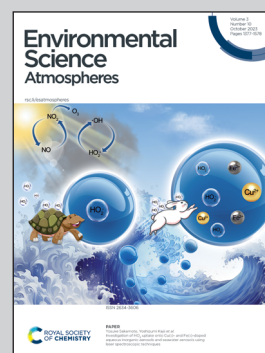


Showcasing research from Professor Tianlei Zhang's laboratory, School of Chemistry & Environment Science, Shaanxi University of Technology, Hanzhong, China.

A possible atmospheric source of HNO_3 : the ammonolysis reaction of $t\text{-N}_2\text{O}_4$ in the presence of water monomer, water dimer, and sulfuric acid

The effect of H_2O , $(\text{H}_2\text{O})_2$ and H_2SO_4 on the ammonolysis of $t\text{-N}_2\text{O}_4$ to form HNO_3 was studied by a quantum chemical method and Master equation rate calculations. Results reveal that the ammonolysis of $t\text{-N}_2\text{O}_4$ with $(\text{H}_2\text{O})_2$ and H_2SO_4 are barrierless or nearly barrierless reactions. Considering the effective rate constant, $(\text{H}_2\text{O})_2$ outperforms the other catalysts in the range of 280-320 K (0 km). Moreover, the effect of H_2SO_4 is obvious at higher altitudes of 5-30 km. In general, this work will give new insights into how the neutral and acidic catalysts affect the formation of HNO_3 .

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A possible atmospheric source of HNO₃: the ammonolysis reaction of *t*-N₂O₄ in the presence of water monomer, water dimer, and sulfuric acid†

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Although the ammonolysis of *t*-N₂O₄ is one of the potential sources of HNO₃ formation, the available studies have only focused on its naked reaction. Herein, the effect of important neutral and acidic trace gases, water monomer, water dimer, and sulfuric acid, on the formation of HNO₃ from the ammonolysis of *t*-N₂O₄ was studied by the quantum chemical method of CCSD(T)/aug-cc-pVTZ//B3LYP-D3/6-311++G(3df,2pd) and the Master equation/Rice–Ramsperger–Kassel–Marcus (ME/RRKM) rate calculations. The quantum chemical results revealed that the ammonolysis of *t*-N₂O₄ with (H₂O)₂ and H₂SO₄ are barrierless or nearly barrierless reactions, potentially lowering the energy barrier to 3.4–4.1 kcal mol⁻¹. The calculated effective rate constant illustrates that (H₂O)₂ (100% RH) dominates over H₂O and H₂SO₄ within the range of 280–320 K (0 km), with an effective rate constant that is 1–3 orders of larger magnitude, whereas H₂SO₄ (10⁸ mol cm⁻³) is the most favorable catalyst within the troposphere between 5 and 30 km. However, the contributions of H₂O, (H₂O)₂, and H₂SO₄ are not apparent in the gas-phase ammonolysis of *t*-N₂O₄ within the range of 213–320 K and 0–30 km because their effective rate constants were at least 4 orders of magnitude lower than the corresponding rate constant of the ammonolysis of *t*-N₂O₄. In general, the current findings shed fresh light on neutral (H₂O and (H₂O)₂) and acidic (H₂SO₄) catalysts that not only affect energy barriers but also have an impact on the ammonolysis of *t*-N₂O₄ in neutral and acidic conditions.

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Environmental significance

Nitrogen tetroxide (N₂O₄) is considered to be a dimer of nitrogen dioxide (NO₂) and plays an important role in the formation of acid rain. The fact is that the ammonolysis of *t*-N₂O₄ is one of the potential sources of HNO₃ formation; thus, the effect of water monomer, water dimer, and sulfuric acid on the ammonolysis of *t*-N₂O₄ was studied by quantum chemical method and Master equation rate calculations. The quantum chemical results reveal that the ammonolysis of *t*-N₂O₄ with (H₂O)₂ and H₂SO₄ are barrierless or nearly barrierless reactions. In terms of the effective rate constant, (H₂O)₂ outperforms the other catalysts in the temperature range 280–320 K (0 km). Moreover, the effect of H₂SO₄ on the ammonolysis reaction of *t*-N₂O₄ is obvious at higher altitudes of 5–30 km. In general, this work will give a new insights into how the neutral and acidic catalysts affect the formation of HNO₃.

1. Introduction

Nitrogen dioxide (NO₂),¹ as one of the most significant NO_x, is not only a precursor to the photochemical formation of ozone in the troposphere,² but it can also contribute to the formation of photochemical smog and cause significant health and

environmental hazards.³ Nitrogen tetroxide (N₂O₄), a dimer of nitrogen dioxide (NO₂),⁴ can be used for nitration, nitrosation, and oxidation.⁵ In addition, N₂O₄ is an oxidizing agent for auto-igniting fuels and plays an important role in the formation of acid rain.⁶ Due to the fact that N₂O₄ is a highly toxic chemical species that hinders experimental studies,⁷ quantum calculations are a trend in current research to probe the N₂O₄-related reaction mechanisms. As the most major loss route of N₂O₄ in the atmosphere, the hydrolysis of N₂O₄ is potentially important in the lower atmosphere and plays an important role in the formation of HONO, a major source of OH pollution in the urban atmosphere.^{8,9}

Several investigations have shown that the less stable *t*-N₂O₄ (*trans*-N₂O₄) is substantially more reactive than *s*-ONONO₂ (symmetric N₂O₄); hence, it was selected as a starting point for

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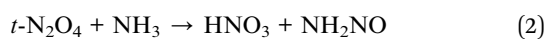
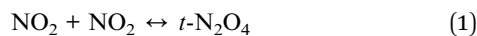
† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d3ea00095h>

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studying the hydrolysis of NO₂ dimers.^{9–12} The reaction barrier for the hydrolysis of *t*-N₂O₄ to form HONO was in the range of 10.8–11.9 kcal mol⁻¹ at different theoretical levels.^{9,11,12} Several groups reported that using H₂O and (H₂O)₂ as catalysts stabilized the reactant complexes by 3.5–6.9 kcal mol⁻¹ and reduced the energy barrier to 6.2 kcal mol⁻¹. Consequently, Zhang *et al.*⁹ revealed that, for the hydrolysis of *t*-N₂O₄, the H₂SO₄ catalyst is more effective than H₂O and (H₂O)₂ catalysts, resulting in not only a higher binding energy of 15.0 kcal mol⁻¹ for the reactant complex but also a lower energy barrier of 3.8 kcal mol⁻¹.

In the atmosphere, the hydrolysis of *t*-N₂O₄ to produce HONO is the most major loss route of *t*-N₂O₄.^{10,11} As a complement to the loss of *t*-N₂O₄, the ammonolysis reaction of *t*-N₂O₄ can form HNO₃,¹³ which could be competitive with the main source of HNO₃, the gas-phase reaction of NO₂ with the hydroxyl (OH) radical^{18,14} during the day and the hydrolysis reaction of N₂O₅^{15–17} at night, in polluted areas of NH₃. The ammonolysis of *t*-N₂O₄ (shown in eqn (1) and (2)) investigated by Lin *et al.*¹³ reveals that the energy barrier of the *t*-N₂O₄ + NH₃ reaction was determined to be 5.3 kcal mol⁻¹ and the corresponding rate constant at low temperature was not pressure-dependent. As far as we know, the effect of neutral and acidic gases on the ammonolysis of *t*-N₂O₄, which plays a significant catalytic role in hydrogen abstraction reactions,^{9,15,18–37} has not been explored.



As in the previous studies on N₂O₄ + H₂O,^{11,12} HO₂ + NO₂,³⁸ H₂CO + NH₃,³⁹ and SO₂ + NO₂⁴⁰ reactions, water molecules were found to play an essential role in enhancing the stability of pre-reactive complexes and lowering the apparent activation energies of the transition states. In addition to water monomer, some recent works addressed the potential role of water dimer,^{18,41} which may play a significant catalytic role in hydrogen abstraction reactions because its concentrations can reach 9×10^{14} mol cm⁻³.^{42,43} Aside from water monomer and water dimer, acidic^{9,15,30–37} gas species in the atmosphere may also be effective in lowering the energy barriers for hydrogen transfer reactions^{30–34} and atmospheric hydrolysis reactions^{9,15,35–37} in the gas phase. The presence of H₂SO₄^{37,44} in the atmosphere was considered to be a more effective catalyst than neutral catalysts,^{25,28–30,33,45} which not only greatly reduces the energy barriers^{27,28,33} but also facilitates the transfer of hydrogen,^{30–33,46} and thus, H₂SO₄ was regarded as either good acceptors or good donors of H in the catalytic gas reactions. These situations stimulated our interest in studying the ammonolysis of *t*-N₂O₄ by neutral (H₂O and (H₂O)₂) and acidic (H₂SO₄) gases.

In this work, using global minimum searching combined with quantum chemical methods, we first obtained the stable structures of the reactant complexes of *t*-N₂O₄⋯NH₃⋯X (X = H₂O, (H₂O)₂, and H₂SO₄). The ammonolysis of *t*-N₂O₄ in the presence of X was then studied using the stable molecular clusters *t*-N₂O₄⋯NH₃⋯X. Finally, the effective rate constant for

the ammonolysis of *t*-N₂O₄ with X was estimated at temperatures ranging from 213 to 320 K and altitudes ranging from 0 to 30 km.

2. Calculation details

2.1 Electronic structure calculations

The molecular geometries of the isolated reactants, pre-reactive complexes, transition states, post-reactive complexes, and products of the ammonolysis reaction of *t*-N₂O₄ without and with X were optimized using the B3LYP-D3 method^{47–53} with 6-311++G(3df,2pd) basis set in Gaussian 09 suites.⁵⁴ The D3 method has been reliably utilized to describe the noncovalent interaction and the equilibrium structure of atmospheric clusters and reactions.^{51,55} Notably, the keyword “stable = opt” is added in the calculations at the B3LYP-D3/6-311++G(3df,2pd) level to ensure that all the geometries are stable. Frequency calculations were calculated at the same level for all stationary points to check that all transition states have the same character as a first-order saddle point with a single imaginary frequency and that other stationary points correspond to minima on potential energy surfaces (PESs). The scaling factor employed to adjust the ZPEs was 0.969.^{56,57} To ensure that the optimized transition state connects the desired pre- and post-reactive complexes, intrinsic reaction coordinate (IRC) calculations^{58–60} were performed at the same level. To improve the accuracy of the relative energies, single-point energy calculation was performed at the CCSD(T)/aug-cc-pVTZ^{61–63} level by using the Gaussian 09 software.⁵⁴ It was noted that the *T*₁ diagnostic values for closed-shell in Table S3† were 0.02 less than the standard value,^{64,65} showing the multireference calculations for recovering non-dynamical correlation were not a problem, and the single reference method of CCSD(T)/aug-cc-pVTZ is reliable to single point energy calculation.

Global minimum searching combined with quantum chemical methods was employed to obtain the most stable structures of the reactant complexes of *t*-N₂O₄⋯NH₃⋯X. Initially, 500 structures with low energies were auto-produced by ABCluster software^{66,67} with TIP4P^{68,69} model for H₂O, (H₂O)₂, and CHARMM⁷⁰ force field for *t*-N₂O₄, NH₃, and H₂SO₄. Then, pre-optimized by the semi-empirical method of PM7⁷¹ in MOPAC 2016.⁷² Next, the structures with the N(*t*-N₂O₄)⋯N(NH₃) interaction of electron donor–acceptor (EDA) and facilitating the transfer of hydrogen atom from *t*-N₂O₄ to NH₃ were selected to optimize at the B3LYP-D3/6-311+G(d,p) level. Subsequently, 50 isomers with an order of electronic energies were chosen to optimize at the level of B3LYP-D3/6-311+G(2d,2p). Finally, the global minimum isomers within 5.0 kcal mol⁻¹ (the electric energy) were re-optimized at the B3LYP-D3/6-311++G(3df,2pd) level.

2.2 Rate constant calculations

The rate constants for the ammonolysis reaction of *t*-N₂O₄ without and with X were calculated in two steps. First, the high-pressure-limit (HPL) rate constants were calculated by using the VRC-VTST calculations in Polyrate 2017 software.⁷³ The details



of VRC-VTST can be seen in Table S7 of the ESI.† Then, based on the HPL rate constants, the rate constants for the ammonolysis reaction of $t\text{-N}_2\text{O}_4$ without and with X at different temperatures and pressures were calculated using the Master Equation Solver for Multi-Energy Well Reactions (MESMER) program.⁷⁴ The Inverse Laplace Transform (ILT) approach was used to analyze the barrierless step from distinct reactants to the pre-reactive complex.^{75,76} Meanwhile, the transition step from the pre-reactive complex to the post-reactive complex occurring through the transition state was applied to the RRKM theory.^{77,78} ILT methods and RRKM theory can be, respectively, expressed in eqn (3) and (4).

$$k(E) = \frac{W(E - E_0)}{h\rho(E)} \quad (3)$$

$$k^\infty(\beta) = \frac{1}{Q(\beta)} \int_0^\infty k(E)\rho(E)\exp(-\beta E)dE \quad (4)$$

In eqn (3) and (4), where $W(E - E_0)$ is the rovibrational sum of states (SOS) at the optimized transition state (TS) geometry, E_0 is the reaction threshold energy, h is Planck's constant, $\rho(E)$ is the density of rovibrational states of the reactant, and $Q(\beta)$ is the corresponding canonical partition function. Moreover, the electronic geometries, vibrational frequencies, and rotational constants were calculated at the B3LYP-D3/6-311++G(3df,2pd) level, and single-point energy calculations were refined at the CCSD(T)/aug-cc-pVTZ level for the modeling. The one-dimensional asymmetric Eckart potential was used to treat the tunneling effect in the RRKM calculation. In addition, the Lennard-Jones (L-J) parameters $\epsilon/k_B = 71.4$ K and $\sigma = 3.798$ Å were used for N_2 ,⁷⁹ $\epsilon/k_B = 200.0$ K and $\sigma = 3.50$ Å were used for $t\text{-N}_2\text{O}_4$,⁸⁰ while $\epsilon/k_B = 481.0$ K and $\sigma = 2.92$ Å were estimated for NH_3 .⁸⁰

3. Results and discussions

The pre-reactive complex in each reaction channel was denoted by “IM” followed by a number, whereas the transition state and post-reactive complexes were denoted by “TS” and “IMF”, respectively. Species in the presence of H_2O , $(\text{H}_2\text{O})_2$, and H_2SO_4 were denoted by the suffixes “WM”, “WD”, and “SA”.

3.1 Mechanism and rate constants for the ammonolysis reaction of $t\text{-N}_2\text{O}_4$

The ammonolysis reaction of $t\text{-N}_2\text{O}_4$ has been extensively investigated from the theoretical viewpoint.¹³ Here, this reaction has been reinvestigated at the CCSD(T)/aug-cc-pVTZ//B3LYP-D3/6-311++G(3df,2pd) level to check the catalytic effect of X. Our results shown in Fig. 1 were found to be very mechanistically and energetically similar to the work reported by Lin *et al.*¹³ All the relative energy values qualitatively matched (see Table S2†). As seen in Fig. 1, the $t\text{-N}_2\text{O}_4 + \text{NH}_3$ reaction occurred through a ring formation mechanism, resulting in the formation of a six-membered ring complex $t\text{-N}_2\text{O}_4 \cdots \text{NH}_3$ with a binding energy of $0.5 \text{ kcal mol}^{-1}$. Then, the terminal O1 atom of $t\text{-N}_2\text{O}_4$ abstracts the H atom of NH_3 along with the N-N bond formation to form the product complex $\text{HNO}_3 \cdots \text{NH}_2\text{NO}$. From an energy point of view, the barrier height of the $t\text{-N}_2\text{O}_4 + \text{NH}_3$ reaction was $3.7 \text{ kcal mol}^{-1}$, revealing that $t\text{-N}_2\text{O}_4$ can easily react with NH_3 in the gas phase.

The calculated rate constants for the ammonolysis reaction of $t\text{-N}_2\text{O}_4$ are listed in Table 1. In the ammonolysis reaction of $t\text{-N}_2\text{O}_4$, the hindered internal rotation (HIR)^{77,81–86} correction at 760 Torr has a moderate effect, increasing the rate constants by a factor of 1.25 to 1.28. Meanwhile, the almost unchanged rate constants for the ammonolysis reaction of $t\text{-N}_2\text{O}_4$ at different atmospheric pressures revealed that the pressure (10–760 Torr)⁸⁷ has little effect on the ammonolysis reaction of $t\text{-N}_2\text{O}_4$ within the temperature range of 280–320 K.^{88,89}

3.2 Mechanism and rate constants for the ammonolysis reaction of $t\text{-N}_2\text{O}_4$ with H_2O and $(\text{H}_2\text{O})_2$

Fig. 2a and b show the ammonolysis reaction of $t\text{-N}_2\text{O}_4$ assisted by H_2O (Channel WM) and $(\text{H}_2\text{O})_2$ (Channel WD), where both H_2O and $(\text{H}_2\text{O})_2$ served as a “bridge” to promote hydrogen atom transfer from the N3 atom of NH_3 to the terminal O1 atom of $t\text{-N}_2\text{O}_4$. In the case of Channel WM, the reaction can occur either (a) between NH_3 and monohydrated $t\text{-N}_2\text{O}_4$ ($t\text{-N}_2\text{O}_4 \cdots \text{H}_2\text{O}$) or (b) between hydrated NH_3 ($\text{NH}_3 \cdots \text{H}_2\text{O}$) and $t\text{-N}_2\text{O}_4$. The binding energy of $t\text{-N}_2\text{O}_4 \cdots \text{H}_2\text{O}$ was $2.9 \text{ kcal mol}^{-1}$, which was consistent with the previously calculated value of $3.5 \text{ kcal mol}^{-1}$ at the CCSD(T)-F12a/cc-pVDZ-F12//M06-2X/6-311++G(3df,2pd) level.⁹

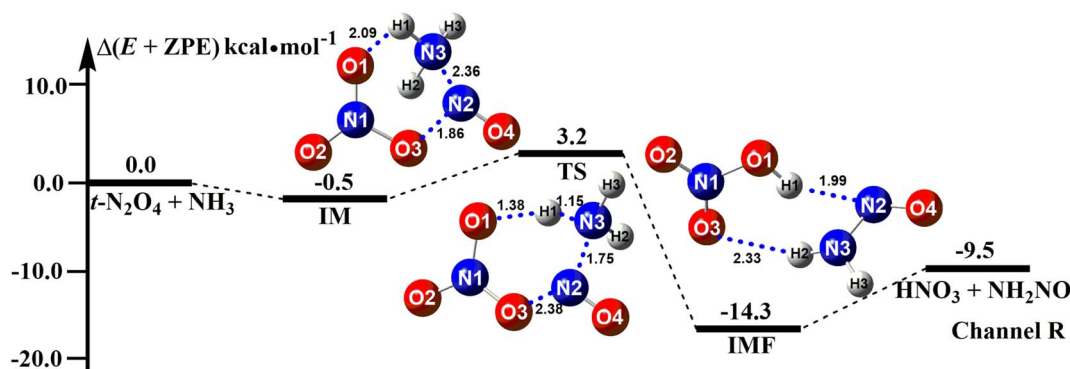


Fig. 1 Potential energy profiles for the $t\text{-N}_2\text{O}_4 + \text{NH}_3 \rightarrow \text{HNO}_3 + \text{NH}_2\text{NO}$ reaction at the CCSD(T)/aug-cc-pVTZ//B3LYP-D3/6-311++G(3df,2pd) level.



Table 1 The rate constants (k_R) ($\text{cm}^3 \text{mol}^{-1} \text{s}^{-1}$) for the $t\text{-N}_2\text{O}_4 + \text{NH}_3$ reaction without and with HIR treatments within the temperature range of 280–320 K and pressure range of 10–760 Torr

T (K)	HIR impact ^a			Pressure impact				
	With HIR treatments	Without HIR treatments	Factor ^b	10 Torr	50 Torr	100 Torr	300 Torr	760 Torr
280	6.67×10^{-17}	5.20×10^{-17}	1.28	5.03×10^{-17}	5.14×10^{-17}	5.16×10^{-17}	5.18×10^{-17}	5.20×10^{-17}
290	7.81×10^{-17}	6.13×10^{-17}	1.27	5.90×10^{-17}	6.07×10^{-17}	6.10×10^{-17}	6.12×10^{-17}	6.13×10^{-17}
298	8.80×10^{-17}	6.95×10^{-17}	1.27	6.65×10^{-17}	6.88×10^{-17}	6.92×10^{-17}	6.94×10^{-17}	6.95×10^{-17}
300	9.06×10^{-17}	7.17×10^{-17}	1.26	6.84×10^{-17}	7.09×10^{-17}	7.13×10^{-17}	7.16×10^{-17}	7.17×10^{-17}
310	1.04×10^{-16}	8.32×10^{-17}	1.26	7.84×10^{-17}	8.20×10^{-17}	8.26×10^{-17}	8.30×10^{-17}	8.32×10^{-17}
320	1.19×10^{-16}	9.57×10^{-17}	1.25	9.40×10^{-17}	9.40×10^{-17}	9.55×10^{-17}	9.55×10^{-17}	9.57×10^{-17}

^a HIR impact represents the hindered internal rotations treatment. ^b Factor denotes the rate ratio between with HIR treatments and without HIR treatments.

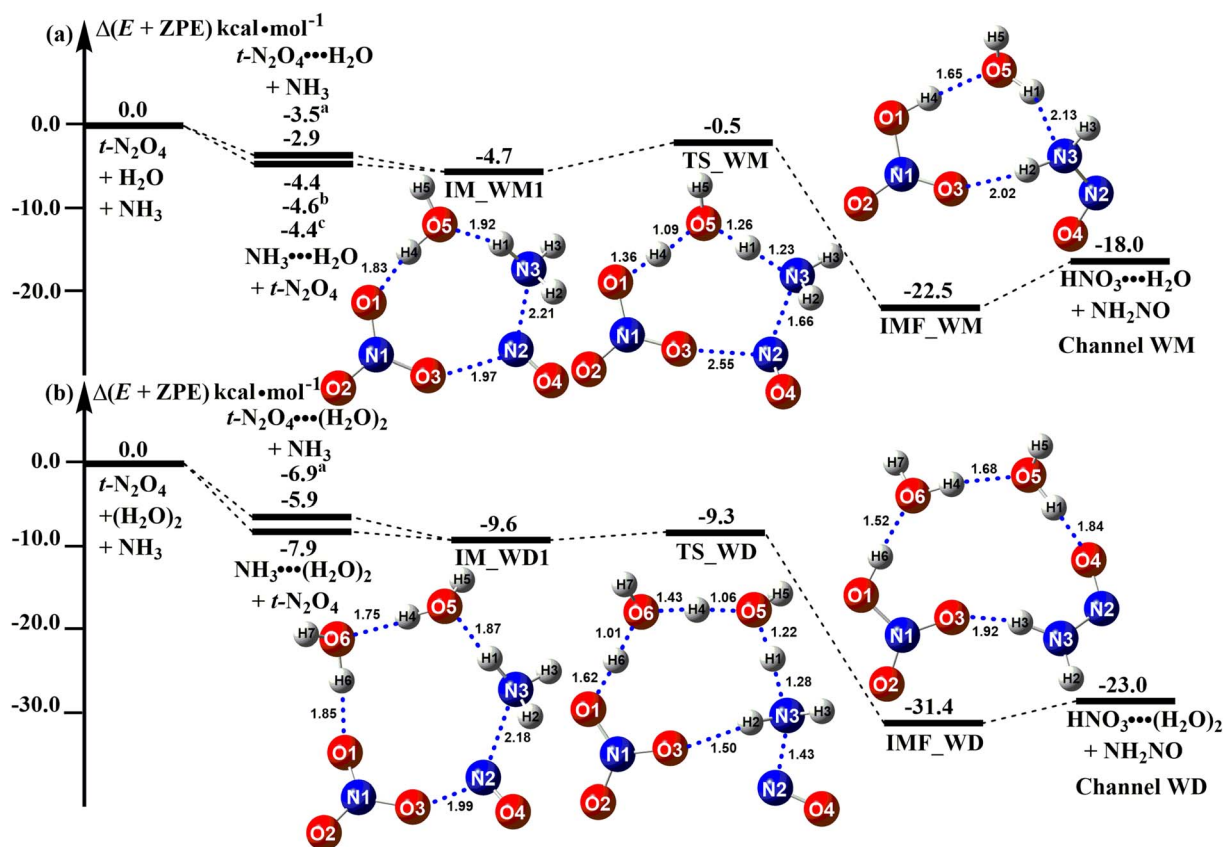


Fig. 2 Potential energy profiles for the $t\text{-N}_2\text{O}_4 + \text{NH}_3 \rightarrow \text{HNO}_3 + \text{NH}_2\text{NO}$ reaction catalyzed by H_2O and $(\text{H}_2\text{O})_2$ at the CCSD(T)/aug-cc-pVTZ//B3LYP-D3/6-311++G(3df,2pd) level (a–c) denotes the values respectively reported from ref. 9, 90, and 91.

The binding energy of $\text{NH}_3 \cdots \text{H}_2\text{O}$ was 4.4 kcal mol^{-1} , which agreed well with the calculated values of 4.4–4.6 kcal mol^{-1} .^{90,91} The stability of $\text{NH}_3 \cdots \text{H}_2\text{O}$ was 1.5 kcal mol^{-1} higher than that of $t\text{-N}_2\text{O}_4 \cdots \text{H}_2\text{O}$. So, the ammonolysis of $t\text{-N}_2\text{O}_4$ in the presence of H_2O mainly takes place *via* the collision of $\text{NH}_3 \cdots \text{H}_2\text{O}$ with $t\text{-N}_2\text{O}_4$ to form the quasi-planar eight-membered ring reactant complex IM_WM1. The energy of the reactant complex IM_WM1 was 4.7 kcal mol^{-1} lower than that of the isolated reactants $t\text{-N}_2\text{O}_4 + \text{H}_2\text{O} + \text{NH}_3$. In the complex IM_WM1, H_2O played the roles of a single acceptor and donor of hydrogen

bonds. After the formation of the complex IM_WM1, the reaction proceeded to form a hydrogen-bonded complex, $\text{HNO}_3 \cdots \text{H}_2\text{O} \cdots \text{NH}_2\text{NO}$ (denoted IMF_WM), through the transition state TS_WM, with an energy barrier of 4.2 kcal mol^{-1} .

The reaction $t\text{-N}_2\text{O}_4 + \text{NH}_3 + (\text{H}_2\text{O})_2$ can be initiated by the reactant complex IM_WD1, which can be formed from $t\text{-N}_2\text{O}_4 \cdots (\text{H}_2\text{O})_2 + \text{NH}_3$ or $t\text{-N}_2\text{O}_4 + \text{NH}_3 \cdots (\text{H}_2\text{O})_2$. It is clear from Fig. 2b that NH_3 was most likely bound to $(\text{H}_2\text{O})_2$ prior to $t\text{-N}_2\text{O}_4$. The reactant complex IM_WD1 has a quasi-planar structure similar to that of complex IM_WM1 and can be regarded as H_2O in



IM_WM1, which was replaced by $(\text{H}_2\text{O})_2$. The binding energy of IM_WD1 was $1.7 \text{ kcal mol}^{-1}$ from $t\text{-N}_2\text{O}_4 + \text{NH}_3 \cdots (\text{H}_2\text{O})_2$. After complex IM_WD1, the ammonolysis reaction of $t\text{-N}_2\text{O}_4$ with $(\text{H}_2\text{O})_2$ can form the product complex $\text{HNO}_3 \cdots (\text{H}_2\text{O})_2 \cdots \text{NH}_2\text{NO}$ (labeled as IMF_WD) through the transition state TS_WD with an energy barrier of $0.3 \text{ kcal mol}^{-1}$. Three hydrogen atom transfer mechanisms occurred at TS_WD, as well as the simultaneous formation of the N(2)–N(3) bond. In comparison to H_2O in Channel WM, $(\text{H}_2\text{O})_2$ in Channel WD played a more obvious catalytic role in promoting the ammonolysis reaction of $t\text{-N}_2\text{O}_4$. When $(\text{H}_2\text{O})_2$ was used as a catalyst in Channel WD, it stabilized the reactant complex by a further $4.9 \text{ kcal mol}^{-1}$ and decreased the reaction barrier by $3.9 \text{ kcal mol}^{-1}$. The more pronounced catalytic effect of $(\text{H}_2\text{O})_2$ could be attributed to two factors. On the one hand, $(\text{H}_2\text{O})_2$ may improve the $\text{N}(t\text{-N}_2\text{O}_4) \cdots \text{N}(\text{NH}_3)$ interaction compared to H_2O . For example, in the reactant complex IM_WD1, the strengthening of the $\text{N}(t\text{-N}_2\text{O}_4) \cdots \text{N}(\text{NH}_3)$ interaction was shown by the shortening of the bond distance $\text{N}(2) \cdots \text{N}(3)$ (2.18 \AA , shown in Fig. 2b), which is less than the corresponding value in the reactant complex IM_WM1 (2.21 \AA , shown in Fig. 2a). On the other hand, when H_2O was replaced by $(\text{H}_2\text{O})_2$, however, the transition state extended from an eight-member ring (TS_WM) to a ten-member ring (TS_WD). This structural change reduces the ring tension of the transition state to a certain extent, lowering the reaction energy barrier.

As shown in Table 2, within the range of 213–320 K and 0–30 km,^{88,89} the rate constant for the ammonolysis reaction of $t\text{-N}_2\text{O}_4$ assisted by H_2O (k_{WM}) was predicted to be 3.94×10^{-20} to $1.93 \times 10^{-19} \text{ cm}^3 \text{ mol}^{-1} \text{ s}^{-1}$, which was 2–3 orders of magnitude lower than that of the naked ammonolysis reaction of $t\text{-N}_2\text{O}_4$. The calculated rate constant for the ammonolysis reaction of $t\text{-N}_2\text{O}_4$ assisted by $(\text{H}_2\text{O})_2$ (k_{WD}) was 3.66×10^{-16} to $1.98 \times 10^{-15} \text{ cm}^3 \text{ mol}^{-1} \text{ s}^{-1}$, which was 1–2 orders of magnitude greater than the naked ammonolysis reaction of $t\text{-N}_2\text{O}_4$. The ammonolysis reaction of $t\text{-N}_2\text{O}_4$ assisted by $(\text{H}_2\text{O})_2$ was more kinetically favorable than the ammonolysis reaction of $t\text{-N}_2\text{O}_4$ with H_2O , with a rate constant that was 3–5 orders of magnitude greater.

3.3 Mechanism and rate constants for the ammonolysis reaction of $t\text{-N}_2\text{O}_4$ with H_2SO_4

As shown in Fig. S3,[†] nine geometrical isomers of the reactant complex $t\text{-N}_2\text{O}_4 \cdots \text{NH}_3 \cdots \text{H}_2\text{SO}_4$ (labeled as IM_SAn, $n = 1-9$) were found at the B3LYP-D3/6-311++G(3df,2pd) level, with complex IM_SA1 being the most stable. Based on complex IM_SA1, Fig. 3 presents the potential energy surface (PES) for the ammonolysis of $t\text{-N}_2\text{O}_4$ in the presence of H_2SO_4 (Channel SA). In the case of Channel SA, the reaction can occur (a) between NH_3 and $t\text{-N}_2\text{O}_4 \cdots \text{H}_2\text{SO}_4$ or (b) between $t\text{-N}_2\text{O}_4$ and $\text{NH}_3 \cdots \text{H}_2\text{SO}_4$. The binding energy of $\text{NH}_3 \cdots \text{H}_2\text{SO}_4$ was $14.7 \text{ kcal mol}^{-1}$, which was in good agreement with the previously reported value.⁹² The large binding energy of $\text{NH}_3 \cdots \text{H}_2\text{SO}_4$ indicates that in the reaction of $t\text{-N}_2\text{O}_4 + \text{NH}_3 + \text{H}_2\text{SO}_4$, NH_3 is easily bound to the isolated H_2SO_4 . In this sense, the ammonolysis reaction of $t\text{-N}_2\text{O}_4$ with H_2SO_4 mainly takes place *via* the collision of $t\text{-N}_2\text{O}_4$ with $\text{NH}_3 \cdots \text{H}_2\text{SO}_4$, resulting in the reactant complex IM_SA1.

The energy of the reactant complex IM_SA1 was $15.0 \text{ kcal mol}^{-1}$ lower than that of the separate reactants $t\text{-N}_2\text{O}_4$, NH_3 , and H_2SO_4 . In the complex IM_SA1, H_2SO_4 served as a single donor and acceptor of hydrogen bonds to form a ring-like structure with the binary complex of $t\text{-N}_2\text{O}_4 \cdots \text{NH}_3$. The stability of complex IM_SA1 increased by $5.4-10.3 \text{ kcal mol}^{-1}$ when compared to the reactant complexes IM_WM1 and IM_WD1, with the distance of the $\text{N}(t\text{-N}_2\text{O}_4) \cdots \text{N}(\text{NH}_3)$ bond reduced by $0.01-0.04 \text{ \AA}$. After the formation of the complex IM_SA1, Channel SA proceeded through the transition state TS_SA to form a ten-membered ring hydrogen-bonded complex $\text{HNO}_3 \cdots \text{H}_2\text{SO}_4 \cdots \text{NH}_2\text{NO}$ (labeled as IMF_SA). Similar to the transition state TS_WM described above, TS_SA was in the middle of a double hydrogen atom transfer process, with the H_2SO_4 moiety serving as a bridge for the hydrogen transfer. From the viewpoint of the energy barrier height, Channel SA was a barrierless process. In comparison to H_2O and $(\text{H}_2\text{O})_2$, H_2SO_4 could lower the energy barrier, at least by $0.7-4.6 \text{ kcal mol}^{-1}$. Complex IMF_SA showed a ten-membered ring structure. It had a binding energy of $28.0 \text{ kcal mol}^{-1}$ to the separate reactants $t\text{-N}_2\text{O}_4$, NH_3 , and H_2SO_4 , which was $13.0 \text{ kcal mol}^{-1}$ lower than the

Table 2 Calculated rate constants (k , $\text{cm}^3 \text{ mol}^{-1} \text{ s}^{-1}$) for the $t\text{-N}_2\text{O}_4 + \text{NH}_3$ reaction with H_2O , $(\text{H}_2\text{O})_2$, and H_2SO_4 calculated by master equation within the range of 213–320 K and 0–30 km^{a,b}

Altitude	T (K)	k_{R}	k_{WM}	k_{WD}	k_{SA}
0 km	280	5.20×10^{-17}	1.16×10^{-19}	6.05×10^{-16}	1.49×10^{-16}
	290	6.13×10^{-17}	1.33×10^{-19}	5.29×10^{-16}	1.50×10^{-16}
	298	6.95×10^{-17}	1.48×10^{-19}	4.76×10^{-16}	1.50×10^{-16}
	300	7.17×10^{-17}	1.52×10^{-19}	4.64×10^{-16}	1.51×10^{-16}
	310	8.32×10^{-17}	1.72×10^{-19}	4.11×10^{-16}	1.52×10^{-16}
	320	9.57×10^{-17}	1.93×10^{-19}	3.66×10^{-16}	1.54×10^{-16}
5 km	259.3	3.59×10^{-17}	8.59×10^{-20}	8.30×10^{-16}	1.48×10^{-16}
10 km	229.7	1.92×10^{-17}	5.32×10^{-20}	1.40×10^{-15}	1.51×10^{-16}
15 km	212.6	1.24×10^{-17}	3.94×10^{-20}	1.98×10^{-15}	1.56×10^{-16}
20 km	215.5	1.28×10^{-17}	4.15×10^{-20}	1.86×10^{-15}	1.55×10^{-16}
25 km	218.6	1.37×10^{-17}	4.38×10^{-20}	1.74×10^{-15}	1.54×10^{-16}
30 km	223.7	1.55×10^{-17}	4.79×10^{-20}	1.57×10^{-15}	1.53×10^{-16}

^a k_{R} , k_{WM} , k_{WD} , and k_{SA} were respectively denoted the rate constants for the $t\text{-N}_2\text{O}_4 + \text{NH}_3$, $t\text{-N}_2\text{O}_4 + \text{NH}_3 \cdots \text{H}_2\text{O}$, $t\text{-N}_2\text{O}_4 + \text{NH}_3 \cdots (\text{H}_2\text{O})_2$, and $t\text{-N}_2\text{O}_4 + \text{NH}_3 \cdots \text{H}_2\text{SO}_4$ reactions. ^b The 0–30 km data were reported from ref. 88 and 89.



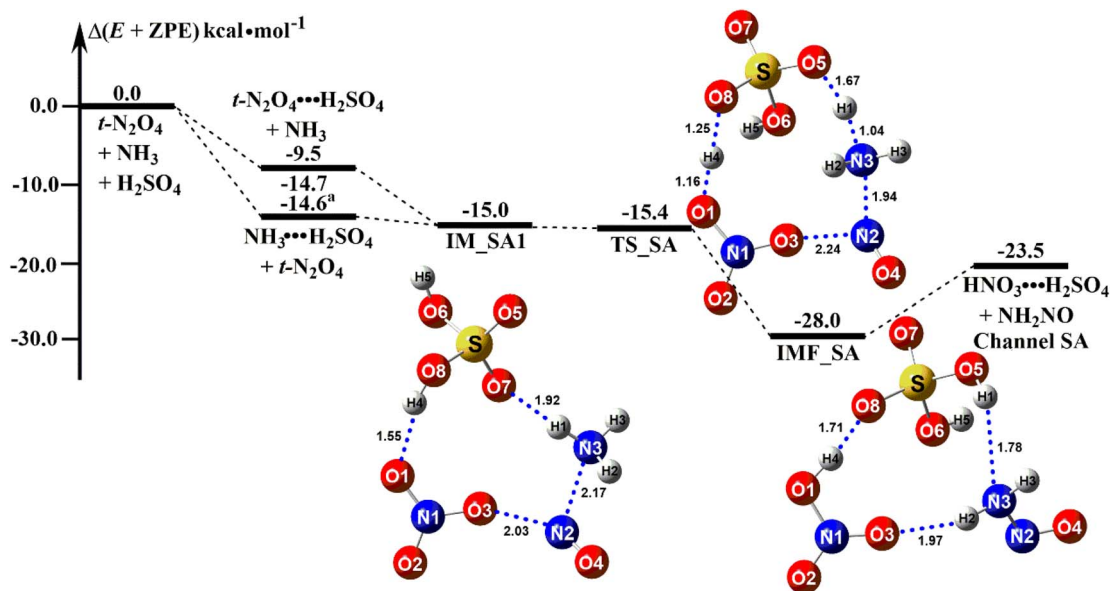


Fig. 3 Potential energy profiles for the $t\text{-N}_2\text{O}_4 + \text{NH}_3 \rightarrow \text{HNO}_3 + \text{NH}_2\text{NO}$ reaction catalyzed by H_2SO_4 at the CCSD(T)/aug-cc-pVTZ//B3LYP-D3/6-311+G(3df,2pd) level (a) The values were reported from ref. 92.

reactant complex IM_SA1. The rate constant of the $t\text{-N}_2\text{O}_4 + \text{NH}_3 \cdots \text{H}_2\text{SO}_4$ reaction is one order of magnitude more than that of the ammonolysis reaction of $t\text{-N}_2\text{O}_4$ without a catalyst, as shown in Table 2. This result revealed that H_2SO_4 in the $t\text{-N}_2\text{O}_4 + \text{NH}_3 \cdots \text{H}_2\text{SO}_4$ reaction plays a favorable catalytic role in promoting the ammonolysis reaction of $t\text{-N}_2\text{O}_4$. Thus, the ammonolysis reaction of $t\text{-N}_2\text{O}_4$ assisted by $(\text{H}_2\text{O})_2$ and H_2SO_4 was more kinetically favorable than the reaction with H_2O .

3.4 Kinetics and implication in atmospheric chemistry

According to previous reports,^{93–96} it is clear that the rate constant listed in Table 2 is insufficient to predict the atmospheric importance of the ammonolysis reaction of $t\text{-N}_2\text{O}_4$ assisted by $(\text{H}_2\text{O})_2$ and H_2SO_4 . To understand the atmospheric effect of $(\text{H}_2\text{O})_2$ and H_2SO_4 on the ammonolysis reaction of $t\text{-N}_2\text{O}_4$, we introduced the effective rate constants (k') to calculate the relative efficiency of neutral and acid trace gases affecting the atmospheric reaction^{97–100} and to compare the rate constant for the naked reaction.

On the basis of the rate constant for Channels WM, WD, and SA, the equilibrium constant for the bimolecular formation ($\text{NH}_3 \cdots \text{H}_2\text{O}$, $\text{NH}_3 \cdots (\text{H}_2\text{O})_2$, and $\text{NH}_3 \cdots \text{H}_2\text{SO}_4$) and the

concentrations of H_2O , $(\text{H}_2\text{O})_2$, and H_2SO_4 are listed in Tables S4–6,[†] as stated in eqn (5)–(7).

$$k'_{\text{WM}} = k_{\text{WM}} \times K_{\text{eq}}(\text{NH}_3 \cdots \text{H}_2\text{O}) \times [\text{H}_2\text{O}] \quad (5)$$

$$k'_{\text{WD}} = k_{\text{WD}} \times K_{\text{eq}}(\text{NH}_3 \cdots (\text{H}_2\text{O})_2) \times [(\text{H}_2\text{O})_2] \quad (6)$$

$$k'_{\text{SA}} = k_{\text{SA}} \times K_{\text{eq}}(\text{NH}_3 \cdots \text{H}_2\text{SO}_4) \times [\text{H}_2\text{SO}_4] \quad (7)$$

where k_{WM} , k_{WD} , and k_{SA} are the bimolecular rate constants for Channel WM, Channels WD, and SA, respectively; $K_{\text{eq}}(\text{NH}_3 \cdots \text{H}_2\text{O})$, $K_{\text{eq}}(\text{NH}_3 \cdots (\text{H}_2\text{O})_2)$, and $K_{\text{eq}}(\text{NH}_3 \cdots \text{H}_2\text{SO}_4)$ are the equilibrium constants for the formation of complexes $\text{NH}_3 \cdots \text{H}_2\text{O}$, $\text{NH}_3 \cdots (\text{H}_2\text{O})_2$, and $\text{NH}_3 \cdots \text{H}_2\text{SO}_4$. $[\text{H}_2\text{O}]$, $[(\text{H}_2\text{O})_2]$, and $[\text{H}_2\text{SO}_4]$ represent the concentrations of H_2O , $(\text{H}_2\text{O})_2$, and H_2SO_4 taken from previous reports.^{44,87} The k' for the ammonolysis reaction of $t\text{-N}_2\text{O}_4$ with X at 0 km altitude and at different altitudes (5–30 km) in the troposphere was calculated.

3.4.1 Zero kilometer altitude. As seen in Table 3, the calculated value of k'_{SA} can compete with k'_{WD} at 280 K. With increasing temperature, the calculated k'_{SA} was $\sim 1\text{--}3$ orders of magnitude smaller than the values of k'_{WD} , showing that the ammonolysis reaction of $t\text{-N}_2\text{O}_4$ was superior to that of an acidic

Table 3 The effective rate constants (k' , $\text{cm}^3 \text{molecule}^{-1} \text{s}^{-1}$) for the $t\text{-N}_2\text{O}_4 + \text{NH}_3$ reaction assisted by H_2O , $(\text{H}_2\text{O})_2$, and H_2SO_4 within the temperature range of 280–320 K^{a,b} (0 km)

Catalysts	T/K	280 K	290 K	298 K	300 K	310 K	320 K
k'_{WM}	100% RH ^c	4.29×10^{-22}	6.96×10^{-22}	1.02×10^{-21}	1.11×10^{-21}	1.70×10^{-21}	2.49×10^{-21}
k'_{WD}	100% RH ^c	2.45×10^{-21}	3.68×10^{-21}	5.16×10^{-21}	5.46×10^{-21}	7.70×10^{-21}	1.02×10^{-20}
k'_{SA}	$[\text{H}_2\text{SO}_4]^c = 10^8 \text{ mol cm}^3$	1.39×10^{-21}	5.49×10^{-22}	2.72×10^{-22}	2.32×10^{-22}	1.03×10^{-22}	4.71×10^{-23}

^a k'_{WM} , k'_{WD} , and k'_{SA} were respectively denoted the effective rate constants for the $t\text{-N}_2\text{O}_4 + \text{NH}_3 \cdots \text{H}_2\text{O}$, $t\text{-N}_2\text{O}_4 + \text{NH}_3 \cdots (\text{H}_2\text{O})_2$, and $t\text{-N}_2\text{O}_4 + \text{NH}_3 \cdots \text{H}_2\text{SO}_4$ reactions. ^b The values of temperature were reported from ref. 88 and 89. ^c The values of concentrations were reported from ref. 44 and 87.



Table 4 The effective rate constants (k' , $\text{cm}^3 \text{ molecule}^{-1} \text{ s}^{-1}$) for the $t\text{-N}_2\text{O}_4 + \text{NH}_3$ reaction with H_2O , $(\text{H}_2\text{O})_2$, and H_2SO_4 at different altitudes in troposphere^a

Altitude (km) ^b	T (K) ^b	P (torr) ^b	k'_{WM}	k'_{WD}	k'_{SA}	$k'_{\text{SA}}/k_{\text{tot}}$
5	259.3	406.75	5.62×10^{-23}	1.50×10^{-22}	1.81×10^{-21}	5.05×10^{-5}
10	229.7	202.16	2.12×10^{-23}	1.74×10^{-22}	9.47×10^{-21}	4.93×10^{-4}
15	212.6	91.20	1.38×10^{-25}	2.95×10^{-26}	3.74×10^{-21}	3.02×10^{-4}
20	215.5	41.04	6.12×10^{-26}	4.52×10^{-27}	4.00×10^{-22}	3.11×10^{-5}
25	218.6	19.00	3.05×10^{-26}	8.56×10^{-28}	2.71×10^{-21}	1.97×10^{-4}
30	223.7	8.36	1.33×10^{-26}	1.12×10^{-28}	7.88×10^{-21}	5.06×10^{-4}

^a k'_{WM} , k'_{WD} , and k'_{SA} were respectively denoted the effective rate constants for the $t\text{-N}_2\text{O}_4 + \text{NH}_3 \cdots \text{H}_2\text{O}$, $t\text{-N}_2\text{O}_4 + \text{NH}_3 \cdots (\text{H}_2\text{O})_2$, and $t\text{-N}_2\text{O}_4 + \text{NH}_3 \cdots \text{H}_2\text{SO}_4$ reactions; $k_{\text{tot}} = k_{\text{R}} + k'_{\text{WD}} + k'_{\text{SA}}$. ^b The values of altitude, temperature and pressure were reported from ref. 87, 88 and 89.

(H_2SO_4) catalyst in the presence of neutral (H_2O and $(\text{H}_2\text{O})_2$) catalysts. However, the H_2SO_4 -catalyzed reaction can be neglected because its calculated k'_{SA} was at least 4 orders of magnitude lower than the corresponding value of k_{R} in the naked reaction of the ammonolysis reaction of $t\text{-N}_2\text{O}_4$. This indicated that the contributions of H_2SO_4 to the ammonolysis reaction of $t\text{-N}_2\text{O}_4$ in atmospheric chemistry are not obvious within the temperature range of 280–320 K (at 0 km altitude).

3.4.2 Higher altitudes. The k' for the ammonolysis reaction of $t\text{-N}_2\text{O}_4$ with H_2O , $(\text{H}_2\text{O})_2$, and H_2SO_4 were calculated within the 5–30 km altitude range, and the calculated k' is listed in Table 4. It can be seen in Table 4 that the contribution of H_2SO_4 was most obvious in the catalysts of H_2O , $(\text{H}_2\text{O})_2$, and H_2SO_4 within the altitude range of 5–30 km, since the value of k'_{SA} was larger by 1–8 orders of magnitude than that of k'_{WD} . In order to quantitatively assess the impact of H_2SO_4 on the ammonolysis reaction of $t\text{-N}_2\text{O}_4$, the total rate constant k_{tot} can be calculated using eqn (8). The branching ratio for $k'_{\text{SA}}/k_{\text{tot}}$ in Table 4 was calculated to be 3.11×10^{-5} – 5.06×10^{-4} at 5–30 km. This indicates that the contribution of the H_2SO_4 -assisted ammonolysis reaction of $t\text{-N}_2\text{O}_4$ can be negligible in atmospheric gas-phase chemistry.

$$k_{\text{tot}} = k_{\text{R}} + k'_{\text{WD}} + k'_{\text{SA}} \quad (8)$$

4. Summary and conclusions

In this work, the favorable route for the ammonolysis reaction of $t\text{-N}_2\text{O}_4$ in the presence of neutral (H_2O and $(\text{H}_2\text{O})_2$) and acidic (H_2SO_4) catalysts was investigated using the quantum chemical method of CCSD(T)/aug-cc-pVTZ//B3LYP-D3/6-311++G(3df,2pd) and the master equation. The calculated results show that the energy barrier for the ammonolysis reaction of $t\text{-N}_2\text{O}_4$ increased when H_2O was present, but when $(\text{H}_2\text{O})_2$ was present, the reaction energy barrier decreased to $0.3 \text{ kcal mol}^{-1}$, which was $3.4 \text{ kcal mol}^{-1}$ lower than the ammonolysis reaction of $t\text{-N}_2\text{O}_4$ without the catalyst, especially when H_2SO_4 was directly involved in the reaction that is a barrierless process. In terms of the effective rate constant, $(\text{H}_2\text{O})_2$ outperforms the other catalysts in the temperature range of 280–320 K (0 km). Moreover, the effect of H_2SO_4 on the ammonolysis reaction of $t\text{-N}_2\text{O}_4$ is obvious at higher altitudes of 5–30 km. Overall, this work will

give a new insight into how the neutral and acidic catalysts affect the formation of HNO_3 from the ammonolysis reaction of $t\text{-N}_2\text{O}_4$. As HNO_3 is an important source of acid rain, the present work will provide a potential formation pathway for HNO_3 , which plays a crucial role in the formation of acid rain.

Author contributions

Ruxue Mu: investigation, data curation, visualization, writing – original draft. Weixin Zhou: supervision, writing – review and editing, project administration. Zhaozhao Hong: visualization, data curation. Rui Wang: writing – review and editing, project administration. Quan Liu: visualization, data curation. Qiang Zhang: formal analysis. Min Jiang: formal analysis. Balaganesh Muthiah: supervision, writing – review and editing. Tianlei Zhang: writing – review and editing, project administration.

Conflicts of interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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