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Photolysis of nitrophenols in gas phase and aqueous environment: a potential daytime source for atmospheric nitrous acid (HONO)[†]

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The daytime atmospheric nitrous acid (HONO) source has attracted considerable attention due to its important role in determining the concentrations of hydroxyl radicals (OH[•]), O₃, and secondary organic aerosols (SOAs). The widespread nitrophenols (NPs) are potential HONO sources; however, the relevant reaction mechanism has not been explored yet. We employ quantum mechanical calculations combined with the statistical RRKM rate theory to explore the photochemical reaction paths of two common NPs (2-NP and 4-NP) to form HONO in the gas and aqueous phases. It is revealed that both NPs have strong sunlight absorption, and photolysis occurs on the excited T₁ state. Both NPs can easily dissociate under light in both gas and aqueous phases, where OH[•] and NO are the major products rather than HONO. Due to the higher solubility, 4-NP is found to more easily generate HONO in aqueous solution, which experiences the intermolecular excited state hydrogen transfer with vicinal water molecules. Kinetics shows that the relative humidity (RH) hardly promotes the gas-phase photolysis, and the temperature slightly affects the reaction rates. However, as a minor product, HONO generations from gas-phase 2-NP and aqueous 4-NP have high photolysis frequencies of 5.73×10^{-5} and $5.25 \times 10^{-6} \text{ s}^{-1}$, respectively, indicating that the hydrolysis of NPs can be important sources for atmospheric HONO.

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Environmental significance

HONO has a great impact on atmospheric chemistry, and its origins have received intensive attention. As one of the abundant components in the heavily polluted atmosphere, nitrophenols are considered as a potential photochemical source of HONO. Using quantum mechanical calculations combined with the statistical rate theory, we unravel the possible photochemical pathway of nitrophenols, especially for HONO formation, which provides significant implications for understanding the atmospheric photochemical processes.

1. Introduction

Nitrous acid (HONO) plays a significant role in atmospheric chemistry,^{1–3} which can produce 20–80% of hydroxyl radicals (OH[•]) in tropospheric reactions during the heavily polluted daytime.^{4–8} The produced OH[•] are important oxidants in the atmosphere and have a great impact on the formation of secondary organic aerosols (SOAs).⁹ The sources of HONO in the atmosphere are rather diverse, and nitrogen oxides (NO₃[–], NO₂, etc.) are commonly accepted.^{10–19} In recent years, researchers have observed HONO generated from atmospheric organic compounds (AOCs) in particulate matters.²⁰ However, AOCs as the source of HONO is still not clear. As one of the typical AOCs, the aromatic compounds containing nitro groups, which widely

exist in forest, rural, urban, and marine areas, have been considered as one of the important potential sources of HONO.^{21–24} Such compounds are also known to be harmful to living organisms.^{25–27}

As the simplest nitro-aromatic compounds, nitrophenols (NPs), which can be formed from the oxidation of phenols under high NO_x conditions, as well as from the combustions of fossil fuel and biomass, have received intensive research attention due to their great contributions to the formation of SOAs.^{21,28–32} NPs also show highly photo-absorption properties to solar radiation, which accounts for most of the visible part (50–80%, $\lambda > 400 \text{ nm}$) and $\sim 4\%$ of the ultraviolet part ($\lambda \sim 370 \text{ nm}$), and contribute to photochemical reactions.^{33–36}

A high concentration of NPs has been observed in both gas phase (0.2–52 ng m^{–3}) and particle phase (0.08–768 ng m^{–3}).^{23,37–41} Furthermore, the concentration of NPs in winter is higher than it in summer.^{39,42,43} The nitro group of NP molecules is commonly located at the *ortho*- or *para*-position to the hydroxyl group, corresponding to 2-nitrophenol (2-NP) and 4-nitrophenol (4-NP), respectively. The concentration of 4-NP is slightly greater than

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that of 2-NP in the atmosphere.^{44,45} Pioneer studies have given strong evidence that 2-NP could be a photochemical source for HONO and OH[•]. For example, Sangwan *et al.* found that 2-NP has strong absorption in the range of near-UV ($\lambda = 295\text{--}400\text{ nm}$),^{46,47} indicating that the direct photolysis of 2-NP by sunlight is possible.⁴⁸ Bejan *et al.* detected ~ 100 pptv HONO formation within an hour in the presence of ~ 1.0 ppbv of 2-NP in the gas phase under irradiation ($\lambda = 300\text{--}500\text{ nm}$),⁴⁹ and Nitta *et al.* used time-resolved photoelectron spectroscopy with 29.5 eV probe pulses to find that 2-NP can split off HONO in 0.5–1 ps.⁵⁰ On the other hand, 4-NP can also be a potential photochemical source for HONO formation. Some experimental research studies have proved that 4-NP can generate HONO in the condensed phase, *e.g.*, acidic aqueous solution and the environment with high relative humidity (RH).^{51,52} Since 4-NP can form an intermolecular hydrogen bond with a water molecule, it may interact with water vapor to promote the HONO formation.

Although previous observations have given evidence on HONO production from the photochemical reaction of NPs,^{47,49,51,52} details of the reaction mechanism, such as the reaction pathway, the rate-determining steps (RDSs), the role of water vapor in the reaction, and whether the reaction can happen in the gas phase or aqueous solution are still unclear. In recent theoretical studies, researchers have agreed that NPs easily generate triplet states after photoexcitation. As a result, the photolysis reactions and HONO formation have a high possibility of occurring in the triplet manifolds (especially the T_1 state).^{53,54} Here, we employ quantum mechanical calculations to comprehensively reveal the possible photoreaction pathways of the two typical NPs (2-NP and 4-NP), especially for HONO generation, as well as evaluate its reaction rates in both unimolecular reaction and the reaction with the participation of water molecules (mimicking the high moist atmosphere and aqueous solution). The obtained results provide important novel mechanistic insight into the originality of atmospheric HONO.

2. Computational methods

2.1 Electronic structure calculations

The electronic energies on the unimolecular reaction paths on the S_1 states are obtained by the relaxed scanning technique at the state-averaged complete active space self-consistent field (CASSCF) level of theory, in which the equal weights are used, and then followed by the single-point energy calculations of two lowest singlets and two lowest triplet states with the complete active space second-order perturbation theory (CASPT2).^{55,56} The active spaces for 2-NP and 4-NP are (10e, 10o) and (12e, 11o), respectively, as shown in Fig. S1, ESI† and the rationality of such active spaces was proved by previous works.^{57–59} In addition, the ionization potential-electron affinity (IPEA) shift is set to zero;⁶⁰ the imaginary shift technique (0.2 a.u.) is employed to avoid the intruder-state issue in the CASPT2 calculation.⁶¹ The def2-TZVPP basis sets⁶² are used in the CAS calculations. All of the CASPT2 and CASSCF calculations are carried out with ORCA 4.2.1 package.^{63,64}

Besides the unimolecular reactions, we also explore the more complex photochemical reactions of NPs, such as water-assisted reactions. To reduce the computational cost, density functional theory (DFT) is carried out to compute the reaction pathways in the ground state (S_0) and lowest triplet excited state (T_1), including optimizing the molecular geometries for the equilibrium states (ESs), transition states (TSs), and minimum energy crossing points (MECPs) between the S_0 and T_1 states. The TSs and MECPs are checked to connect the reactants and products using the intrinsic reaction coordinate (IRC) calculations, and the TSs are recognized to have only one imaginary frequency. Optimization of geometries for MECP between the T_1 and S_0 states is carried out by the sobMECP program,⁶⁵ which is the modified version of the Harvey's MECP code.⁶⁶ In all the DFT calculations, the B3LYP/6-311++G(d,p) level of theory^{67–73} is used for all the geometry optimization and energies computation, which is implemented using the Gaussian 09 program.⁷⁴

In addition, linear response time-dependent density functional theory (LR-TDDFT)⁷⁵ is employed to predict the spectroscopic properties of NPs, such as the vertical excitation energies (VEEs) and spin-orbit coupling (SOC) integrals. The TD-B3LYP/6-311++G(d,p) level of theory,^{67–73} which gives the most accurate excitation energy of NPs among several testing functionals (compared in Table S1, ESI†), is employed for these computations. The SOC integrals calculation is implemented in the PySOC program.⁷⁶

2.2 Reaction rate constant calculation

Rice–Ramsperger–Kassel–Marcus (RRKM) theory is used to model the kinetics for the unimolecular photolysis channels of NPs on the T_1 surface, including the photolysis of NPs and water-NP hydrated complexes. RRKM theory assumes that a transition state for passing from the reactant to the product, and rapid intramolecular vibrational energy redistribution among the degrees of freedom of a molecule. The RRKM microcanonical rate constant, $k(E)$, which represents the statistical rate under collision-free conditions,^{77–80} is given by

$$k(E) = \frac{\sigma N^*(E - E_0)}{h\rho(E)} \quad (1)$$

where σ is the reaction path degeneracy, h is Planck's constant, and N^* is the sum of states at the equilibrium configuration with E_0 , which represents the relative energy of this configuration with respect to the reactant whose rovibrational density of states is $\rho(E)$.⁸¹ By neglecting the rotational motion and calculating the sum and density of states classically,⁸⁰ eqn (1) is simplified by

$$k(E) = \sigma \frac{\prod_{i=1}^s \nu_i}{\prod_{i=1}^{s-1} \nu_i^*} \left(\frac{E - E_0}{E} \right)^{s-1} \quad (2)$$

where ν_i and ν_i^* are the vibrational frequencies of the reactants and TSs, respectively.

To further study the temperature effect of the photolysis reaction, the canonical rate constant, $k(T)$, which assumes the energy levels are populated according to the thermal equilibrium, is followed by⁸⁰



$$k(T) = \int_{E_0}^{\infty} k(E)P(E)dE = \sigma \frac{k_B T}{h} \frac{Q^\ddagger}{Q} \exp\left(-\frac{E_0}{k_B T}\right) \quad (3)$$

where k_B is the Boltzmann's constant, $P(E)$ is the probability in the Boltzmann distribution, and Q^\ddagger and Q are vibrational-rotational partition functions for the activated complex and the reactant, respectively. Eqn (3) is often referred to as the unimolecular rate constant in the high-pressure limit. Similar to eqn (2), the canonical rate constant $k(T)$ is described as⁸⁰

$$k(T) = \sigma \frac{\prod_{i=1}^s \nu_i}{\prod_{i=1}^{s-1} \nu_i^\ddagger} \exp\left(-\frac{E_0}{k_B T}\right) \quad (4)$$

The rate constant of intersystem crossing (ISC) is derived from the Fermi's golden rule,^{82–86} which is according to the following formula

$$k_{ISC}^{nm} = \frac{2\pi}{\hbar} \langle S_n | \hat{H}^{SO} | T_m \rangle^2 \times \text{FCWD} \quad (5)$$

where $\langle S_n | \hat{H}^{SO} | T_m \rangle$ is the SOC integrals between the S_n and T_m states, and FCWD is the Franck–Condon (FC) weighted density of states based on the Marcus–Levich–Jortner theory.⁸⁶ Since temperature has little effect on the ISC rate constant,^{87–89} the vibrational and rotational effect can be ignored. The FCWD can be simplified by the Lorentzian or Gaussian function. Then, k_{ISC}^{nm} can be written in terms of a simplified formula as⁹⁰

$$k_{ISC}^{nm} = \frac{2\pi}{\hbar} \langle S_n | \hat{H}^{SO} | T_m \rangle^2 \times \frac{1}{\pi} \frac{\gamma}{\Delta E_{ST}^2 + \gamma^2} \quad (6)$$

The total (experimental observed) ISC rate constant in state S_n is the sum of ISC to any state T_m , which is given by⁸⁶

$$k_{ISC}^n = \sum_m k_{ISC}^{nm} \quad (7)$$

In addition, the bimolecular rate constants for HONO formation from 4-NP in the gas phase are evaluated using the transition state theory (TST) with the Wigner quantum tunneling correction,^{91–93}

$$k(T) = \left(1 + \frac{1}{24} \left(\frac{\hbar \text{Im}(\nu^\ddagger)}{k_B T}\right)^2\right) \times \frac{k_B T}{h} \exp\left(-\frac{\Delta G^\ddagger}{RT}\right) \quad (8)$$

where R is the gas constant, ΔG^\ddagger is the Gibbs free energy changes between the reactant and TS, and $\text{Im}(\nu^\ddagger)$ is the imaginary frequency of TS.

3. Results and discussion

3.1 Excited-state properties

To understand the possibility of a photoreaction in the atmosphere and the initial population of NPs after photo-excitation, we first study the light absorption properties of NPs. The previously reported absorption spectra of NPs showed a broad absorption band in the range of $\lambda = 300$ – 400 nm, indicating NPs can efficiently absorb sunlight in the lower atmosphere.^{46,94} At the level of theory used here (TD-B3LYP/6-311++G(d,p)), the values of VEE for

$S_0 \rightarrow S_1$ of 2-NP and 4-NP are predicted to be $79.8 \text{ kcal mol}^{-1}$ ($\lambda = 358.08 \text{ nm}$) and $87.9 \text{ kcal mol}^{-1}$ ($\lambda = 325.62 \text{ nm}$), respectively, which are consistent with the estimated experimental and theoretical values of $\sim 81.6 \text{ kcal mol}^{-1}$ ($\lambda = 350 \text{ nm}$).^{46,94} As listed in Table S2 and Fig. S2 (ESI†), the VEE of 2-NP is significantly red-shifted compared to that of 4-NP. This is because the intramolecular hydrogen bonds between the hydroxyl group and the nitro group of 2-NP lowers the relative energies of the lowest unoccupied molecular orbital (LUMO) π^* and the subsequent transition.^{95,96} It is noteworthy to mention that a similar red shift is also observed in the spectra of nitrogen, oxygen, and halogen substituted molecules,^{96–98} allowing this class of species to absorb and react at the longer wavelengths of the UV range that are more available in the troposphere.

Water vapors in the atmosphere may also have an impact on the excited state properties of NPs since the oxygen atoms of the NP molecules can be combined with water molecules through hydrogen bonding interaction, as shown in Fig. S3 (ESI†). However, as listed in Table S3 (ESI†), the interaction between NPs and water is relatively weak due to the positive binding free energies, as well as the small changes in VEEs ($< 0.2 \text{ kcal mol}^{-1}$) of the hydrated NP complexes to the lowest two excited states (T_1 and S_1) at the FC point compared to the isolated molecules. Moreover, it is believed that a thermal fluctuation can further cause the absorption of NPs to respond to a larger wavelength range. Therefore, both the S_1 and the T_1 states of NPs can be efficiently populated by light with a wavelength of $> 300 \text{ nm}$ (corresponding to the energy of $95.3 \text{ kcal mol}^{-1}$).

In addition, it is found that the energies of the S_1 and T_4 excited states at the FC points are close ($\Delta E \leq 1.4 \text{ kcal mol}^{-1}$, TD-B3LYP/6-311++G(d,p) level, see Table S2, ESI†) for both 2-NP and 4-NP, leading to large SOC integrals of 10.46 cm^{-1} and 39.41 cm^{-1} , respectively (Table S4, ESI†). Such small energy intervals and large SOC integrals indicate a high possibility for the ISC of NPs from the singlet state to triplet state.

Since the C–N fission could be a key step for the photolysis of NPs, the corresponding potential energy surfaces (PESs) on the S_1 states of 2-NP and 4-NP are computed at the CASPT2//CASSCF level, as shown in Fig. 1. Due to the large dissociation barriers ($> 30 \text{ kcal mol}^{-1}$) on the S_1 state, NPs can hardly react on the S_1 state, followed by a rapid ISC. In addition, a barrierless process from 2-NP to a hydrogen-shifted 2-NP in the S_1 state (namely aci-2-NP) is found, as well as the large S_0 – T_1 gaps ($\sim 80 \text{ kcal mol}^{-1}$ for 2-NP and 4-NP; $\sim 30 \text{ kcal mol}^{-1}$ for the aci-2-NP*, CASPT2 level), indicating that the excited-state hydrogen transfer (ESHT) is coupled to ISC for 2-NP. Such result can be supported by the recent experiments and simulation observations, which showed the ISC of NPs and ESHT of 2-NP can occur within a picosecond.^{50,58,59,99} Besides, the aci-2-NP has two isomers on the S_0 state and three isomers on the T_1 state based on the geometry optimization at the B3LYP/6-311++G(d,p) level (see Fig. S4, ESI†). In addition, this barrierless ESHT only occurs intramolecularly, so that the vicinal water molecule does not affect this process (see Fig. S5, ESI†). Thus, water vapors should have little influence on the photochemical reactions (both the ESHT and C–N bond fission) of NPs due to the weak water–NP interaction.



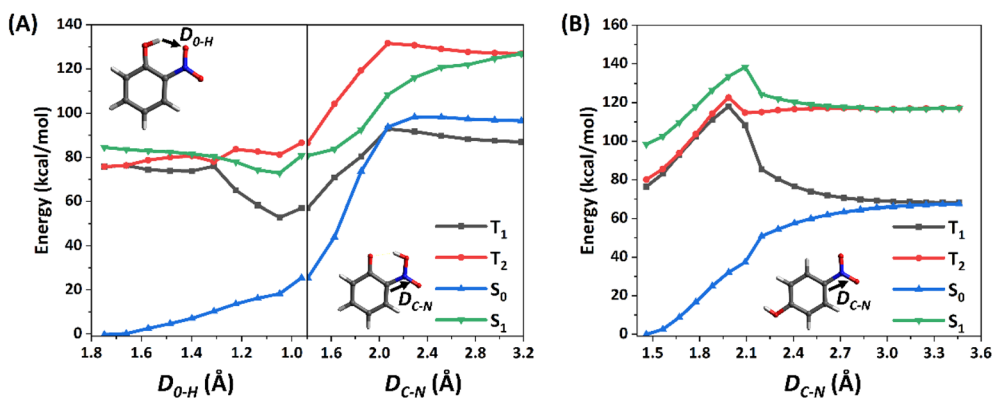


Fig. 1 Computed energy profiles (in kcal mol⁻¹) for the reaction pathways of (A) 2-NP, and (B) 4-NP in the lowest two singlet and triplet excited states (relaxed scan in the S₁ state at CASPT2//CASSCF/def2-TZVPP level). See text for discussion.

3.2 Photochemical reactions pathways of 2-NP

Both pioneer research studies and the above discussions have shown that 2-NP is favored for ISC coupled with the ESHT under the light condition. Such process can occur within a picosecond; therefore, the photoreaction of 2-NP can occur on the T₁ state.^{50,58,59,99} As shown in Fig. 2, the energy gap between the S₁ and T₄ states is only ~0.06 eV, corresponding to the ISC rate of $3.92 \times 10^{10} \text{ s}^{-1}$, which is close to the calculated total observable ISC rate $k_{\text{ISC1}}(\text{obs})$. This result illustrates that the S₁ state of 2-NP should firstly transit to T₄, and then hop back to the T₁ state. Since the hopping process from T₄ to T₁ is fast, we consider that the photochemical reactions mainly take place on the lowest

triplet state (T₁). In previous studies, 2-NP has the ability to produce both HONO and OH[•].^{46,47,49,50,53} Thus, we separately study the photochemical reactions from the aci-2-NP isomers initiated by the C–N or O–N bond fission, and determine the possible reaction pathways in view of the energetic and kinetics aspect.

In the experimental research, HONO formation from 2-NP is a pure gas-phase experiment without water.⁴⁹ Therefore, the photolysis of 2-NP can be a unimolecular reaction, as shown in Fig. 2, which is examined by the intrinsic reaction coordinate (IRC) methodology. It is believed that the ESHT easily occurs. Thus, the photolysis reaction starts from the aci-2-NP isomer.

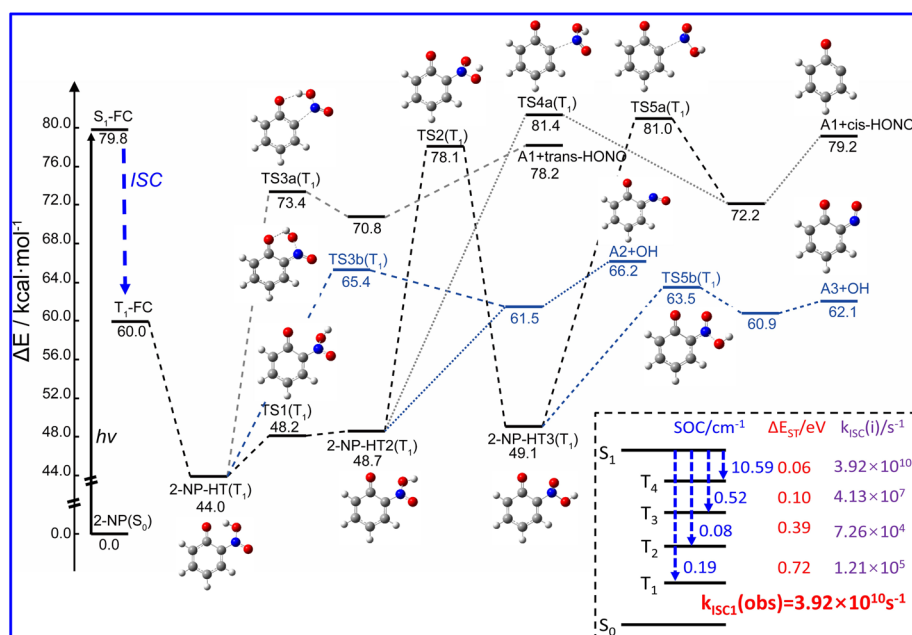


Fig. 2 Computed photolysis pathways (at B3LYP/6-311++G(d,p) level) of the 2-NP molecule in the gas phase for the formation of HONO and OH[•]. Key parameters for the ISC of 2-NP at the FC point are shown in the bottom right corner, including the SOC integrals (in cm⁻¹), ΔE_{ST} (in eV), ISC rate constant $k_{\text{ISC}(i)}$ (in s⁻¹) between the lowest singlet state (S₁) and lowest four triplet states (T_n, n = 1, 2, 3, 4), as well as the observed ISC rates $k_{\text{ISC1}}(\text{obs})$ (in s⁻¹), which are the sums of each ISC channel. The zero-point energy (ZPE) correction is included in the energy values of ΔE. See text for discussions.



For unimolecular decomposition, both S_0 and T_1 states are able to generate HONO, as shown in Fig. S7 and S8.† However, although the IRC calculation proves a connection between 2-NP (or aci-2-NP) and HONO, it still cannot represent the real reaction pathway because the IRC neglects the vibrational and kinetic effects. To illustrate the rationality of the reaction pathway, we analyze the dynamical trajectories accomplished by the Born–Oppenheimer molecular dynamics (BOMD) simulation starting from the structure of the transition state. For each transition state, 25 trajectories are simulated within 400 fs. The results show that among the four isomers (2-NP, 2-NP-HT1, 2-NP-HT2, 2-NP-HT3), only 2-NP-HT2 and 2-NP-HT3 can form HONO on the T_1 state, and the rest of the trajectories all roll back to the reactant, as shown in Fig. S7 and S8.† Thus, the HONO formation can only appear on the T_1 state, although there is still some possibility for hopping back to the S_0 state. The S_0 state is found to hardly contribute to the HONO formation.

Then, we consider the unimolecular reaction starting from aci-2-NP, and find that only aci-2-NP on the T_1 state can form HONO *via* C–N bond fission under UV light irradiation with the energy of TS being 73.4, 81.4, and 81.0 kcal mol^{−1} for 2-NP-HT1, 2-NP-HT2 and 2-NP-HT3, respectively (Fig. 2). This energy barrier of the TS is close to the $S_0 \rightarrow S_1$ vertical excitation energy (79.8 kcal mol^{−1}). After C–N bond decomposition, a van der Waals complex between two fragments is formed at first, and then the HONO molecule is released. Surprisingly, the energy of the isolated A1 radical and *trans*-HONO is still higher than that of TS3a, which illustrates that the 2-NP-HT1(T_1) can hardly release HONO into the atmosphere, and only the other two isomers are available. Therefore, the 2-NP-HT1(T_1) isomer is not likely to generate HONO, which is consistent with the BOMD results. After releasing HONO, as shown in Fig. S9,† biradical

fragments A1 (open-shell electronic structure) may rearrange into a five-membered-ring molecule (P2), but this process is not very easy (with a barrier of ~14.0 kcal mol^{−1}). Since the ground state of A1 is a triplet state, a bimolecular hydrolysis reaction may occur. With the presence of a water molecule, A1 can be easily converted to catechol (P1) with a small energy barrier of ~4.0 kcal mol^{−1}.

Another photochemical pathway is the OH[•] formation path. Among all the TSs in the dissociation pathways shown in Fig. 2, the OH[•] formation pathways have lower energies of transition states than the HONO formation paths, which illustrate that 2-NP is more likely to produce OH[•] instead of HONO through gas phase photolysis. After producing OH[•], A2/A3 radicals (isomers) may further react with water, as shown in Fig. S10.† This reaction pathway has the chance to release NO and further react with OH[•], which was previously produced to generate HONO. However, this process seems not likely to happen due to the large energy barrier for the NO release (>38 kcal mol^{−1}).

Water vapor plays a crucial role in atmospheric chemistry, thus we further study the photolysis channels for HONO and OH[•] formation in the presence of water as shown in Fig. 3. We find that water vapor can lower the energy of TS ($\Delta E < 5$ kcal mol^{−1}) in generating HONO and OH[•] compared to the unimolecular reaction. However, the kinetics results show that the participation of water vapor does not have a promoting effect on the kinetics of photolysis for 2-NP, even leading to 1–3 orders of magnitude lower reaction rate constant, as listed in Table 1. Thus, although the energy of transition state is lowered, the reaction is not accelerated by the presence of water vapor. Furthermore, the binding free energies of the 2-NP and aci-2-NP isomers with water is positive, which shows that it is difficult to combine the reactant with water. We also find that the reaction rate constants are increased with excitation wavelength, and the

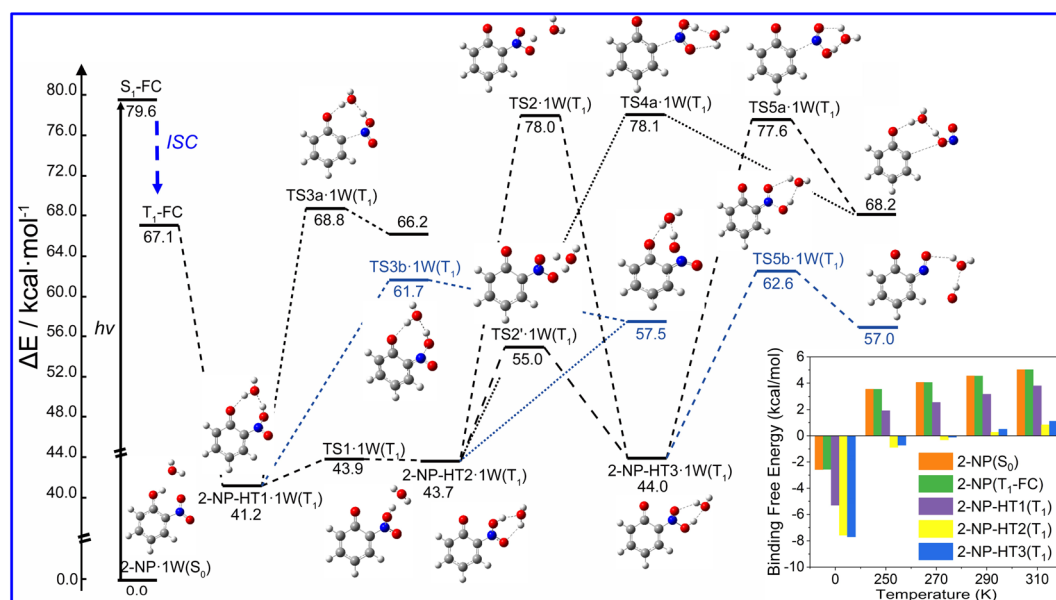


Fig. 3 Computed photolysis pathways (at B3LYP/6-311++G(d,p) level) of the 2-NP hydrated complex for HONO and OH[•] formation. The binding free energies between the 2-NP isomers with a water molecule on the S_0 and T_1 states are shown in the bottom right corner. The ZPE correction is included in the values of ΔE . See text for discussions.



Table 1 Computed RRKM microcanonical rate constants $k(E)$ (in s^{-1}) as the function of internal energy of 70–100 kcal mol^{-1} (corresponding to wavelength range of 280–410 nm) for each elementary photolysis channel of 2-NP, and its hydrated complexes for HONO and OH^{\bullet} formation

Reaction path	Internal energy (kcal mol^{-1})			
	70.0	80.0	90.0	100.0
2-NP-HT1 \rightarrow HONO	1.07×10^5	2.91×10^6	3.15×10^7	1.91×10^8
2-NP-HT2 \rightarrow HONO	1.36×10^5	7.11×10^6	1.19×10^8	9.85×10^8
2-NP-HT3 \rightarrow HONO	1.20×10^5	5.29×10^6	7.90×10^7	6.04×10^8
2-NP-HT1 \rightarrow OH^{\bullet}	5.33×10^8	4.12×10^9	1.88×10^{10}	6.09×10^{10}
2-NP-HT3 \rightarrow OH^{\bullet}	3.45×10^{10}	1.15×10^{11}	2.87×10^{11}	5.88×10^{11}
2-NP-HT1·1W \rightarrow HONO	3.22×10^3	1.25×10^5	1.80×10^6	1.37×10^7
2-NP-HT2·1W \rightarrow HONO	1.36×10^2	2.93×10^4	1.30×10^6	2.20×10^7
2-NP-HT3·1W \rightarrow HONO	3.44×10^2	5.78×10^4	2.18×10^6	3.30×10^7
2-NP-HT1·1W \rightarrow OH^{\bullet}	2.50×10^7	2.67×10^8	1.56×10^9	6.11×10^9
2-NP-HT3·1W \rightarrow OH^{\bullet}	4.55×10^8	3.65×10^9	1.74×10^{10}	5.83×10^{10}

HONO formation rate is 2–5 order of magnitude lower than it is in OH^{\bullet} formation with and without water vapor, which agrees with the previous discussions.

Therefore, the direct photolysis of 2-NP for HONO and OH^{\bullet} formation experiences a unimolecular reaction. OH^{\bullet} is more likely to be generated than HONO, and all the photolysis reactions are nearly irrelevant with the participation of water vapor, which is consistent with the previous experimental results.^{49,50}

3.3 Photochemical reactions pathways of 4-NP

Unlike 2-NP, 4-NP cannot form intramolecular hydrogen bonds, wherein the HONO-moiety cannot be produced under the light irradiation. Here, 4-NP stands for a class of molecules with non-intermolecular hydrogen bonds.

The gas-phase photochemical reactions of 4-NP also follows the paths shown in Fig. 4. Similar to the reaction of 2-NP, the reaction of 4-NP starts from releasing NO_2 or NO. From the energetic view, the energy of TS for NO formation ($74.3 \text{ kcal mol}^{-1}$) is $\sim 8.0 \text{ kcal mol}^{-1}$ lower than that of NO_2 formation ($82.0 \text{ kcal mol}^{-1}$), which can be further reduced by water vapor. The energies of TSs are all below the $S_0 \rightarrow S_1$ transition energy ($87.8 \text{ kcal mol}^{-1}$). Thus, the reactions are able to occur at this energy of irradiation. ISC is also an important photophysical property for 4-NP. We do not account for the transition to the triplet states above the S_1 state. Therefore, there are only three ISC channels for the gas-phase 4-NP with the total ISC rate of $8.68 \times 10^9 \text{ s}^{-1}$, which is shown in Fig. 4(c).

After NO_2 release, H_2O vapor can react with B1 to generate phenol and OH^{\bullet} , then the oxidation of phenol by OH^{\bullet} and NO_2 further generate HONO (Fig. 4(b) and S13†). This reaction pathway may lead to HONO formation from 4-NP in the gas phase, but may not be highly probable to occur due to the small concentration of water in the atmosphere and the positive binding free energy between 4-NP and water molecules. Furthermore, although O_2 is abundant and reactive in the atmosphere, it does not react with B1 due to the larger barrier than that in the hydrolysis of B1 (Fig. 4(b)). Moreover, as shown in Fig. S14,† in the presence of H_2O and $\text{HO-C}_6\text{H}_4$ (B2) fragments, NO can be further transferred into HON, but not efficiently (with a barrier of $\sim 30 \text{ kcal mol}^{-1}$). By contrast, the

formation of HONO from B1 (with a barrier of $\sim 6.0 \text{ kcal mol}^{-1}$) can occur. Except for hydrolysis, the oxygenation of B2 may be more reactive to form hydroquinone and O_2H , but still has the energy barrier of $\sim 18 \text{ kcal mol}^{-1}$ (Fig. S14†). Therefore, only NO_2 and NO could be the major products of the gas-phase photochemistry of 4-NP.

Since 4-NP is soluble due to the intermolecular hydrogen bonding interaction with water, 4-NP can also react in the aqueous phase. In the experiments of solution photochemistry,^{51,52} HONO is formed with a higher rate in the acid environment, where 4-NP can maintain its neutral form, instead of anion. Unlike the gas-phase reactions, the bimolecular reaction is also possible due to the large concentration of water molecules. Fig. 5 shows the PES for photochemistry of 4-NP in solution computed by the solvation model based on density (SMD). As shown in Fig. 5(a), the properties of ISC of 4-NP in solution is similar to that of the gas phase result, and all the total ISC rates are $\sim 10^9 \text{ s}^{-1}$, indicating that water molecules have little affect on the photophysical properties of 4-NPs. The results of the potential energy surfaces (Fig. 5(b)) indicate that the energy of TS for bimolecular reaction is greatly reduced ($\Delta E = 3\text{--}13 \text{ kcal mol}^{-1}$) compared to that of the unimolecular paths, and all paths can efficiently occur under the 70–100 kcal mol^{-1} ($\lambda = 280\text{--}410 \text{ nm}$) irradiation. Due to the large concentration of water molecules in solution, the dissolved 4-NP molecules may undergo intermolecular ESHT. It is found that at least three water molecules are needed to realize this process due to the large separation distance between the hydroxyl and nitro groups in 4-NP molecules. Here, the ISC of molecules after photoexcitation occurs in the FC region, and the subsequent reaction takes place on the T_1 state. As shown in Fig. 5(c), the unimolecular ESHT of the 4-NP·3W hydrated complex has a large reaction energy barrier ($>10 \text{ kcal mol}^{-1}$) on all excited state energy surfaces, so the previous ISC hypothesis is reasonable. As shown in Fig. 5(c) and (d), the energy barriers for the intermolecular ESHT of 4-NP in solution are $\sim 10 \text{ kcal mol}^{-1}$ (unimolecular reaction) and $1.1 \text{ kcal mol}^{-1}$ (bimolecular reaction), which indicates that 4-NP can easily transfer a hydrogen atom to generate aci-4-NP in solution. Then, we consider the dissociation of aci-4-NP on the T_1 state. It is found that the dissociation



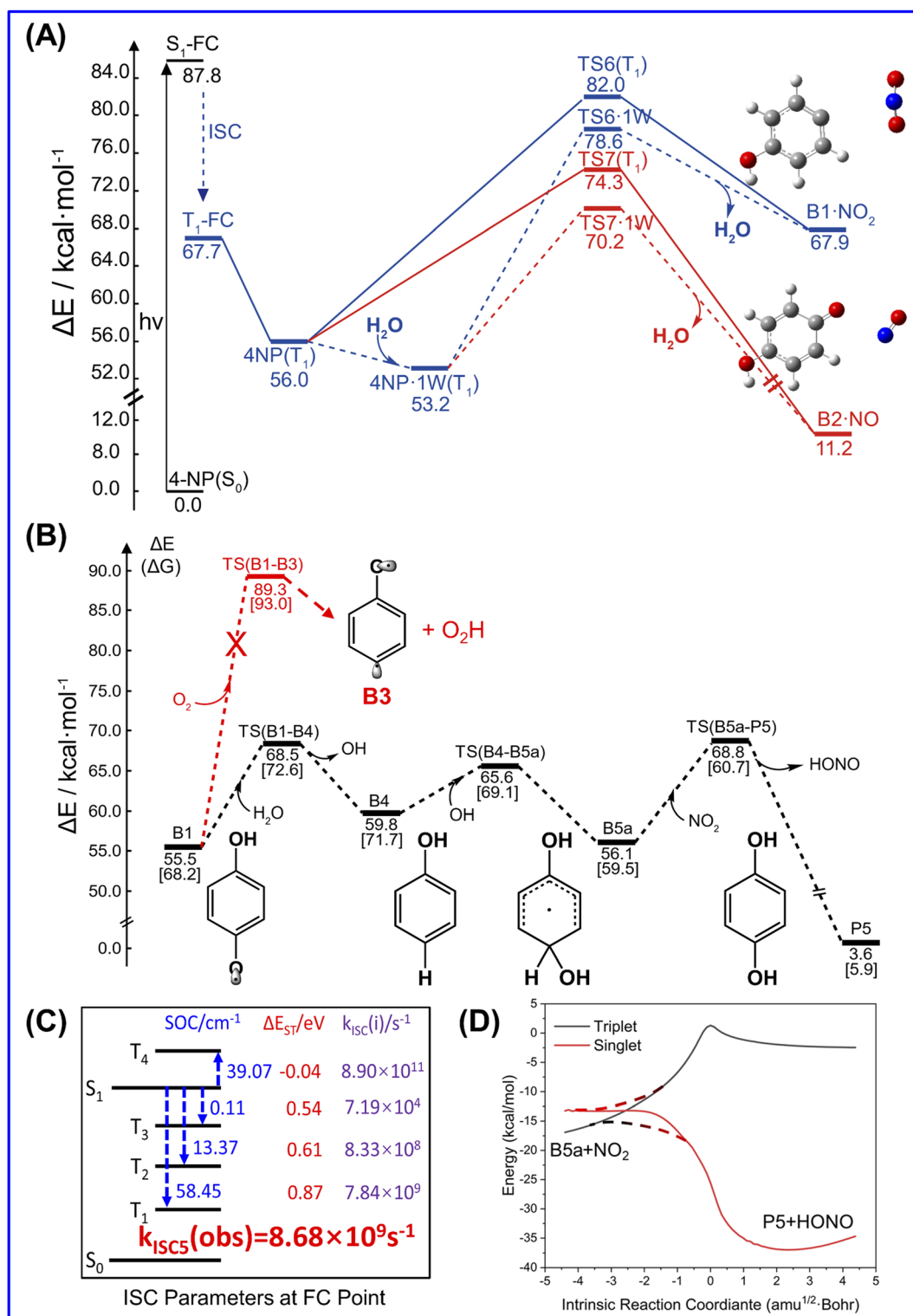


Fig. 4 Photochemical mechanism of 4-NP in the gas phase. (A) Computed photolysis pathways (at B3LYP/6-311++G(d,p) level) of 4-NP for NO₂ and NO formation in the gas phase. (B) Computed gas-phase reaction pathways (at B3LYP/6-311++G(d,p) level) for HONO formation in the presence of O₂ or water vapor from the B1 radical intermediate, which is generated from the C–N bond fission of 4-NP (C) Key parameters for ISC of 4-NP (gas) at the FC point, including SOC integrals (in cm⁻¹), ΔE_{ST} (in eV), ISC rate constant *k*_{ISC}(*i*) (in s⁻¹) between the lowest singlet state (S₁) and lowest four triplet states (T_{*n*}, *n* = 1, 2, 3, 4), as well as the observed ISC rates *k*_{ISC5}(obs) (in s⁻¹), which are the sums of the ISC channels. (D) The IRC of the singlet and triplet states for hydrogen transfer between B5a and NO₂ fragments. The ZPE correction is included in the energy values of ΔE. See text for discussions.



of the N–O bond of this molecule is very difficult since the energy barriers of the unimolecular and hydrated complex are ~ 33 and ~ 35 kcal mol $^{-1}$, respectively. However, it is very easy to generate HONO through dissociating the C–N bond with the energy barrier of only ~ 14 kcal mol $^{-1}$, which is less than the barriers in the bimolecular generation of HONO and unimolecular formation of NO $_2$ and NO. Thus, we can conclude that the dissociation of aci-4-NP on the T $_1$ state is the main source of HONO generation from 4-NP.

We further compute the rate constants of each elementary photolysis channel in both gas and aqueous phases by RRKM theory with the excitation/internal energy of 70–100 kcal mol $^{-1}$ (corresponding to light irradiation of 280–410 nm), as listed in Table 2. For gas-phase reactions, the unimolecular photolysis to form NO $_2$ and NO have the rate constants of 10^6 – 10^8 s $^{-1}$ and 10^8 – 10^9 s $^{-1}$, respectively. Meanwhile, in the presence of water molecules, the rate constants of NO $_2$ and NO formation are 10^4 – 10^7 s $^{-1}$ and 10^7 – 10^8 s $^{-1}$, respectively. From this issue, NO is more prone to generate from the photolysis of 4-NP, and water molecules have no apparent influence on the photolysis rates. For the photolysis in solution, as a comparison, NO $_2$ and NO formation have the rate constants of 10^5 – 10^8 s $^{-1}$ and 10^7 – 10^9 s $^{-1}$, respectively. The photolysis from the hydrated complex is also unfavorable compared to the unimolecular reaction. Therefore, NO $_2$ and NO are only favorable to generate in the gas phase rather than aqueous phase.

Then, we consider the rate constants of several possible reaction channels for the formation of HONO in solution. For the bimolecular reaction, the rate constant of HONO formation is 10^3 – 10^4 s $^{-1}$. However, when an extra water molecule is added, its rate constant decreases to 10^0 – 10^1 s $^{-1}$. It can be concluded that the additional water molecule does not have advantage to the reaction. In addition, the rate constant of intermolecular ESHT of 4-NP with the existence of at least three water molecules is 10^6 – 10^7 s $^{-1}$. Meanwhile, the rate constant of the dissociation of aci-4-NP to form HONO is 10^3 – 10^7 s $^{-1}$, and to generate OH \cdot is 10^9 – 10^{11} s $^{-1}$. With a water molecule present, the rate constant for HONO and OH \cdot formation is 10^5 – 10^8 s $^{-1}$ and 10^9 – 10^{10} s $^{-1}$. Here, we have found that water molecules can promote the dissociation of aci-4-NP, which leads to HONO formation in solution. That is to say, the formation of HONO from 4-NP in solution does not present a bimolecular reaction, but is similar to 2-NP molecule, *i.e.*, carries out an ESHT. This is then followed by unimolecular photolysis, which is contrary with the experimental suggestions.^{51,52} To summarize, these results indicate that the photochemical reaction of 4-NP preferably generates NO $_2$ and NO in gas phase, while it generates HONO in aqueous solution.

3.4 Kinetics and atmospheric implications for HONO formation

To evaluate the contribution to HONO formation of NPs in the real atmosphere, we analyze the reaction kinetics of NPs to release

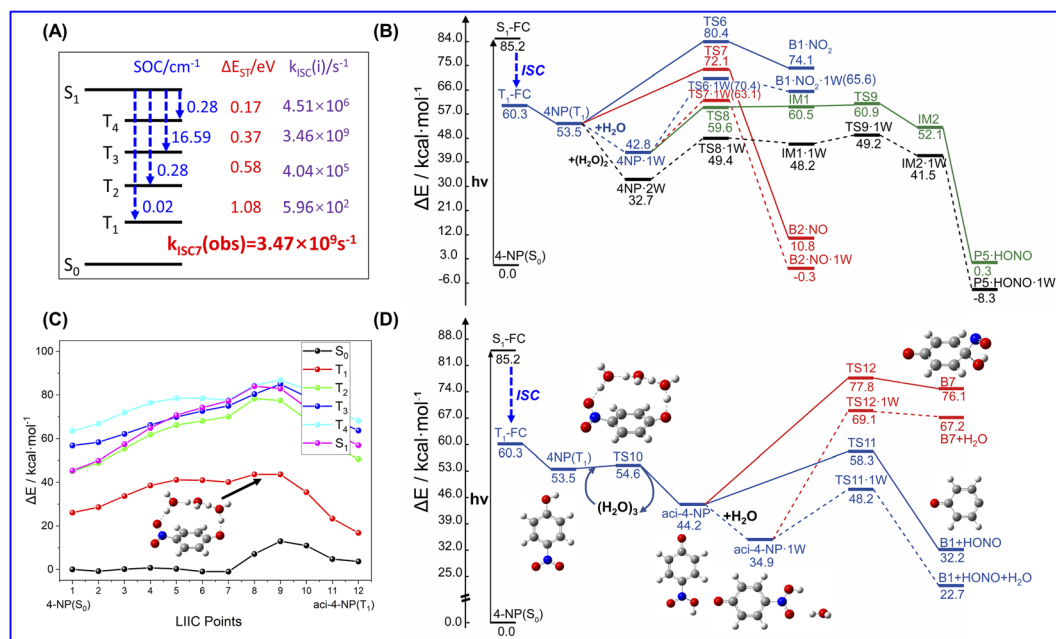


Fig. 5 Photochemical mechanism of 4-NP in aqueous phase. (A) Key parameters for ISC of 4-NP (aq) at the FC point, including SOC integrals (in cm $^{-1}$), ΔE_{ST} (in eV), ISC rate constant $k_{ISC}(i)$ (in s $^{-1}$) between the lowest singlet state (S_1) and lowest four triplet states (T_n , $n = 1, 2, 3, 4$), as well as the observed ISC rates $k_{ISC6}(obs)$ (in s $^{-1}$), which are the sums of the ISC channels. (B) Computed photolysis pathways (at B3LYP/6-311++G(d,p) level) of 4-NP for NO $_2$, NO and bimolecular HONO formation in aqueous phase. (C) Computed intermolecular ESHT pathway (at TD-B3LYP//B3LYP/6-311++G(d,p) level) of 4-NP in aqueous phase. Each geometry in the potential energy surfaces are obtained by the linear-interpolated internal coordinates (LIIC) technique. (D) Computed photolysis pathways (at B3LYP/6-311++G(d,p) level) of aci-4-NP in water solution initiated by intermolecular ESHT. The ZPE correction is included in the energy values of ΔE . All of the energies are calculated with the SMD solvation model. See text for discussions.



HONO via the photochemical reactions. The typical concentrations of NPs in the atmosphere are $\sim 2.0 \text{ ng m}^{-3}$ (8.60×10^6 molecules per cm^3) for 2-NP and $\sim 5.0 \text{ ng m}^{-3}$ (2.16×10^7 molecules per cm^3) for 4-NP.⁴⁵ It is notable that the concentration of 4-NP can be up to $\sim 768 \text{ ng m}^{-3}$ in heavily polluted weather.^{100,101} The concentration of water vapor is a function of temperature and RH (see ESI† for details). Here, we regard the aqueous phase as water vapor with the concentration at 300 K and 100% RH. All of the rate constants of each elementary reaction are calculated using RRKM theory under the collision-free condition.

The computational details for the time evolution of the photolysis products using the micro-kinetic analysis and related results are summarized in Section S1, ESI.† Firstly, we study the photolysis kinetics under the excitation energy of 70–100 kcal mol^{-1} (corresponding to the wavelength range of 280–410 nm) for 2-NP and 4-NP, as well as the reactions involving water. Almost all of the concentrations and reaction rates of the photolysis products are increased with the increase of the excitation energy (Fig. S16, S17 and S19†). For the gas-phase photolysis of 2-NP, as shown in Table S11,† the yield of OH• is much higher than that of HONO with and without water, demonstrating the branching ratio of $\sim 1:10^{-4}$. Thus, OH• is the main product of the gas-phase photolysis of 2-NP. The relative populations of OH• and HONO from the water-assisted reactions are 10^{-6} to 10^{-4} and 10^{-10} to 10^{-6} , respectively. These values are much lower than the relative populations in the unimolecular reactions of 0.89–0.99 and 10^{-6} to 10^{-4} , respectively, which should be attributed to the low binding free energy between 2-NP and water (Tables S3 and S10, ESI†). In addition, based on the micro-kinetics, we extract the rate constants for OH• and HONO formation from the unimolecular reactions (OH•: 10^7 – 10^9 s^{-1} ; HONO: 10^4 – 10^6 s^{-1}), which are 1–2 orders of magnitude higher than those in the water-assisted reactions.

Thus, the main channel of the photolysis of 2-NP should be a unimolecular reaction, which does not rely on the humidity of the atmosphere.

The photolysis of 4-NP in the gas phase has a similar situation (results are listed in Table S12†), where the water vapors do not promote the reaction and NO is the major product with the formation rate of 10^7 – 10^8 s^{-1} (NO_2 : 10^5 – 10^7 s^{-1}). Considering the following hydrolysis of the B1 fragment, as listed in Table 3, the final relative populations (under an excitation energy of 80 kcal mol^{-1}) of NO_2 , NO, and HONO are 2.16×10^{-2} , 0.6, and 1.8×10^{-17} , respectively, which shows that almost no HONO can be produced from the photolysis of 4-NP in the gas phase. On the other hand, when 4-NP is dissolved in aqueous solution, the relative populations of the products are significantly changed. The final relative populations (under an excitation energy of 80 kcal mol^{-1}) of NO_2 , NO, and HONO become 2.59×10^{-3} , 5.85×10^{-2} , and 8.50×10^{-5} , respectively (Table 3), in which the content of HONO is remarkably increased, although it is still a minor product. Comparing the final relative populations of HONO formed by the aqueous bimolecular reaction of 4-NP and dissociation of aci-4-NP (Table S14, ESI†), the latter reaction path produces more HONO, because the intermolecular ESHT of 4-NP is more favorable than the bimolecular reaction. In aqueous solution, the formation rates of NO_2 , NO and HONO are found to be 10^5 – 10^8 , 10^7 – 10^8 , and 10^2 – 10^5 s^{-1} , respectively (Table S14†).

The photolysis frequency (J , in s^{-1}) is often regarded as one of the key parameters in atmospheric photochemical reaction research, which is defined as

$$J = \int \phi(\lambda) \sigma(\lambda) F(\lambda, \Theta) d\lambda \quad (9)$$

Table 2 Computed RRKM microcanonical rate constant $k(E)$ (in s^{-1}) as the function of the internal energy of 70–100 kcal mol^{-1} (corresponding to a wavelength range of 280–410 nm) for each elementary photolysis channel of 4-NP and its hydrated complex to form NO_2 , NO and HONO in the gas and aqueous phases

Reaction path	Internal energy (kcal mol^{-1})			
	70.0	80.0	90.0	100.0
Gas-phase photolysis				
4-NP $\rightarrow \text{NO}_2$	1.81×10^6	2.72×10^7	1.97×10^8	8.97×10^8
4-NP $\rightarrow \text{NO}$	1.47×10^8	7.55×10^8	2.58×10^9	6.69×10^9
4-NP·1W $\rightarrow \text{NO}_2$	2.35×10^4	5.94×10^5	6.32×10^6	3.86×10^7
4-NP·1W $\rightarrow \text{NO}$	1.14×10^7	7.25×10^7	2.91×10^8	8.58×10^8
Aqueous-phase photolysis				
4-NP $\rightarrow \text{NO}_2$	5.87×10^5	1.03×10^7	8.37×10^7	4.12×10^8
4-NP $\rightarrow \text{NO}$	4.30×10^7	2.33×10^8	8.23×10^8	2.20×10^9
4-NP·1W $\rightarrow \text{NO}_2$	2.03×10^4	8.09×10^5	1.18×10^7	9.05×10^7
4-NP·1W $\rightarrow \text{NO}$	3.62×10^7	3.79×10^8	2.18×10^9	8.42×10^9
4-NP·1W $\rightarrow \text{HONO}$	1.55×10^3	4.88×10^3	1.12×10^4	2.13×10^4
4-NP·2W $\rightarrow \text{HONO}$	3.76×10^0	1.42×10^1	3.73×10^1	7.79×10^1
4-NP·3W $\rightarrow \text{Aci-4-NP} \cdot 3\text{W}$	3.44×10^6	1.27×10^7	3.43×10^7	7.52×10^7
Aci-4-NP $\rightarrow \text{HONO}$	3.19×10^3	2.03×10^5	3.85×10^6	3.47×10^7
Aci-4-NP $\rightarrow \text{OH}^\bullet$	8.23×10^9	2.69×10^{10}	6.59×10^{10}	1.33×10^{11}
Aci-4-NP·1W $\rightarrow \text{HONO}$	4.31×10^5	9.97×10^6	1.00×10^8	5.85×10^8
Aci-4-NP·1W $\rightarrow \text{OH}^\bullet$	1.58×10^9	6.15×10^9	1.72×10^{10}	3.87×10^{10}



Table 3 The relative populations of OH[•], HONO, NO₂ and NO from the photolysis of 2-NP in the gas phase, as well as 4-NP in both gas and solution phases under the excitation energy of 80 kcal mol⁻¹ (356 nm)

	Photolysis product			
	OH [•]	HONO	NO ₂	NO
2-NP (gas)	0.98	1.00×10^{-3}	N.A.	N.A.
4-NP (gas)	N.A.	1.80×10^{-17}	2.16×10^{-2}	6.00×10^{-1}
4-NP (solution)	N.A.	8.50×10^{-5}	2.59×10^{-3}	5.85×10^{-2}

where $\phi(\lambda)$ is the quantum yield of photolysis obtained from the ratio of the reaction rates, $\sigma(\lambda)$ is the photo-absorption cross-section of a molecule, $F(\lambda, \theta)$ is the actinic flux of light source, and θ is the zenith angle. The photo-absorption cross-sections for 2-NP and 4-NP are 1.03×10^{-17} cm² per molecules (250 K), 8.86×10^{-18} cm² per molecules (300 K); and 3.02×10^{-18} cm² per molecules (250 K), 2.80×10^{-18} cm² per molecules (300 K), respectively (Fig. S2, ESI[†]). The actinic flux is $\sim 1 \times 10^{14}$ photons per cm² per nm per s) at 358 nm, which is slightly changed with the variation of the zenith angle and wavelength of light.^{102,103} In addition, we consider the actinic flux as a constant value in the 280–410 nm irradiation. Then, based on these parameters, the photolysis frequency for HONO formation from 2-NP in the gas phase and 4-NP in aqueous solution can be calculated. As shown in Fig. 6 and listed in Table 4, the rate constants, quantum yields, and photolysis frequency for HONO formation are calculated through the micro-kinetic studies. Under the excitation energy of 70–100 kcal mol⁻¹ (corresponding to the wavelength of 280–410 nm), the rate constants are 10^4 – 10^6 s⁻¹ and 10^2 – 10^6 s⁻¹ for 2-NP and 4-NP, respectively. 2-NP has a higher HONO formation rate than 4-NP, which finds a local maximum at the excitation energy of 80 kcal mol⁻¹ (~ 356 nm). These results follow a similar law with the photo-absorption cross-sections of NPs. In addition, the quantum yields of 2-NP (10^{-5} to 10^{-3} s⁻¹) are larger than that of 4-NP (10^{-4} to 10^{-3}) in the lower excitation energy region. Photolysis frequencies of 2-NP and 4-NP are 10^{-8} to 10^{-6} s⁻¹ and

10^{-9} to 10^{-6} s⁻¹, respectively. In the real atmosphere and experimental research, we only focus on the photolysis frequency within an excitation energy range. By summing up each individual value, the total photolysis frequencies under the excitation energy of 70–100 kcal mol⁻¹ are 5.73×10^{-5} and 5.25×10^{-6} s⁻¹ for 2-NP and 4-NP, respectively, which are in good agreement with the experimentally reported values.^{49,52} It can be seen that 2-NP is more favorable in producing HONO than 4-NP in the real atmosphere.

We further consider the temperature effect on the photolysis reactions based on the canonical rate constants, which can be used to describe the ability to overcome the energy barriers between the minimum and transition state, including excitation energies at different temperatures. It is noteworthy that we only use this model to describe the temperature effect qualitatively, rather than provide a real population of each species at different temperatures. As listed in Table S15 (Fig. S16–S19[†]), a higher temperature is favorable for photolysis reaction. Therefore, more HONO originating from the photolysis of NPs should be observed at higher temperatures (*i.e.* summer season). In addition, the energy distribution of the reactant does not shift to higher energy levels with the increase of temperature, as shown in the calculated absorption spectra (Fig. S2[†]). Therefore, the temperature effect in the photolysis of nitrophenols can only affect the kinetics of the reactions.

Moreover, the air–water interface (such as the aerosol surface) is an important place for the reaction to occur. However, we just studied the photochemical reaction of NPs in gas and solution since the reaction channel at the air–water interfaces should not be different from the aqueous phase except for the concentration of water. However, based on the previous discussions, the concentration of water molecules hardly influences the photochemical reaction rates of NPs. Therefore, the reaction kinetics at the air–water interfaces should be similar to the reaction in solution.

To summarize, the kinetics analysis shows that OH[•] and NO can be efficiently produced by the gas-phase photolysis of 2-NP

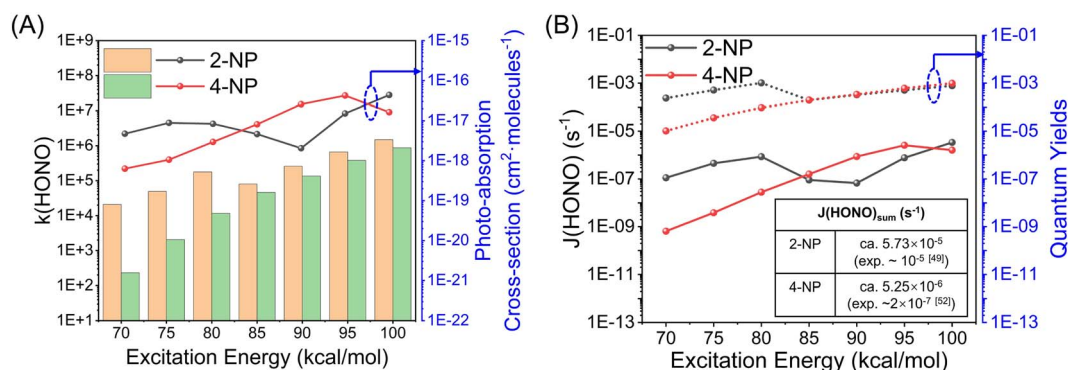


Fig. 6 Kinetics for HONO formation: (A) Microcanonical rate constants for HONO formation, and photo-absorption cross-section of 2-NP (gas phase) and 4-NP (aqueous phase); (B) quantum yields (dash lines) and photolysis frequencies (solid lines) for the HONO formation of 2-NP (gas phase) and 4-NP (aqueous phase). These results are obtained from the micro-kinetics simulations based on the RRKM theory as a function of the excitation energy of 70–100 kcal mol⁻¹ (corresponding to a wavelength range of 280–410 nm). Furthermore, the calculated and experimental measurement of the photolysis frequencies for HONO formation from 2-NP and 4-NP with an excitation energy range of 70–100 kcal mol⁻¹ is listed.

Table 4 Quantum yields of HONO formation as the function of excitation energy of 70–100 kcal mol^{−1} (corresponding to a wavelength range of 280–410 nm) from 2-NP (gas phase) and 4-NP (aqueous phase)

	Excitation energy (kcal mol ^{−1})						
	70.0	75.0	80.0	85.0	90.0	95.0	100.0
2-NP	2.36×10^{-4}	5.10×10^{-4}	1.02×10^{-3}	1.95×10^{-4}	3.26×10^{-4}	5.06×10^{-4}	7.77×10^{-4}
4-NP	1.02×10^{-5}	3.61×10^{-5}	9.47×10^{-5}	1.97×10^{-4}	3.37×10^{-4}	6.07×10^{-4}	9.85×10^{-4}

and 4-NP, and the temperature has minimal influence on the formation rates. Although as the minor products, the gas-phase photolysis reactions still have some contributions to generate both HONO and NO₂. On the other hand, in the condensed aqueous phase, HONO can be efficiently produced from the photolysis of 4-NP with the assistance of water, and a higher temperature apparently benefits the reaction. Based on these results, we can conclude that HONO is more likely to be produced from 2-NP in the gas phase and 4-NP in the aqueous solution. In addition, according to our existing knowledge, a direct experimental comparison of these two molecules in photochemical reactions is still lacking. Therefore, we are looking forward to more precise experimental evidence to compare these two molecules directly on photochemical HONO production.

4. Conclusion

In conclusion, we employ quantum mechanical methods to predict the photoreaction pathways of HONO generation from two typical atmospheric NPs, *e.g.*, 2-NP and 4-NP. The results show that NPs have high solar absorption efficiency in the wavelength range of 300–400 nm, as well as a high probability of ISC, indicating that the photolysis reactions of these molecules occur in the T₁ state. For 2-NP, the photolysis is more likely to occur in the gas phase, where the ESHT rapidly occurs after light irradiation, and then HONO and OH[•] are released. The kinetics study shows that 2-NP and 4-NP can both easily dissociate after light irradiation in the gas phase, while OH[•] and NO are the major products rather than HONO. In addition, since 4-NP can dissolve in water, the photolysis of 4-NP in the condensed aqueous phase is much easier to produce HONO than in the gas phase, which also experiences an ESHT instead of the bimolecular reactions. However, HONO is still a minor product in the aqueous photolysis of 4-NP. These results demonstrate that HONO is a minor product from both water-insoluble NPs (WINP, such as 2-NP) and water-soluble NPs (WSNP, such as 4-NP). However, NP molecules still have considerable contributions to HONO generation in the real atmosphere. Therefore, the photolysis of WINP in the gas phase and WSNP in the aqueous phase can be a significant source for atmospheric HONO, and the gas phase photolysis may be more important.

Author contributions

The manuscript was written through contributions of all authors. All authors have given approval to the final version of the manuscript.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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References

- J. Wang, J. Li, J. Ye, J. Zhao, Y. Wu, J. Hu, D. Liu, D. Nie, F. Shen, X. Huang, D. Huang, D. Ji, X. Sun, W. Xu, J. Guo, S. Song, Y. Qin, P. Liu, J. R. Turner, H. C. Lee, S. Hwang, H. Liao, S. T. Martin, Q. Zhang, M. Chen, Y. Sun, X. Ge and D. J. Jacob, *Nat. Commun.*, 2020, **11**, 2844.
- W. Zhang, S. Tong, C. Jia, L. Wang, B. Liu, G. Tang, D. Ji, B. Hu, Z. Liu, W. Li, Z. Wang, Y. Liu, Y. Wang and M. Ge, *Environ. Sci. Technol.*, 2020, **54**, 12870–12880.
- Y. Liu, Y. Zhang, C. Lian, C. Yan, Z. Feng, F. Zheng, X. Fan, Y. Chen, W. Wang, B. Chu, Y. Wang, J. Cai, W. Du, K. Daellenbach, J. Kangasluoma, F. Bianchi, J. Kujansuu, T. Petäjä, X. Wang, B. Hu, Y. Wang, M. Ge, H. He and M. Kulmala, *Atmos. Chem. Phys.*, 2020, **20**, 13023–13040.
- A. Kukui, M. Legrand, S. Preunkert, M. Frey, R. Loisil, J. Roca, B. Jourdain, M. King, J. France and G. Ancellet, *Atmos. Chem. Phys.*, 2014, **14**, 12373–12392.
- L. Xue, R. Gu, T. Wang, X. Wang, S. Saunders, D. Blake, P. Louie, C. Luk, I. Simpson, Z. Xu, Z. Wang, Y. Gao, S. Lee, A. Mellouki and W. Wang, *Atmos. Chem. Phys.*, 2016, **16**, 9891–9903.
- C. Young, R. Washenfelder, J. Roberts, L. Mielke, H. Osthoff, C. Tsai, O. Pikelnaya, J. Stutz, P. Veres, A. Cochran, T. VandenBoer, J. Flynn, N. Grossberg, C. Haman, B. Lefer, H. Stark, M. Graus, J. Gouw, J. Gilman, W. Kuster and S. Brown, *Environ. Sci. Technol.*, 2012, **46**, 10965–10973.
- J. Zheng, X. Shi, Y. Ma, X. Ren, H. Jabbour, Y. Diao, W. Wang, Y. Ge, Y. Zhang and W. Zhu, *Atmos. Chem. Phys.*, 2020, **20**, 5457–5475.
- J. Nan, S. Wang, Y. Guo, Y. Xiang and B. Zhou, *Atmos. Environ.*, 2017, **154**, 167–178.
- J. Brean, R. Harrison, Z. Shi, D. Beddows, W. Acton, C. Hewitt, F. Squires and J. Lee, *Atmos. Chem. Phys.*, 2019, **19**, 14933–14947.



- 10 C. Han, W. Yang, Q. Wu, H. Yang and X. Xue, *Environ. Sci. Technol.*, 2016, **50**, 5017–5023.
- 11 M. Monge, B. D'Anna, L. Mazri, A. Giroir-Fendler, M. Ammann, D. Donaldson and C. George, *Proc. Natl. Acad. Sci. U. S. A.*, 2010, **107**, 6605–6609.
- 12 M. Brigante, D. Cazoir, B. D'Anna, C. George and D. Donaldson, *J. Phys. Chem. A*, 2008, **112**, 9503–9508.
- 13 C. Ye, N. Zhang, H. Gao and X. Zhou, *Environ. Sci. Technol.*, 2017, **51**, 6849–6856.
- 14 K. Stemmler, M. Ammann, C. Donders, J. Kleffmann and C. George, *Nature*, 2006, **440**, 195–198.
- 15 R. Ammar, M. Monge, C. George and B. D'Anna, *ChemPhysChem*, 2010, **11**, 3956–3961.
- 16 L. Li, Z. Duan, H. Li, C. Zhu, G. Henkelman, J. Francisco and X. Zeng, *Proc. Natl. Acad. Sci. U. S. A.*, 2018, **115**, 7236–7241.
- 17 D. Medeiros and A. Pimentel, *J. Phys. Chem. A*, 2011, **115**, 6357–6365.
- 18 M. Martins-Costa, J. Anglada, J. Francisco and M. Ruiz-López, *J. Am. Chem. Soc.*, 2020, **142**, 20937–20941.
- 19 M. Gen, Z. Liang, R. Zhang, B. Mabato and C. Chan, *Environ. Sci.: Atmos.*, 2022, **2**, 111–127.
- 20 F. Bao, M. Li, Y. Zhang, C. Chen and J. Zhao, *Environ. Sci. Technol.*, 2018, **52**, 6309–6316.
- 21 Z. Wang, J. Zhang, L. Zhang, Y. Liang and Q. Shi, *Atmos. Environ.*, 2021, **246**, 118046.
- 22 M. Mulder, Y. Dumanoglu, C. Efstathiou, P. Kukučka, J. Matejovičová, C. Maurer, P. Přibyllová, R. Prokeš, A. Sofuoglu, S. Sofuoglu, J. Wilson, C. Zetzsch, G. Wotawa and G. Lammel, *Environ. Sci. Technol.*, 2019, **53**, 8914–8924.
- 23 Y. Wang, M. Hu, Y. Wang, J. Zheng, D. Shang, Y. Yang, Y. Liu, X. Li, R. Tang, W. Zhu, Z. Du, Y. Wu, S. Guo, Z. Wu, S. Lou, M. Hallquist and J. Yu, *Atmos. Chem. Phys.*, 2019, **19**, 7649–7665.
- 24 J. Ahad, R. Macdonald, J. Parrott, Z. Yang, Y. Zhang, T. Siddique, A. Kuznetsova, C. Rauert, E. Galarneau, W. Studabaker, M. Evans, M. McMaster and D. Shang, *Environ. Pollut.*, 2020, **266**, 114988.
- 25 D. Traversi, R. Degan, R. Marco, G. Gilli, C. Pignata, S. Villani and R. Bono, *Environ. Int.*, 2009, **35**, 905–910.
- 26 K. Ju and R. Parales, *Microbiol. Mol. Biol. Rev.*, 2010, **74**, 250–272.
- 27 P. Kovacic and R. Somanathan, *J. Appl. Toxicol.*, 2014, **34**, 810–824.
- 28 C. Salvador, R. Tang, M. Priestley, L. Li, E. Tsiligiannis, M. Breton, W. Zhu, L. Zeng, H. Wang, Y. Yu, M. Hu, S. Guo and M. Hallquist, *Atmos. Chem. Phys.*, 2021, **21**, 1389–1406.
- 29 A. Kroflič, J. Anders, I. Drventić, P. Mettke, O. Böge, A. Mutzel, J. Kleffmann and H. Herrmann, *ACS Earth Space Chem.*, 2021, **5**, 1083–1093.
- 30 Z. Finewax, J. Gouw and P. Ziemann, *Environ. Sci. Technol.*, 2018, **52**, 1981–1989.
- 31 M. Harrison, S. Barra, D. Borghesi, D. Vione, C. Arsene and R. Olariu, *Atmos. Environ.*, 2005, **39**, 231–248.
- 32 S. Wang and H. Li, *Environ. Sci. Technol.*, 2021, **55**, 2899–2907.
- 33 R. Hems and J. Abbatt, *ACS Earth Space Chem.*, 2018, **2**, 225–234.
- 34 J. Liu, P. Lin, A. Laskin, J. Laskin, S. Kathmann, M. Wise, R. Caylor, F. Imholt, V. Selimovic and J. Shilling, *Atmos. Chem. Phys.*, 2016, **16**, 12815–12827.
- 35 C. Mohr, F. Lopez-Hilfiker, P. Zotter, A. Prévot, L. Xu, N. Ng, S. Herndon, L. Williams, J. Franklin, M. Zahniser, D. Worsnop, W. Knighton, A. Aiken, K. Gorkowski, M. Dubey, J. Allan and J. Thornton, *Environ. Sci. Technol.*, 2013, **47**, 6316–6324.
- 36 P. Lin, N. Bluvshstein, Y. Rudich, S. Nizkorodov, J. Laskin and A. Laskin, *Environ. Sci. Technol.*, 2017, **51**, 11561–11570.
- 37 F. Ikemori, T. Nakayama and H. Hasegawa, *Atmos. Environ.*, 2019, **211**, 91–102.
- 38 M. Claeys, R. Vermeylen, F. Yasmeeen, Y. Gómez-González, X. Chi, W. Maenhaut, T. Mészáros and I. Salma, *Environ. Chem.*, 2012, **9**, 273–284.
- 39 M. Li, X. Wang, C. Lu, R. Li, J. Zhang, S. Dong, L. Yang, L. Xue, J. Chen and W. Wang, *Sci. Total Environ.*, 2020, **714**, 136760.
- 40 X. Li, L. Jiang, L. Hoa, Y. Lyu, T. Xu, X. Yang, Y. Iinuma, J. Chen and H. Herrmann, *Atmos. Environ.*, 2016, **145**, 115–127.
- 41 J. Luttke, V. Scheer, K. Levens, G. Wünsch, J. Cape, K. Hargreaves, R. Storeton-West, K. Acker, W. Wiprecht and B. Jones, *Atmos. Environ.*, 1997, **31**, 2637–2648.
- 42 Y. Zhang, L. Müller, R. Winterhalter, G. Moortgat, T. Hoffmann and U. Pöschl, *Atmos. Chem. Phys.*, 2010, **10**, 7859–7873.
- 43 W. Yuan, R. Huang, L. Yang, T. Wang, J. Duan, J. Guo, H. Ni, Y. Chen, Q. Chen, Y. Li, U. Dusek, C. O'Dowd and T. Hoffmann, *Atmos. Chem. Phys.*, 2021, **21**, 3685–3697.
- 44 M. Harrison, M. Heal and J. Cape, *Atmos. Chem. Phys.*, 2005, **5**, 1679–1695.
- 45 Y. Zhang, L. Muller, R. Winterhalter, G. Moortgat, T. Hoffmann and U. Pöschl, *Atmos. Chem. Phys. Discuss.*, 2010, **10**, 13253–13286.
- 46 M. Sangwan and L. Zhu, *J. Phys. Chem. A*, 2016, **120**, 9958–9967.
- 47 M. Sangwan and L. Zhu, *J. Phys. Chem. A*, 2018, **122**, 1861–1872.
- 48 J. Chen, J. Wenger and D. Venables, *J. Phys. Chem. A*, 2011, **115**, 12235–12242.
- 49 I. Bejan, Y. Aal, I. Barnes, T. Benter, B. Bohn, P. Wiesen and J. Kleffmann, *Phys. Chem. Chem. Phys.*, 2006, **8**, 2028–2035.
- 50 Y. Nitta, O. Schalk, H. Igarashi, S. Wada, T. Tsutsumi, K. Saita, T. Taketsugu and T. Sekikawa, *J. Phys. Chem. Lett.*, 2021, **12**, 674–679.
- 51 F. Barsotti, T. Bartels-Rausch, E. Laurentis, M. Ammann, M. Brigante, G. Mailhot, V. Maurino, C. Minero and D. Vione, *Environ. Sci. Technol.*, 2017, **51**, 7486–7495.
- 52 W. Yang, D. You, C. Li, C. Han, N. Tang, H. Yang and X. Xue, *Environ. Sci. Technol. Lett.*, 2021, **8**, 747–752.
- 53 S. Cheng, C. Zhou, H. Yin, J. Sun and K. Han, *J. Chem. Phys.*, 2009, **130**, 234311.
- 54 L. Vereecken, H. K. Chakravarty, B. Bohn and J. Lelieveld, *Int. J. Chem. Kinet.*, 2016, **48**, 785–795.



- 55 K. Andersson, P. Malmqvist, B. Roos, A. Sadlej and K. Wolinski, *J. Phys. Chem.*, 1990, **94**, 5483–5488.
- 56 K. Andersson, P. Malmqvist and B. Roos, *J. Chem. Phys.*, 1992, **96**, 1218–1226.
- 57 C. Xu, F. Gu and C. Zhu, *Phys. Chem. Chem. Phys.*, 2018, **20**, 5606–5616.
- 58 C. Xu, L. Yu, C. Zhu, J. Yu and Z. Cao, *Sci. Rep.*, 2016, **6**, 26768.
- 59 C. Xu, L. Yu, C. Zhu and J. Yu, *J. Phys. Chem. A*, 2015, **119**, 10441–10450.
- 60 G. Ghigo, B. Roos and P. Malmqvist, *Chem. Phys. Lett.*, 2004, **396**, 142–149.
- 61 N. Forsberg and P. Malmqvist, *Chem. Phys. Lett.*, 1997, **274**, 196–204.
- 62 F. Weigend and R. Ahlrichs, *Phys. Chem. Chem. Phys.*, 2005, **7**, 3297–3305.
- 63 F. Neese, *Wiley Interdiscip. Rev.: Comput. Mol. Sci.*, 2012, **2**, 73–78.
- 64 F. Neese, *Wiley Interdiscip. Rev.: Comput. Mol. Sci.*, 2017, **8**, e1327.
- 65 T. Lu, *The sobMECP Program*, 2016, website: <https://sobereva.com/286>, date of access: 12/05/2022.
- 66 J. Harvey, M. Aschi, H. Schwarz and W. Koch, *Theor. Chem. Acc.*, 1998, **99**, 95–99.
- 67 A. D. Becke, *J. Chem. Phys.*, 1993, **98**, 5648–5652.
- 68 C. Lee, W. Yang and R. G. Parr, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1998, **37**, 785–789.
- 69 S. H. Yosko, L. Wilk and M. Nusair, *Can. J. Phys.*, 1980, **58**, 1200–1211.
- 70 A. D. McLean and G. S. Chandler, *J. Chem. Phys.*, 1980, **72**, 5639–5848.
- 71 K. Raghavachari, J. S. Binkley, R. Seeger and J. A. Pople, *J. Chem. Phys.*, 1980, **72**, 650–654.
- 72 T. Clark, J. Chandrasekhar, G. W. Spitznagel and P. v. R. Schleyer, *J. Comput. Chem.*, 1983, **4**, 294–301.
- 73 M. J. Frisch, J. A. Pople and J. S. Binkley, *J. Chem. Phys.*, 1984, **80**, 3265–3269.
- 74 M. Frisch, *et al.*, *Gaussian 09, Revision D.01*, Gaussian, Inc., Wallingford, CT, 2010.
- 75 M. Marques, C. Ullrich, F. Nogueira, A. Rubio, K. Burke and E. Gross, *Time-dependent Density Functional Theory*, Springer, Berlin, 2006.
- 76 X. Gao, S. Bai, D. Fazzi, T. Niehaus, M. Barbatti and W. Thiel, *J. Chem. Theory Comput.*, 2017, **13**, 515.
- 77 G. Cui and W.-H. Fang, *ChemPhysChem*, 2011, **12**, 1351–1357.
- 78 K. Takahashi, K. L. Plath, R. T. Skodje and V. Vaida, *J. Phys. Chem. A*, 2008, **112**, 7321–7331.
- 79 A. W. Harrison, M. F. Shaw and W. J. De Bruyn, *J. Phys. Chem. A*, 2019, **123**, 8109–8121.
- 80 N. E. Henriksen and F. Y. Hansen, Chapter 7: Unimolecular Reactions, *Theories of Molecular Reaction Dynamics: the Microscopic Foundation of Chemical Kinetics*, Oxford University Press, 2008.
- 81 T. Baer and W. L. Hase, *Unimolecular Reaction Dynamics: Theory and Experiments*, Oxford University Press, 1996.
- 82 P. A. M. Dirac, *Proc. R. Soc. London, Ser. A*, 1927, **114**, 243–265.
- 83 V. Lawetz, G. Orlandi and W. Siebrand, *J. Chem. Phys.*, 1972, **56**, 4058–4072.
- 84 G. W. Robinson and R. P. Frosch, *J. Chem. Phys.*, 1963, **38**, 1187–1203.
- 85 E. Y. T. Li, T. Y. Jiang, Y. Chi and P.-T. Chou, *Phys. Chem. Chem. Phys.*, 2014, **16**, 26184–26192.
- 86 K. Schmidt, S. Brovelli, V. Coropceanu, D. Belijonne, J. Cornil, C. Bazzini, T. Caronna, F. Meinardi, Z. Shuai and J.-L. Bredas, *J. Phys. Chem. A*, 2007, **111**, 10490–10499.
- 87 X.-F. Song, Z.-W. Li, W.-K. Chen, Y.-J. Gao and G. Cui, *Inorg. Chem.*, 2022, **61**, 7673–7681.
- 88 X.-F. Song, L.-Y. Peng, W.-K. Chen, Y.-J. Gao, W.-H. Fang and G. Cui, *Chem.-Eur. J.*, 2022, e202201782.
- 89 X.-W. Sun, L.-Y. Peng, Y.-J. Gao, Q. Fang and G. Cui, *J. Phys. Chem. A*, 2022, **126**, 4176–4184.
- 90 K. Shizu and H. Kaji, *J. Phys. Chem. A*, 2021, **125**, 9000–9010.
- 91 J. L. Steinfeld, J. S. Francisco and W. L. Hase, *Chemical Kinetics and Dynamics*, Prentice-Hall, 2nd edn, 1999.
- 92 H. Eyring, *J. Chem. Phys.*, 1935, **3**, 107–115.
- 93 E. Wigner, *Z. Phys. Chem., Abt. B*, 1932, **19**, 203–216.
- 94 P. Grammaticakis, *Bull. Soc. Chim. Fr.*, 1951, **18**, 220–226.
- 95 R. Haddon, *J. Am. Chem. Soc.*, 1980, **102**, 1807–1811.
- 96 P. Nagy, *Int. J. Mol. Sci.*, 2014, **15**, 19562–19633.
- 97 K. Mackeprang, S. Schröder and H. Kjaergaard, *Chem. Phys. Lett.*, 2013, **582**, 31–37.
- 98 J. Ladarević, B. Božić, L. Matović, *et al.*, *Dyes Pigm.*, 2019, **162**, 562–572.
- 99 H. Ernst, T. Wolf, O. Schalk, N. Gonzalez-García, A. Boguslavskiy, A. Stolow, M. Olzmann and A. Unterreiner, *J. Phys. Chem. A*, 2015, **119**, 9225–9235.
- 100 S. Caumo, M. Claeys, W. Maenhaut, R. Vermeylen, S. Behrouzi, M. Shalamzari and P. Vasconcellos, *Atmos. Environ.*, 2016, **145**, 272–279.
- 101 X. Li, L. Jiang, L. Hoa, Y. Lyu, T. Xu, X. Yang, Y. Iinuma, J. Chen and H. Herrmann, *Atmos. Environ.*, 2016, **145**, 115–127.
- 102 E. Eckstein, D. Perner, Ch. Brühl and T. Trautmann, *Atmos. Chem. Phys. Discuss.*, 2002, **2**, 1939–1977.
- 103 Y. Kanaya, Y. Kajii and H. Akimoto, *Atmos. Environ.*, 2003, **37**, 2463–2475.

