influences the formation of the OH radical, and can be oxidized to nitric acid and nitrate, thereby increasing wet and dry nitrogen deposition in ecosystems (Pilegaard, 2013). Notably, the positive effect of nitrogen fertilizer on HONO emissions has been consistently verified (Wang et al., 2021). Xue et al. (2019) reported an
- 105 extraordinarily high HONO flux of 1515 ng N m⁻² s⁻¹ under excessive fertilized conditions, which greatly exceeded the emissions from unfertilized farmland and even surpassed laboratory results. This underscores the significant potential for Nr emissions originating from agricultural soil. Therefore, it is imperative to comprehend the fluxes within agricultural ecosystems to elucidate the mechanisms of Nr

production and loss. The lack of field data on HONO fluxes in paddy fields, coupled with the ambiguous

- 110 impacts of agricultural management activities, hinders our understanding of soil–atmosphere exchange mechanisms. Laboratory studies have also demonstrated HONO and NO emissions at high water content (Wang et al., 2021; Wu et al., 2019), and the anaerobic denitrification in oxygen-limited environments can be an important source of HONO (Bhattarai et al., 2021; Wang et al., 2021; Wu et al., 2019). This highlights the necessity to investigate further the effects of flooded paddy fields and agricultural practices 115 on soil HONO emissions.
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In this study, the soil–atmosphere exchange processes were investigated using the AG method in conjunction with the BroadBand Cavity Enhanced Absorption Spectrometer (BBCEAS) system and NO_x analyzer in paddy fields located in the Huaihe River Basin. The variations in HONO and NO_x levels and fluxes were evaluated across various agricultural management processes from June to July, 120 corresponding to the paddy growing season. Additionally, a particular focus was placed on investigating the sources of HONO during the rotary tillage period and its contribution to atmospheric oxidizing capacity.

2. Materials and methods

2.1 Measurement site

- 125 The field campaign was performed at the Shouxian National Climatological Observatory (32°25′ N, 116°47′ E; 25 m above sea level), located 9 km south of Shouxian, Anhui Province (Fig. S1). This location represents a typical rice–wheat rotation ecosystem in the Huang–Huai agro-ecological region, which serves as the primary grain production area in China, contributing to 18 % of the nation's total grain production. Additionally, it is responsible for 76.3 % of the country's total nitrogen fertilizer 130 application (Cao et al., 2019). The site covers a 17 ha field and is dedicated to the cultivation of rice– wheat rotation. It serves as an experimental site for studying surface–atmosphere exchange. The site is situated amidst other agricultural fields, with a less traffic road to the north (250 m). The prevailing climate in the region is subtropical monsoon, characterized by distinct seasons, with high temperatures and rainfall occurring in the same season. The average annual temperature is 14.8 °C, and the average
- 135 annual precipitation is 905 mm.

2.2 Experimental design

The flux measurement was conducted from 1 June to 14 July 2021, immediately following the winter wheat harvest on 31 May 2021. The tillage process took place over 11 days from 2 June to 13 June, followed by flooding irrigation, and fertilization with compound fertilizer $(N-P_2O_5-K_2O 15\% -$

140 15 %–15 %) of 67.5 kg N ha⁻¹ before 22 June 2021. Consequently, the surface was a mixture of bare soil and sparse winter wheat residues before irrigation (June 13th, 9:00), while the soil became waterlogged after flooding irrigation. The paddy seedlings were transplanted on 26 and 27 June at a density of 1.8 \times 10⁵ plants ha-1 , growing from 0.14 m to approximately 0.22 m during the whole campaign. Additionally, irrigation was employed post-paddy transplantation to mitigate water deficiency during the growth phase, 145 thereby preventing the potential mortality of paddy seedlings. The 46 %–N urea solution of 69 kg N ha- $¹$ was applied as top-dressing on 10 July.</sup>

The concentrations of HONO and $NO₂$ in the ambient air were measured using a homemade BBCEAS instrument with a time resolution of 1 min and detection limits of 54 pptv (2*σ*) for HONO and 98 pptv (2σ) for NO₂. The measurement uncertainty was 8.7 % and 8.1 % for HONO and NO₂, 150 respectively. Further details regarding BBCEAS, such as its principle, instrument parameters and quality control, are described in detail elsewhere (Duan et al., 2018; Tang et al., 2019). NO was measured by custom-built chemiluminescence (Model 42iTL, Thermo Scientific, USA), and O³ were measured with Thermo Scientific Model 49i, with detection limits of 50 pptv for NO and 500 pptv for $O₃$, respectively. The measurement of soil temperature and moisture, as well as meteorological and micrometeorological 155 parameters, are presented in Text S1.

Trace gas profiles of HONO, NO, and NO₂ were obtained using inlets positioned at heights of 0.2 m and 1.6 m, which were adjusted to 0.3 m and 1.6 m on 27 June to accommodate the canopy height, consistently exceeding the canopy height throughout the campaign (Fig. 1). Two BBCEAS instruments were used to measure HONO and NO² at different heights, which were intercompared several times 160 throughout the campaign and exhibited excellent agreement (HONO: $R^2 = 0.989$; NO₂: $R^2 = 0.998$) with slopes close to 1 (Fig. S2). A NO_x analyzer for NO measurement was connected to a Teflon solenoid valve to allow sequential measurements at two different heights. All instruments were placed in the thermostated container controlled by an air conditioner, with the external sampling inlets affixed to a small mast. The sampling inlets were oriented away from the mast towards the prevailing wind direction

165 to minimize turbulence disruption. To prevent photolysis and the condensation of water vapor, the PFA inlet lines (7.5 m length with a 6 mm external diameter) were shielded from radiation and slightly heated with heating tape (the heating temperature was about 30 °C).

Figure 1. The aerodynamic gradient measurement set-up for the determination of HONO, NO and NO² fluxes.

170 **2.3 Aerodynamic gradient fluxes of HONO, NO and NO²**

The HONO, NO and NO² fluxes were calculated by the AG method at time intervals of 30 min, which has been elaborated upon in previous studies (Laufs et al., 2017; Meng et al., 2022; Stella et al., 2012) and will be briefly introduced here. The flux (F_χ) of trace gas is calculated from the friction velocity (u_*) and the mixing ratio scaling parameter (χ_*) as follows:

$$
F_{\chi} = -u_* \chi_* \tag{1}
$$

where u_* is calculated from eddy covariance measurements and χ_* is defined from stability-corrected gradient of the scalar mixing ratio (χ) with height (z) as:

$$
\chi_* = \kappa \cdot \frac{\partial \chi}{\partial \left[\ln(z - d) - \Psi_H \left(\frac{z - d}{L} \right) \right]}
$$
 (2)

The fluxes ($F_{\text{HONO,NO and NO}_2}$) of trace gases at geometric mean height can be expressed as:

180
$$
F_{\text{HONO, NO and NO}_2} = -\kappa \cdot u_* \cdot \frac{\partial c(\text{HONO, NO and NO}_2)}{\partial \left[\ln(z-d) - \Psi_H\left(\frac{z-d}{L}\right)\right]}
$$
(3)

where κ is von Kármán constant ($\kappa = 0.4$), *z* is the height above the ground, *d* is zero plane displacement and was taken as $2/3 \cdot h_c$ (h_c is the canopy height), *L* is the Obukhov length and Ψ_H is integrated stability correction function for scalars (Sutton et al., 1993).

Data from all instruments could not always be collected simultaneously for flux calculation due to 185 various factors such as calibration, malfunction, and disturbances from agricultural activities. Consequently, the affected data were excluded when calculating fluxes. The dataset used for the determination of HONO, NO and $NO₂$ fluxes comprised 68 % for HONO, 81 % for NO, and 86 % for NO2. The total uncertainty in the flux is composed of gradient error and friction velocity error (Laufs et al., 2017; Meng et al., 2022). The average uncertainty for HONO, NO, and NO₂ fluxes were 11 %, 16 %,

- 190 and 20 % (median [25 percentile–75 percentile]), respectively. Furthermore, the fluxes were discarded for very stable conditions with low wind speed and friction velocity. It is important to note that HONO, NO and NO₂ are subject to chemical reactions, which could lead to a vertical divergence of flux between the surface and the measurement height. The influence of chemical reactions during turbulent transport was checked utilizing the Damköhler number (*DA*), as detailed in Text S2. The divergence by chemical
- 195 reactions of HONO could be neglected when interpreting the potential sources of HONO and driving factors of HONO flux. The *DA* for the NO-O3-NO² triad generally exhibited values less than 1, however, a sharp increase in flux divergence occurred when the *DA* became greater than 1 (Stella et al., 2012). Additionally, the upward NO₂ flux exhibited a significant correlation $(R = 0.82)$ with NO flux, suggesting that the upward $NO₂$ fluxes could be attributed to the reaction of NO and $O₃$. Consequently, in light of 200 the influence of chemical reactions on the fluxes of NO and $NO₂$, these fluxes (5.9 % for NO flux and
	- 10.5 % for NO² flux) were excluded from subsequent analysis.

3. Results and discussion

3.1 Overview of meteorological and soil parameters

The time series of meteorological parameters throughout the observation period is shown in Fig. 2. 205 The campaign weather was dominated by sunny days, with 64 % of the days having a daily maximum global radiation above 700 W m⁻². The ambient temperature and soil temperature ranged from 17.0 to 36.6 °C and 20.0 to 34.8 °C, with average values of 26.8 ± 3.5 °C and 26.5 ± 2.7 °C, respectively. The relative humidity (RH) and soil water-filled pore space (WFPS) ranged from 22 % to 98 % and 44 % to 88 %, with average values of 77 % \pm 17 % and 69 % \pm 15 %, respectively. The average wind speed was 210 3 m s⁻¹, with a maximum wind speed of 11.0 m s⁻¹ occurring during the rotary tillage period. The PM_{2.5} concentration varied from 1 to 100 μ g m⁻³, with its daily average value remaining below the Chinese National Ambient Air Quality Standard (Class II: 75 μg m⁻³). Intermittent rainfall occurred from June 13 to July 5, with a total precipitation of 186.1 millimeters. Notably, after irrigation in the agricultural field on June 13, WFPS increased from 45 % to 80 %.

Figure 2. Temporal variations of meteorological parameters (wind speed and direction, air temperature, relative humidity and precipitation), soil temperature, WFPS and PM_{2.5} measured from 1 June to 14 July 2021.

3.2 Mixing ratio differences and fluxes of HONO, NO and NO²

The field campaign was performed across various agricultural management activities, including rotary tillage (June 2–13), flooding irrigation (June 13–19), fertilization (June 19–21), paddy cultivation (June 26–27), and top-dressing (July 10). Figure 3 illustrates the time series of HONO, NO, NO² and O³ mixing ratios. Throughout the campaign, the ambient O_3 concentrations varied from 0.54 to 131.57 ppbv, with an average of 48.44 ± 26.29 ppbv. The peak of NO mixing ratios reached 36.02 ppbv during rotary tillage, and the average mixing ratios of NO at lower (0.2/0.3 m) and upper levels (1.6 m) were 0.75 \pm 225 2.21 ppbv and 0.46 ± 1.16 ppbv, respectively. Higher NO mixing ratios were measured at the lower level, likely due to soil NO emissions caused by microbiological activity (Bargsten et al., 2010; Ludwig et al., 2001). Moreover, the average NO₂ mixing ratios were 4.48 ± 4.96 ppbv and 4.75 ± 4.38 ppbv at lower and upper levels, respectively. The synchronous peaks of NO and $NO₂$ and the decrease of $O₃$ (e.g. in the early hours of June 7) indicated that NO release from soil could react rapidly with O_3 to form NO₂. The ambient HONO mixing ratios ranged from below detection limits to 3.60 ppbv at the lower level and 2.36 ppbv at the upper level, with an average of 0.46 ± 0.59 ppbv and 0.37 ± 0.37 ppbv, respectively. The

average HONO/NO_x ratio of 0.079 ± 0.059 was significantly higher than the range for direct emissions

from vehicle exhaust reported in previous studies (0.003–0.018) (Kirchstetter et al., 1996; Kurtenbach et al., 2001; Liang et al., 2017; Liu et al., 2017; Nakashima and Kajii, 2017; Nakashima and Kondo, 2022),

- 235 and was comparable to the value (0.0929) observed in summer agricultural fields in the NCP (Song et al., 2022). Notably, successive HONO peaks were measured during rotary tillage, with HONO mixing ratios reaching 3.60 ppbv at the lower level. These values exceeded those observed during the winter at the same site (Meng et al., 2022) and were comparable to observations at suburban sites in the Pearl River Delta (Li et al., 2012; Su et al., 2008) and rural sites in the NCP (Xue et al., 2020). However,
- 240 HONO levels declined rapidly following flooding irrigation (see Fig. 3 and Table 1). Subsequent to the fertilization and the top-dressing, a noticeable rise in HONO levels was observed, which could be attributed to the increase in HONO release by fertilizer application at high water contents (Tang et al., 2019; Wang et al., 2021; Wu et al., 2019; Xue et al., 2019). Nevertheless, these levels were significantly lower than the mixing ratios observed during rotary tillage.

245 Table 1. The statistical summary of HONO, NO, NO₂, HONO flux, NO flux and NO₂ flux across various agricultural activities spanning from 1 June to 14 July 2021.

Agricultural activities		HONO (ppbv)		N _O (ppbv)		NO ₂ (ppbv)		HONO flux	NO flux	NO ₂ flux
		$0.2/0.3$ m	1.6 _m	$0.2/0.3$ m	1.6 _m	$0.2/0.3$ m	1.6 _m	(nmol m ⁻² s ⁻¹)	(nmol $m^{-2} s^{-1}$)	(nmol m ⁻² s ⁻¹)
Rotary tillage	Ave	0.99	0.69	1.88	0.87	6.26	6.56	0.26	0.47	-0.72
	Min	0.08	0.07	0.07	0.06	0.70	0.90	-0.62	-0.78	-3.50
	Max	3.60	2.36	36.02	17.80	22.24	35.03	1.86	9.12	0.29
Flood irrigation	Ave	0.19	0.18	0.66	0.46	4.62	4.91	0.02	0.18	-0.40
	Min	0.04	0.06	0.13	0.08	1.11	1.11	-0.40	-0.78	-1.93
	Max	0.52	0.44	9.22	2.12	17.31	19.92	0.32	1.18	0.50
Fertilization	Ave	0.24	0.31	0.33	0.28	5.81	6.15	-0.06	0.03	-0.34
	Min	0.08	0.07	0.06	0.05	0.39	0.64	-0.38	-0.13	-1.37
	Max	0.61	0.71	7.02	4.14	30.30	29.07	0.11	0.31	0.07
After fertilization	Ave	0.30	0.36	0.26	0.26	4.58	4.49	-0.05	0.001	-0.19
	Min	0.06	0.05	0.05	0.06	0.69	0.88	-0.41	-0.23	-1.17
	Max	1.05	1.29	4.09	4.61	19.25	20.32	0.26	0.12	0.26
Paddy cultivation and growth	Ave	0.18	0.21	0.42	0.39	3.45	3.76	-0.05	0.02	-0.34
	Min	0.04	0.05	0.05	0.05	0.29	0.53	-0.70	-1.01	-1.93
	Max	0.63	0.69	15.21	12.40	14.32	15.85	0.42	0.44	0.50
Top-dressing	Ave	0.23	0.19	0.24	0.22	2.78	3.02	0.05	0.03	-0.29
	Min	0.05	0.05	0.05	0.05	0.49	0.63	-0.34	-0.37	-3.26
	Max	1.21	0.51	1.57	1.31	9.59	9.49	0.57	1.27	0.09

Note: Ave, Min, and Max represent the average, minimum, and maximum, respectively. The 0.2/0.3 m and 1.6 m represented the lower and upper levels, respectively.

250 **Figure 3.** Time series of O3, NO, NO2, HONO and the fluxes of HONO, NO, and NO² were determined by the aerodynamic gradient method. The mixing ratios of HONO, NO, NO₂ (lower level: 0.2/0.3 m, upper level: 1.6 m), and O³ were measured above a crop rotation field and averaged for 30 min intervals. Periods of agricultural management activities (rotary tillage, flood irrigation, fertilization, after fertilization, paddy cultivation and growth, top-dressing) are denoted at the top of the graph.

255 The fluxes of HONO, NO, and NO² determined by the AG method are illustrated in Fig. 3. Upward fluxes were commonly observed for HONO and NO, while NO² was deposited to the ground. The magnitudes of observed HONO fluxes ranged from -0.70 to 1.86 nmol $m² s⁻¹$, with an average of 0.07 \pm 0.22 nmol m⁻² s⁻¹, which falls within the range of the HONO flux measurements in rural and suburban regions from the literature (see Table 2). The upward HONO fluxes were mostly observed during rotary 260 tillage, reaching up to 1.86 nmol m⁻² s⁻¹. After the irrigation, the increase in soil moisture content (~80 % WFPS) led to a significant reduction in HONO flux. Previous laboratory studies have also demonstrated that lower levels of HONO flux at high water holding capacity, low gas diffusion rates and high solubility could limit the release of HONO from soil (Ermel et al., 2018; Meusel et al., 2018; Wu et al., 2014). The observations before and after irrigation demonstrate the regulatory role of soil moisture in 265 the HONO exchange process, which has been systematically investigated by examining HONO emission flux as a function of soil moisture in previous studies (Mamtimin et al., 2016; Wang et al., 2021). Soil moisture determines whether nitrification or denitrification processes dominate gas emissions and strongly influences the corresponding gas emission rates and concentration compensation points (Cheng, 2013). Several laboratory findings indicate that nitrification under low soil moisture 270 conditions is the dominant process for HONO emissions (Oswald et al., 2013; Scharko et al., 2015), and field observations of HONO have predominantly focused on dryland ecosystems (Ren et al., 2011). Conversely, Wu et al. (2019) demonstrated that soil under high water content (75 %–140 % WHC) can also exhibit substantial emissions of HONO, with an average ratio of the highest HONO flux of wet peak to dry peak being approximately 30 %. However, actual field observations have revealed that 275 HONO fluxes are very low (close to 0) under high water content conditions, which may be attributed to the influence of soil moisture on microbial metabolic activity and gas diffusion in the soil (Hu et al., 2015; Linn and Doran, 1984). Furthermore, Wang et al. (2021) reported the promoting effect of fertilization on HONO flux under high soil moisture conditions (75 %–95 % WHC). Nevertheless, we did not observe this phenomenon in our field experiments with paddy fields. This discrepancy could 280 probably be attributed to the anaerobic or microaerobic conditions created by pre-fertilization irrigation, which exerted a greater inhibitory effect on the nitrification process than the promoting effect of fertilization. Currently, the estimation of HONO flux at the regional scale relies more on laboratory research findings (Gan et al., 2024; Wu et al., 2022). This study highlights discrepancies between laboratory and field observations within the high soil water content range, which pose significant 285 challenges to the uncertainty of estimation results.

The agricultural field acted as a well-known source of atmospheric NO, with an average flux of 0.19 ± 0.53 nmol m⁻² s⁻¹ in this study. Similar to HONO, the upward NO flux was mostly observed during rotary tillage, with a maximum flux of 9.12 nmol $m^2 s^{-1}$ in the early morning (Table 1). This finding is consistent with previous studies exhibiting that tillage increases NO emission (Chatskikh and 290 Olesen, 2007; Fang et al., 2006; Fang and Yujing, 2009; Liu et al., 2005; Pinto et al., 2004; Sehy et al., 2003; Yao et al., 2009; Yamulki and Jarvis, 2002). However, the NO fluxes were close to zero when the paddy field was waterlogged, probably because the nitrification process that dominates NO production in soil was greatly hindered in water-saturated soil and anoxic microsites (Fang and Yujing, 2009). Similarly, we also did not observe significant emissions of HONO under sustained high moisture 295 conditions. The coincidences of peaks in HONO flux and NO flux during rotary tillage suggest that HONO release from soil, similar to NO, is associated with microbial activity in soil (Bargsten et al., 2010; Skiba et al., 1993). Furthermore, the notably elevated fluxes of HONO and NO were observed during rotary tillage in comparison to other phases of agricultural activities (Fig. S3). The higher emission rates of NO and HONO could account for the successive peaks in their concentrations and 300 fluxes. Similar to NO, the emission of HONO from soil could be significantly stimulated by soil tillage. Besides, the average NO₂ flux of -0.42 \pm 0.44 nmol m⁻² s⁻¹ (ranging from -3.50 to 0.50 nmol m⁻² s⁻¹) indicated that the agricultural field acted as a sink for atmospheric $NO₂$ (Fang and Yujing, 2009; Tang et al., 2020).

Table 2 Summary of the maximum and minimum of HONO flux in field measurements over different soil types at

305 remote/rural/suburban sites.

AG: aerodynamic gradient; REA: relaxed eddy accumulation; OTDC: open-top dynamic chamber; HONO fluxa: values in the time series; HONO flux^b: values in the diurnal variations;

3.3 Diurnal profiles of fluxes and HONO source during rotary tillage

- The diurnal variations of NO² photolysis frequency (*J*(NO2)), air temperature, relative humidity, soil 310 temperature, WFPS, NO flux, NO² flux, and HONO flux are illustrated in Fig. 4. The diurnal HONO flux exhibited no discernible diurnal pattern during the whole campaign, which was similar to the diurnal profile observed during BEARPEX 2009 in California (Ren et al., 2011). Significant HONO emissions were primarily observed during rotary tillage, and its daily pattern is depicted in Fig. 4. Upward HONO fluxes were observed throughout the day, with a maximum value of 0.55 nmol $m² s⁻¹$ in the early morning. 315 The distinct HONO emissions were observed in the morning after sunrise. Moreover, the magnitudes of the daytime fluxes $(0.25 \pm 0.13 \text{ nmol m}^2 \text{ s}^{-1})$ were comparable to the nocturnal values $(0.27 \pm 0.13 \text{ nmol})$ $m⁻²$ s⁻¹). The diurnal profile of NO flux exhibited consistent levels of NO emission throughout the day, except for a noticeable peak in the early morning. It is worth noting that the synchronous peak of HONO flux and NO flux was observed in the morning. In contrast to the fluxes of HONO and NO, deposition 320 was the prevailing process for NO₂ flux. A greater downward NO₂ flux of -0.85 \pm 0.27 nmol m⁻² s⁻¹ (-
	- 0.57 ± 0.23 nmol m⁻² s⁻¹ at night) was observed during the daytime, potentially due to an increase in the dry deposition velocity of $NO₂$ during the day.

325 **Figure 4.** (a) Diurnal variations of NO² photolysis frequency (*J*(NO2)), air temperature, relative humidity, soil temperature and WFPS throughout the whole campaign. (b) Diurnal profiles of HONO, NO and NO² fluxes are presented for the whole campaign and rotary tillage. The error bars denoted the standard deviation.

Throughout the rotary tillage period, the emissions of HONO and NO were significant, with the maximum fluxes reaching 1.86 nmol $m^2 s^{-1}$ for HONO and 9.12 nmol $m^2 s^{-1}$ for NO. The concurrent 330 peaks in HONO and NO fluxes indicate that HONO emissions could originate from soil sources, as it is well-established that NO is primarily generated and released from soil microbial processes (Feig et al., 2008; Rende et al., 1989). As shown in Fig. 5, a significant correlation (*R* = 0.77) was observed between the fluxes of HONO and NO during the rotary tillage period, suggesting a shared source for both gases. This could be attributed to the generation and release from soil microbial processes, aligning with the

- 335 results reported by Tang et al. (2020). A gaussian fitting was employed to analyze the variation of HONO and NO fluxes with soil temperature (Fig. S4). It was found that both HONO and NO exhibited maximum emission fluxes at approximately 25 °C and 24 °C, respectively, which is close to the optimal temperature (25 °C) for soil microbial nitrification and denitrification processes (Agehara and Warncke, 2005; Fang and Yujing, 2009). This finding further supports the hypothesis that HONO is generated and released
- 340 from soil biological processes. Additionally, there was an indication of elevation in HONO flux during periods of intense solar radiation in the morning. Although the correlations of HONO flux with NO2, $J(NO₂)$ and $NO₂$ flux were found to be low ($R = 0.28$, 0.12 and 0.25), we observed a significant correlation ($R = 0.60$) between HONO flux and the product of $J(NO_2) \times NO_2$ (Fig. 6), as well as a moderate correlation ($R = 0.41$) with the product of $J(NO₂) \times NO₂$ flux (Fig. S5). This indicates that the
- 345 light-induced NO₂ conversion serves as an important source of HONO during the day. Furthermore, another mechanism of acid displacement can be ruled out, as the major strong acid, HNO₃, is primarily generated during the daytime and subsequently deposited to the ground. Consequently, the peak of the HONO source is expected to occur in the afternoon (Vandenboer et al., 2015). Finally, the results indicate that both mechanisms of soil release from biological processes and light-induced $NO₂$ conversion are 350 likely active, which together affect the diurnal HONO flux pattern.

Figure 5. Correlation of HONO flux with NO flux during rotary tillage.

Figure 6. Correlation of the daytime HONO flux with (a) NO₂, (b) $J(NO_2)$ and (c) the product of $J(NO_2) \times [NO_2]$ during rotary tillage.

360 **3.4 HONO budget during rotary tillage**

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In Section 3.3, we presented the potential sources of HONO flux during rotary tillage by conducting correlation analysis. Here, we will further calculate the specific contributions of HONO sources through budget analysis. The HONO budget can be derived from known HONO sources and sinks, and the potential unknown HONO source during rotary tillage was estimated. The lower-level data that better 365 describe ground source processes were used for budget analysis. In this study, the investigated HONO sources including homogeneous reaction ($P_{\text{OH+NO}}$), heterogeneous reaction of NO₂ on the aerosol surface and the ground surface (P_{aerosol} and P_{ground}). The HONO sinks included the reaction of HONO with OH

 $(L_{\text{OH+HONO}})$, photolysis of HONO (L_{photo}) and dry deposition loss of HONO (L_{den}). The calculation of HONO sources and sinks, as well as the estimates of the mixing layer height (MLH) is described in detail 370 in Text S3 in the Supplemental Material.

$$
\frac{d \text{HONO}}{dt} = \left(P_{\text{OH}+\text{NO}} + P_{\text{unknown}} + P_{\text{aerosol}} + P_{\text{ground}}\right) - \left(L_{\text{OH}+\text{HONO}} + L_{\text{photo}} + L_{\text{dep}}\right) \tag{4}
$$

Simplifying Eq. (4), the dHONO/dt is approximated by $\triangle HONO/\triangle t$. Then Eq. (4) is turned to Eq. (5):

$$
P_{\text{unknown}} = \frac{\triangle HONO}{\triangle t} + L_{\text{OH+HONO}} + L_{\text{photo}} + L_{\text{dep}} - P_{\text{OH+NO}} - P_{\text{aerosol}} - P_{\text{ground}} \tag{5}
$$

The average production and loss rates for the diurnal HONO budget are shown in Fig. 7. The 375 homogeneous reaction of NO and OH accounted for 12.8 % of HONO production, and an average of $P_{\text{OH+NO}}$ was 0.15 \pm 0.10 ppbv h⁻¹. The heterogeneous conversion of NO₂ on the ground surfaces accounted for 12.4 % (0.1 \pm 0.07 ppbv h⁻¹) of HONO production at night. P_{aerosol} (0.01 \pm 0.006 ppbv h⁻¹) was negligible compared to other HONO sources due to its relatively small aerosol surface area (Fig. S6). The photodecomposition (L_{photo}) was the primary sink of HONO during the daytime, with a peak of 380 2.03 ppbv h⁻¹ at 11:00 and an average of 1.44 \pm 0.69 ppbv h⁻¹, while $L_{\text{OH+HONO}}$ was very small and less than 5 % of L_{photo} . The dry deposition of HONO (L_{dep}) was influenced by the mixing layer height (MLH) and dominated the loss of HONO at night, with a rate exceeding 0.6 ppbv h^{-1} . $P_{unknown}$ exhibited the obvious diurnal variation, with higher values during daytime $(1.31 \pm 0.54$ ppbv h⁻¹) and lower values at night (0.53 \pm 0.25 ppbv h⁻¹). The peak of $P_{unknown}$ occurred during 7:00–12:00, with a maximum value of 385 2.18 ppb h⁻¹. The peak value of $P_{unknown}$ is comparable to that measured in Taizhou (2.5 ppbv h⁻¹) (Ye et al., 2023) and larger than in Wangdu $(0.62$ ppbv h⁻¹) (Song et al., 2022), Nanjing $(1.04$ ppbv h⁻¹) (Liu et al., 2019b). Similar to the observed asymmetry around noon reported in previous studies, this could be attributed to the combined effect of solar radiation and the variation of precursor $NO₂$ (Song et al., 2022; Xue et al., 2022). As the significantly larger unknown source strength of HONO during the daytime, we 390 focused on analyzing the unknown source of HONO during the day. Based on the above analysis, we evaluated the contribution of the photo-enhanced heterogeneous pathways.

The coefficients widely adopted in previous studies generally range from 10^{-6} to 10^{-4} (Chen et al., 2023; Liu et al., 2019c; Song et al., 2022; Wong et al., 2013). Here, we used 1×10^{-5} as the photoenhanced uptake coefficients $(\gamma_{a+h\nu})$ and $\gamma_{g+h\nu}$) to calculate the $P_{\text{aerosol}+hv}$ and $P_{\text{ground}+hv}$ (Qin et al., 395 2023; Xue et al., 2020). As shown in Fig. 8, the average $P_{\text{aerosol}+hv}$ and $P_{\text{ground}+hv}$ were 0.02 ± 0.009 ppbv h^{-1} and 0.53 ± 0.50 ppbv h^{-1} during the day, respectively, accounting for 1.4 % and 40.2 % of $P_{unknown}$,

and $P_{\text{aerosol}+hv}$ was a negligible source of daytime HONO formation. The photo-enhanced NO₂ heterogeneous reaction on the surfaces matched the calculated $P_{unknown}$ and well-explained the HONO budget in the morning. Furthermore, the higher photo-enhanced uptake coefficient of 3.5×10^{-5} was 400 adopted as the upper limit for calculating the production of photosensitive conversion of $NO₂$ (Chen et al., 2023). The calculation results demonstrated that the daytime $P_{unknown}$ could be explained when the upper limit of photo-enhanced uptake coefficient was used (Fig. S8).

However, there could be other light-driven reaction pathways to produce HONO in the afternoon, as indicated by the diurnal variation of $P_{unknown}$. Previous studies have demonstrated that the photolysis 405 of pNO3/HNO³ can contribute to HONO production (Chen et al., 2023; Laufs et al., 2017). Recently, Chen et al. (2023) found that the photolysis of $HNO₃$ at the surface interface could well explain the observed $P_{unknown}$ in the afternoon. However, the lack of information about $HNO₃$ concentration does not allow us to directly estimate the contribution of $HNO₃$ photolysis in the present study. Future studies should supplement measurements of $pNO₃/HNO₃$ to better characterize the contribution of this

410 potentially important HONO formation pathway.

Based on the measured fluxes, we also estimated the HONO emission rate from soil (P_{soil}) . The nighttime HONO fluxes ranged from 0.15 nmol $m^2 s^{-1}$ to 0.43 nmol $m^2 s^{-1}$, with corresponding HONO flux rates of 0.32 ppbv h⁻¹ to 0.79 ppbv h⁻¹, which were fully capable to explaining $P_{unknown}$ (ranging from 0.012 to 0.90 ppbv h⁻¹). Therefore, the light-induced HONO sources (photosensitive conversion of NO₂ 415 and photolysis of pNO3/HNO3) and soil emissions could serve together as significant HONO sources in agricultural fields, thereby influencing the overall atmospheric HONO budget.

Figure 7. Diurnal variation of HONO budget during rotary tillage.

420 **Figure 8.** Diurnal variation of light-induced conversion of NO² and HONO flux rate derived from soil emission.

3.5 Implication on the atmospheric oxidizing capacity

The significant increase in atmospheric HONO from agricultural fields can enhance the formation of OH radicals via its photolysis (see the detailed OH production rate calculation in Text S4). Figure 9a exhibited the OH production rates from the photolysis of HONO $(P(OH)_{HONO})$ and $O_3 (P(OH)_{O_3})$. The

- 425 $P(\text{OH})_{\text{HONO}}$ and $P(\text{OH})_{\text{O}_3}$ were found to be 0.82 ppbv h⁻¹ and 1.49 ppbv h⁻¹, respectively, which were significantly higher than the corresponding winter levels at the same site (Fig. S9). The higher $O₃$ concentration in summer plays a primary role in the generation of OH radicals by daytime O_3 photolysis, accounting for 70 % of the total OH production rate. However, the contribution of $P(OH)_{HONO}$, approximately 30 %, is still significant and cannot be ignored.
- 430 During the rotary tillage period, continuous peaks in HONO concentration and flux were observed, with maximum values of 3.06 ppbv and 1.86 nmol m⁻² s⁻¹, respectively. $P(OH)_{HONO}$ and $P(OH)_{O_3}$ were calculated to be 1.42 ppbv h^{-1} and 1.35 ppbv h^{-1} , respectively, accounting for 51 % and 49 % of the total OH production rate (Fig. 9b). $P(OH)_{HONO}$ dominated in the early morning with a value of 2.48 ppbv h⁻¹, while $P(\text{OH})_{\text{O}_3}$ became the main source at midday with a value of 2.74 ppbv h⁻¹. The comparable peak
- 435 magnitude of $P(OH)_{HONO}$ and $P(OH)_{O_3}$ indicates that HONO photolysis is an important source of daytime OH radicals. Furthermore, the peaks of both $P(OH)_{HONO}$ and HONO flux co-occur in the early morning, revealing the significant contribution of agricultural HONO emissions to the regional atmospheric oxidation capacity in the Huaihe River Basin.

Figure 9. Diurnal variation of net OH production rate of the photolysis of HONO $(P(OH)_{HONO})$ and O₃ $(P(OH)_{O_3})$ during (a) the whole campaign and (b) the rotary tillage.

4. Conclusion

The extensive agricultural fields and increased agricultural activities have contributed to certain 445 areas in China becoming hotspots for atmospheric nitrogen oxides, underscoring the increasing importance of regional and global nitrogen budgets. However, the available HONO emission fluxes from agricultural soils are relatively limited. In this study, we utilized the AG method to measure the HONO and NO_x fluxes from agricultural fields in the Huaihe River Basin. For HONO and NO, upward fluxes of 0.07 ± 0.22 and 0.19 ± 0.53 nmol m⁻² s⁻¹ were observed, respectively, while NO₂ exhibited a deposition 450 flux to the ground of -0.42 \pm 0.44 nmol m⁻² s⁻¹. The successive peaks in HONO flux and NO flux were measured during rotary tillage, suggesting a potentially enhanced release of HONO and NO due to soil tillage activities. However, the higher WFPS inhibited the microbial nitrification processes after irrigation, leading to a significant decrease in HONO and NO fluxes. Under this inhibitory effect, no significant peaks in HONO flux were observed after fertilization compared with that during the rotary 455 tillage. Considering limited field observation of HONO flux under high soil water content, future studies

should pay more attention to paddy fields to validate the mechanisms observed in the laboratory.

Significant fluxes were observed during rotary tillage, prompting the investigation into the HONO sources and budget during this period. Biological processes and light-driven NO₂ reactions on the ground surface may both be sources of HONO and influence the local HONO budget. Higher levels of

 $P(OH)_{HONO}$ were observed in the early morning, consistent with the peak emission flux of soil HONO. This reveals the significant contribution of agricultural HONO emissions to the regional atmospheric oxidation capacity in the Huaihe River Basin. Overall, this study provides valuable insights into the dynamics of soil HONO emissions in agricultural fields, elucidating their environmental implications and the role of agricultural activities in the atmospheric chemistry of HONO.

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Data availability. The data described in this manuscript are available at <https://doi.org/10.5281/zenodo.12738765> (Meng et al., 2024) or upon request from the corresponding author [\(mqin@aiofm.ac.cn\)](mailto:mqin@aiofm.ac.cn).

470 *Supplement.* The supplement related to this article is available online at:

Author contributions. MQ and PX designed the experiments. FM, KT, DS, ZL and JD performed the measurements. FM and BH analyzed the data and wrote the manuscript. MQ revised and commented on the paper. YF, YH and TN provided the ancillary data and experimental sites.

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Competing interests. The authors declare that they have no conflict of interest.

Acknowledgements. This work was supported by the National Key Research and Development Program of China (2022YFC3701103), the National Natural Science Foundation of China (U21A2028, 41875154),

480 the Plan for Anhui Major Provincial Science & Technology Project (202203a07020003).

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