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Experimental and Theoretical Study on the OH-Induced Degradation of Piperazine under Simulated Atmospheric Conditions

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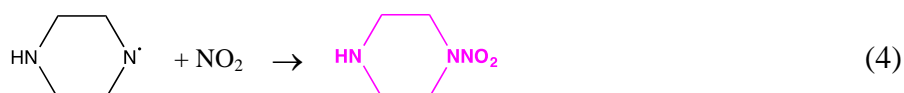
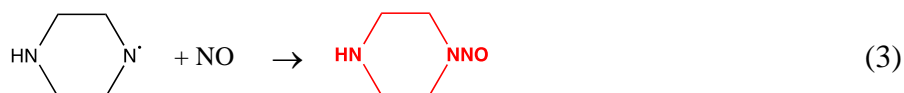
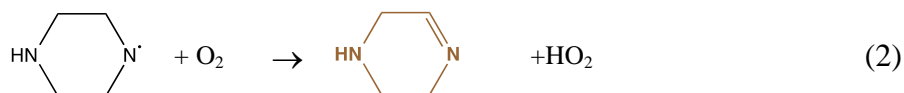
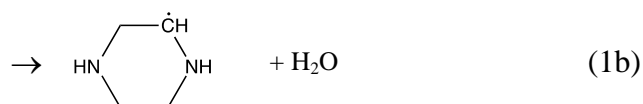
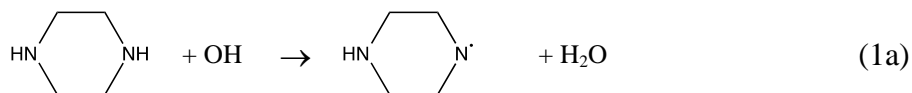
ABSTRACT

The OH-initiated photo-oxidation of piperazine was investigated in chamber experiments at the European Photoreactor (EUPHORE) in Spain and by quantum chemistry methods. The rate coefficient for reaction with OH radicals was determined by the relative rate method to be $k_{\text{OH-piperazine}} = (2.8 \pm 0.6) \times 10^{-10} \text{ cm}^3 \text{ molecule}^{-1} \text{ s}^{-1}$ at $307 \pm 2 \text{ K}$ and $1014 \pm 2 \text{ hPa}$. The reaction was found to proceed via both C-H and N-H abstraction, the latter resulting in the formation of nitroso- and nitro-piperazine. The reaction in the EUPHORE chamber was accompanied by strong particle formation which was induced by an acid-base reaction between photochemically formed nitric acid and the

1 INTRODUCTION

Piperazine (1,4-diazacyclohexane, PZ) is among the amines considered for use in large-scale Carbon capture (CC) to reduce CO₂ emissions from industrial point sources.¹ Tests at the Technology Centre Mongstad (Norway) have established that it is difficult to avoid ppm-level emissions of amines and their process degradation products to the environment during operation of a large-scale capture plant² - the concern being the carcinogenic nitrosamines and nitramines either directly emitted or resulting in the subsequent atmospheric photo-oxidation of the emitted amines.³ The major removal processes of PZ in the atmosphere are uptake in aqueous particles and gas phase reaction with OH radicals during day-time and NO₃ radicals during night-time. The OH rate coefficient for reaction with PZ was recently reported to be $\sim 2.4 \times 10^{-10} \text{ cm}^3 \text{ molecules}^{-1} \text{ s}^{-1}$ at 298 K and to favor C-H abstraction: $k_{\text{N-H}}/(k_{\text{N-H}} + k_{\text{C-H}}) = 0.09 \pm 0.06$.⁴

The PZ nitrosamine (PZ-NO) and nitramine (PZ-NO₂) are both carcinogenic;³ they result in the following sequence of atmospheric gas phase reactions:⁵



Although the O₂ reaction with N-centered amino radicals, R₁R₂N[•], is reported to be around 6 orders of magnitude slower than the corresponding NO and NO₂ reactions,⁶ it is still dominating at most atmospheric conditions, and 1-nitrosopiperazine (PZ-NO) and 1-nitropiperazine (PZ-NO₂) will only be minor products in the natural atmospheric photo-oxidation of PZ. Both compounds were observed, but not quantified, in previous piperazine photo-oxidation experiments in the $\sim 200 \text{ m}^3$ EUPHORE atmospheric chamber,⁷ and in the more recent experiments employing the $\sim 18 \text{ m}^3$ CSIRO indoor smog chamber.⁸

The open literature includes two theoretical studies on the kinetics of the hydrogen abstraction from PZ by OH radicals, in which the branching between the N-H and C-H abstractions 1a and 1b was predicted to be 0.07⁹ and 0.01¹⁰, respectively, at 298 K. The latter theoretical study also includes an investigation of the atmospheric degradation following the initial H-abstraction.

In the present communication we report results from a new series of PZ and PZ-NO₂ photo-oxidation and PZ-NO photolysis experiments in the EUPHORE chamber as well as a quantum chemistry based evaluation of the primary products formed in the OH initiated photo-oxidation of PZ.

2 MATERIALS AND METHODS

2.1 Experimental Methods and Chemicals

A series of experiments was carried out in chamber B of the EUPHORE facility in Valencia, Spain. The facility and analytical methods have recently been reported in detail;¹¹ additional information specific to the present work is found in the [Supporting Information](#).

PZ (Sigma-Aldrich, ReagentPlus®, 99%), acetonitrile (Sigma-Aldrich, LiChrosolv®, ≥99.8%), pyrrole (Sigma-Aldrich, 98 %), 1,3,5-trimethylbenzene (Sigma-Aldrich, 98 %), isoprene (Sigma-Aldrich, analytical standard) and limonene (R-(+), Sigma-Aldrich, analytical standard) were used without further purification. 2-propyl nitrite (isopropyl nitrite, IPN) was synthesized from isopropanol, hydrochloric acid and sodium nitrite, and purified by repeated washing with ice water. The 1:1 nitric acid salt of PZ was prepared by adding a small excess of diluted nitric acid (HNO₃) to diluted PZ followed by rotary evaporation to dryness at 80 °C. Synthesis of 1-nitrosopiperazine and 1-nitropiperazine is described in the [Supporting Information](#).

2.2 Computational Methods

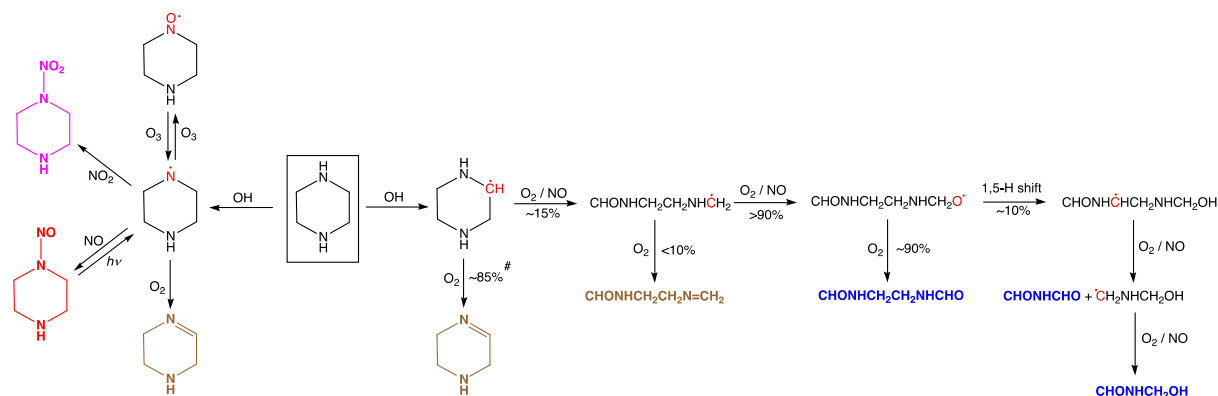
Stationary points on the potential energy surfaces for the atmospheric degradation of piperazine were characterized in M06-2X¹² calculations employing the aug-cc-pVTZ¹³⁻¹⁴ basis set. Pre and post reaction complexes were located by following the intrinsic reaction coordinate¹⁵⁻¹⁸ from the saddle points. Reaction enthalpies and proton affinities were calculated using the G4 model chemistry.¹⁹ Additional dipole moments and isotropic polarizabilities, serving as input to prediction of ion-molecule reaction rate coefficients,²⁰ were obtained in B3LYP/aug-cc-pVTZ calculations, see [Table S1](#) in the [Supporting Information](#). All calculations were performed with Gaussian 09.²¹

Master equation calculations were carried out using the program MESMER 3.0²² (Master Equation Solver for Multi-Energy-well Reactions) to simulate the reactions at atmospheric conditions. The required input parameters for molecules, intermediate species and products were obtained from the ab initio calculations.

3 RESULTS

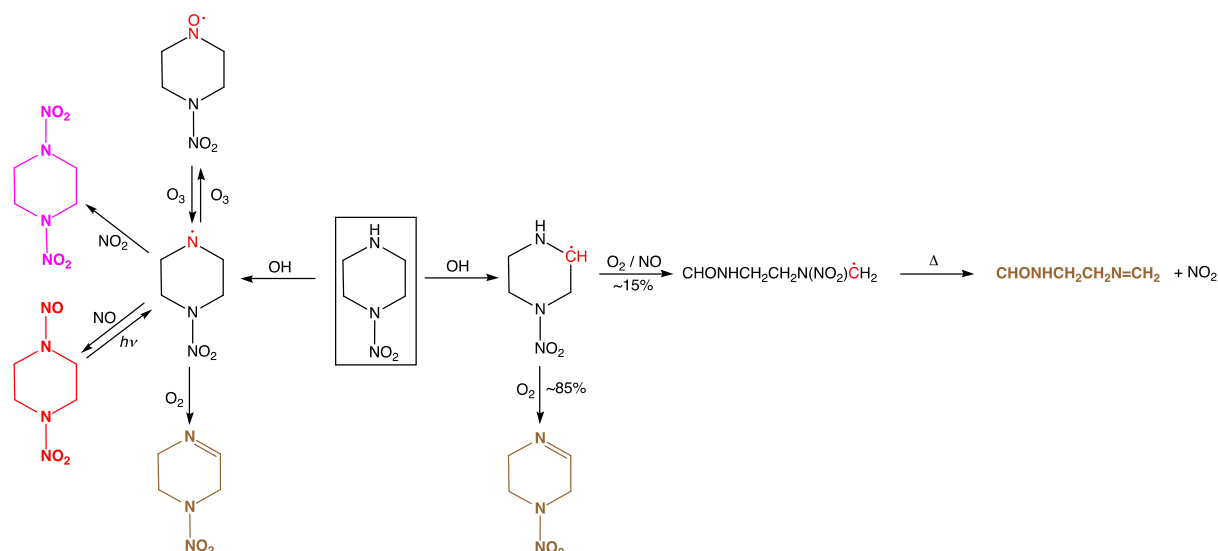
3.1 Computational results

The kinetics of the initial step in the PZ + OH reaction is complicated by PZ existing in two conformations close in energy; a detailed theoretical account of the kinetics is outside the scope of the present work and will be presented elsewhere.²³ Details of the quantum chemistry study of the OH radical initiated atmospheric PZ photo-oxidation are collected in the [Supporting Information](#) (including illustrations of the pivotal potential energy surfaces, [Figures S1 – S5](#), and the associated [Tables S2 – S5](#) containing energies, Cartesian coordinates and vibration-rotation data). The quantum chemistry prediction of the major routes in the atmospheric degradation of piperazine are summarized in [Scheme 1](#). The degradation routes largely concord with those established in previous dimethylamine^{6, 24-25} and diethylamine^{7, 24} photo-oxidation experiments. A recent theoretical study on the OH initiated degradation of piperazine employing the G4 chemistry model did not include the atmospheric fate of the piperazine aminyl radical.¹⁰ The present results also differ with respect to the atmospheric reactions following C-H abstraction from PZ; we find the cyclic alkoxy radical, that ultimately follows C-H abstraction, to be metastable resulting in spontaneous ring opening, and that the major fraction of the resulting CHONHCH₂CH₂NHĊH₂ radical will end up as a diamide. The branching between ring-opening and formation of 1,2,3,6-tetrahydropyrazine (PZ-Im) depends on the NO mixing ratio; master equation calculations employing 5 ppb of NO results in a 15 : 85 branching, whereas a 35 : 65 branching is obtained with 50 ppb of NO.



Scheme 1. Quantum chemistry prediction of the major primary products in the atmospheric photo-oxidation of piperazine. [#] The branching between ring-opening and formation of 1,2,3,6-tetrahydropyrazine (PZ-Im) depends on the NO mixing ratio.

The atmospheric fate of PZ-NO₂ was also investigated, and the major photo-oxidation routes for PZ-NO₂, outlined in Scheme 2, parallel those of PZ (Scheme 1) with one exception – the alkyl-radical formed upon ring-opening ejects NO₂ forming an imine. Details of the quantum chemistry study of the OH radical initiated atmospheric PZ-NO₂ photo-oxidation are collected in the Supporting Information.



Scheme 2. Quantum chemistry prediction of the major primary products in the atmospheric photo-oxidation of 1-nitropiperazine (PZ-NO₂).

3.2 Experimental results

We first report results from new kinetic studies of the PZ + OH reaction. We then present results from PZ-NO photolysis and from PZ-NO₂ photo-oxidation experiments facilitating interpretation of the PZ photo-oxidation experiments. Finally, we present results from studies of the aerosol formed in the PZ photo-oxidation experiments.

3.2.1 Piperazine + OH reaction kinetics

Two relative rate experiments were carried out in the EUPHORE chamber B in which isoprene, limonene, 1,3,5-trimethylbenzene and pyrrole were used as reference compounds. Acetonitrile was added as inert tracer to monitor the apparent dilution by purified air that is constantly added to compensate for leakage and continuous sampling by the air monitors ($k_{\text{OH}+\text{CH}_3\text{CN}} = 2.2 \times 10^{-14} \text{ cm}^3 \text{ molecule}^{-1} \text{ s}^{-1}$ at 298 K). ²⁶ Figure 1 displays the time evolution

of compound-specific PTR-ToF-MS ion signals measured during the second experiment (the first experiment is documented in [Figure S6, Supporting Information](#)). The dilution rate due to air replenishment was $8.6 \times 10^{-6} \text{ s}^{-1}$ in the two experiments; PZ wall loss rates (derived from the reagent decay prior to adding the OH precursor IPN) ranged from 1 to $4 \times 10^{-5} \text{ s}^{-1}$. Initial mixing ratios were ~ 100 ppb for the reference compounds and ~ 200 ppb for PZ. Average OH densities in the EUPHORE chamber during the periods selected for analyses (9:10 – 9:30 and 14:10 – 14:35 UTC) were around $3 \times 10^6 \text{ cm}^{-3}$; average pressure and temperature in the two experiments were 1014 ± 2 mbar and 307 ± 2 K. The temporal profile of PZ recorded by PTR-ToF-MS matches well the one obtained by a home-built high temperature PTR-MS, indicating an adequate rinstrument esponse time ([Figure S7 in the Supporting Information](#)).

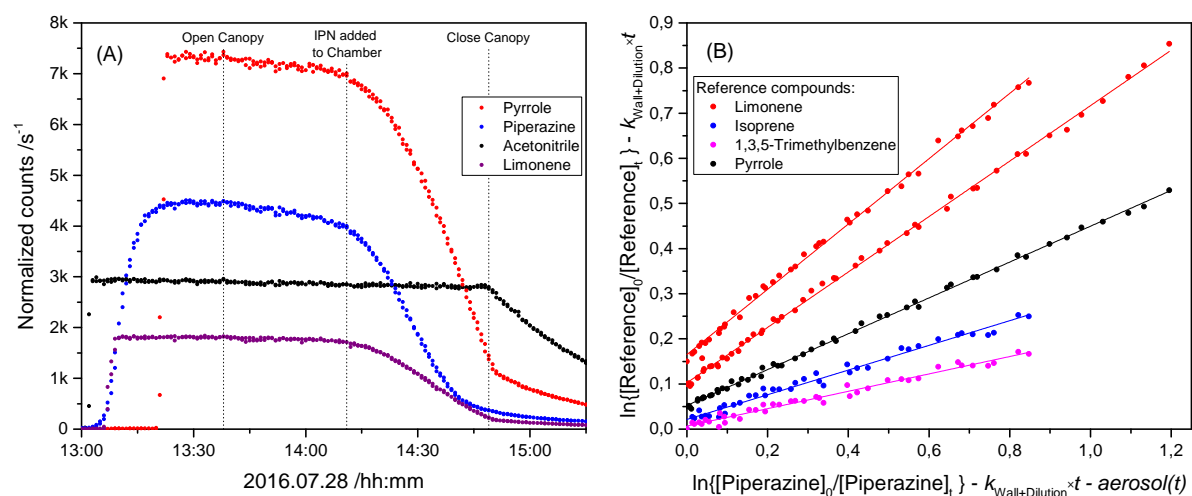


Figure 1. (A): Time evolution of the acetonitrile, pyrrole, piperazine and limonene ion signals at m/z 42.034, 68.050, 87.092 and 137.133, respectively, during the second kinetic experiment on 2016.07.28. (B): Relative rate plot showing the decays of isoprene, limonene, pyrrole and piperazine at 1014 hPa and 307 K in the presence OH radicals. For the sake of clarity the data have been displaced along the abscissa. The data have been corrected for dilution due to chamber air replenishment, for wall loss and for loss to the aerosol, see [Supporting Information](#).

A least-squares fitting of the wall- and dilution loss-corrected data ([Figure S8 in the Supporting Information](#)) results in an average $k_{\text{OH+Piperazine}} = (3.0 \pm 0.6) \times 10^{-10} \text{ cm}^3 \text{ molecule}^{-1} \text{ s}^{-1}$ at 307 ± 2 K and 1014 ± 2 hPa. Significant amounts of PZ are, however, transferred from the gas to the particle phase during the periods selected for analysis. [Figures S9 and S10 \(Supporting Information\)](#) show the time evolutions of aerosol mass and the PZ content, and approximately 6.3 and 1.2 % of PZ were actually lost to the aerosol during the two kinetic experiments. Correction for PZ loss to particles during the kinetic experiments was therefore

implemented in the data analysis ([Supporting Information](#)) resulting in an average $k_{\text{OH+Piperazine}} = (2.8 \pm 0.6) \times 10^{-10} \text{ cm}^3 \text{ molecule}^{-1} \text{ s}^{-1}$ at $307 \pm 2 \text{ K}$ and $1014 \pm 2 \text{ hPa}$. The present result agrees well with those of Onel et al.,⁴ who reported $k(T) = (2.37 \pm 0.03) \times 10^{-10} (T/298)^{-(1.76 \pm 0.08)}$ and $k = (2.25 \pm 0.28) \times 10^{-10} \text{ cm}^3 \text{ molecule}^{-1} \text{ s}^{-1}$ at 307 K from flash photolysis/resonance fluorescence experiments.

3.2.2 1-Nitropiperazine photo-oxidation studies

The atmospheric fate of PZ-NO₂ was investigated in two photo-oxidation experiments – one under high NO₂ levels, the other under high NO levels. [Figure S17](#) in the [Supporting Information](#) shows time profiles of the relevant PTR-ToF-MS ion signals observed during a photo-oxidation experiment demonstrating that protonated PZ-NO₂ fragments severely at the instrumental settings employed: 15% m/z 132.076 (protonated molecule), 38% m/z 86.085 (NO₂ ejection), 30% m/z 85.080 (HNO₂ ejection), 4% m/z 57.059 (ring fragment), and 13% m/z 44.052 (ring fragment). The observed ion signals related to PZ-NO₂ photo-oxidation are summarized in [Table S7](#) in the [Supporting Information](#).

PZ-NO₂ is a very “sticky” compound that transfers relatively fast to particles and wall surfaces ([Figure S17](#)) making quantitative conclusions arduous. In line with the extensive fragmentation of protonated PZ-NO₂, most of the ion signals observed during the photo-oxidation corresponds to fragments. It is, however, clear that 1,2-dinitropiperazine (m/z 177.055 ([PZ-(NO₂)₂]⁺H⁺) and fragment ions) is a major product, and that also 1-nitro-4-nitrosopiperazine (m/z 161.064 ([NO₂-PZ-NO]⁺H⁺) and fragment ions) is formed in the experiments. Surprisingly, the expected key product, 1-nitro-1,2,3,6-tetrahydropyrazine (NO₂-PZ-Im), is not detected as the protonated ion at m/z 130.061. Assuming a similar fragmentation of protonated PZ-Im as observed for protonated PZ-NO₂, fragment ions are expected at m/z 84.069 (NO₂ ejection), 83.061 (HNO₂ ejection), 55.042 (CH₂CH₂NHCH⁺), and 42.034 (CH₂CH₂N⁺). Of these, the m/z 83.061, 55.042 and 42.034 are observed.

3.2.3 1-Nitrosopiperazine photolysis studies

Nitrosamines have a characteristic $n \rightarrow \pi^*$ transition in the UV-A region and photolyse rapidly in natural sunlight; the quantum yield to photo-dissociation of (CH₃)₂NNO following S₀ → S₁($n\pi^*$) excitation at 363.5 nm was reported to be 1.03 ± 0.10 ,²⁷⁻²⁸ and theory shows that the excited S₁ state is repulsive leading to swift dissociation following excitation.²⁹ Previous experiments at EUPHORE show that nitroso-dimethylamine, -diethylamine, –

methylethylamine, –methylpropylamine, –methylbutylamine, –morpholine, –piperazine, –piperidine and –pyrazine all photolyze equally fast with $j_{\text{Nitrosamine}}/j_{\text{NO}_2} = 0.34 \pm 0.04$.^{7, 30}

Three photolysis experiments were carried out in the EUPHORE chamber B under conditions that did not allow quantification of the solar radiation. Cyclohexane was added to the chamber (~2 ppm) to enable an estimate of the amount of OH radicals formed following PZ-NO photolysis; the derived OH radical mixing ratio varied between 1 and $4 \times 10^5 \text{ cm}^{-3}$ (for details, see the [Supporting Information](#)). Taking $k_{\text{PZ-NO+OH}} \approx k_{\text{PZ+OH}}$ as a conservative estimate of the PZ-NO reactivity towards OH places an upper level of 2% PZ-NO removal by OH radicals during the photolysis.

[Table 1](#) summarises the observed PTR-TOF-MS ion signals relevant to the PZ-NO photolysis experiments; a more complete list of ion signals observed in the present experiments is found in [Table S6](#) in the [Supporting Information](#), which also includes data obtained in our previous studies.⁷ [Figure S15](#) in the [Supporting Information](#) illustrates the raw ion signal time profiles; the ion signals growing in upon photolysis fall in 2 categories: (1) the m/z 44.052, 83.061, 85.080, 86.085 and 132.074 having time profiles typical of primary products, and (2) the less intense m/z 46.030, 74.024, 81.045 and 97.045 with time profiles more resembling those of secondary products.

Table 1. Selected ion signals observed during 1-nitrosopiperazine photolysis experiments.

m/z	Ion sum formula	Interpretation
44.052	$\text{C}_2\text{H}_6\text{N}^+$	Fragment of $[\text{PZ-NO}]\text{H}^+$, $[\text{PZ-NO}_2]\text{H}^+$ and $[\text{PZ-Im}]\text{H}^+$
83.061	$\text{C}_4\text{H}_7\text{N}_2^+$	H_2 elimination from $[\text{PZ-Im}]\text{H}^+$
85.080	$\text{C}_4\text{H}_9\text{N}_2^+$	Fragment of $[\text{PZ-Im}]\text{H}^+$, $[\text{PZ-NO}]\text{H}^+$, $[\text{PZ-NO}_2]\text{H}^+$
86.085	$\text{C}_4\text{H}_{10}\text{N}_2^+$	Fragment of $[\text{PZ-NO}]\text{H}^+$, $[\text{PZ-NO}_2]\text{H}^+$
116.079	$\text{C}_4\text{H}_{10}\text{N}_3\text{O}^+$	$[\text{PZ-NO}]\text{H}^+$
132.074	$\text{C}_4\text{H}_{10}\text{N}_3\text{O}_2^+$	$[\text{PZ-NO}_2]\text{H}^+$

$[\text{PZ-NO}]\text{H}^+$ fragmentation: m/z 116.079 (68% protonated molecule); m/z 86.085 (11% NO ejection); m/z 85.080 (17% HNO ejection); m/z 44.052 (4% ring scission). $[\text{PZ-NO}_2]\text{H}^+$ fragmentation: m/z 132.076 (17% protonated molecule); m/z 86.085 (40% NO_2 ejection); m/z 85.080 (32% HNO_2 ejection); m/z 44.052 (11% ring scission).

Two primary products are expected following nitrosamine photolysis: the imine (PZ-Im, 1,2,3,6-tetrahydropyrazine) and the nitramine PZ- NO_2 , [Scheme 1](#). Attempts to synthesise PZ-Im were futile, and fragmentation of protonated PZ-NO and PZ- NO_2 complicates the unambiguous identification of PZ-Im by PTR-TOF-MS. Inspection of the ion signals observed in the PZ-NO photolysis experiments during the time period before opening the chamber canopy (see [Figure S11](#)) reveals the $[\text{PZ-NO}]\text{H}^+$ fragmentation; the fragmentation of $[\text{PZ-NO}_2]\text{H}^+$ was determined in separate experiments with the same instrument parameters,

see later. In summary, the ion signals at m/z 44.052, 85.080 and 86.085 all have an origin in both PZ-NO and PZ-NO₂, [Table 1](#). That is, the expected ion signal of protonated PZ-Im at m/z 85.080 (C₄H₉N₂⁺) is also a major fragment of both protonated PZ-NO and PZ-NO₂.

The PZ-NO and PZ-NO₂ fragmentation corrected ion signals are presented in [Figure 2](#), from which it can be seen that there remains significant ion intensities at m/z 44.052 and 85.080, whereas m/z 86.085 is fully accounted for by PZ-NO and PZ-NO₂. The m/z 44.052 (C₂H₆N⁺) is obviously also a ring scission fragment of [PZ-Im]H⁺ and m/z 83.061 (C₄H₇N₂⁺) is explained by H₂-loss from [PZ-Im]H⁺.

[Figure 2](#) also includes the derived volume mixing ratios of PZ-NO, PZ-NO₂ and PZ-Im. The mass balance in the photolysis experiments is only around 60% and the discrepancy is primarily due to “missing” PZ-Im, which in part can be explained by partitioning to surfaces and subsequent hydrolysis and/or condensation reactions. The corresponding CHARON PTR-TOF-MS results (see later) suggests a minor presence of PZ-imines in co-existing aerosols.

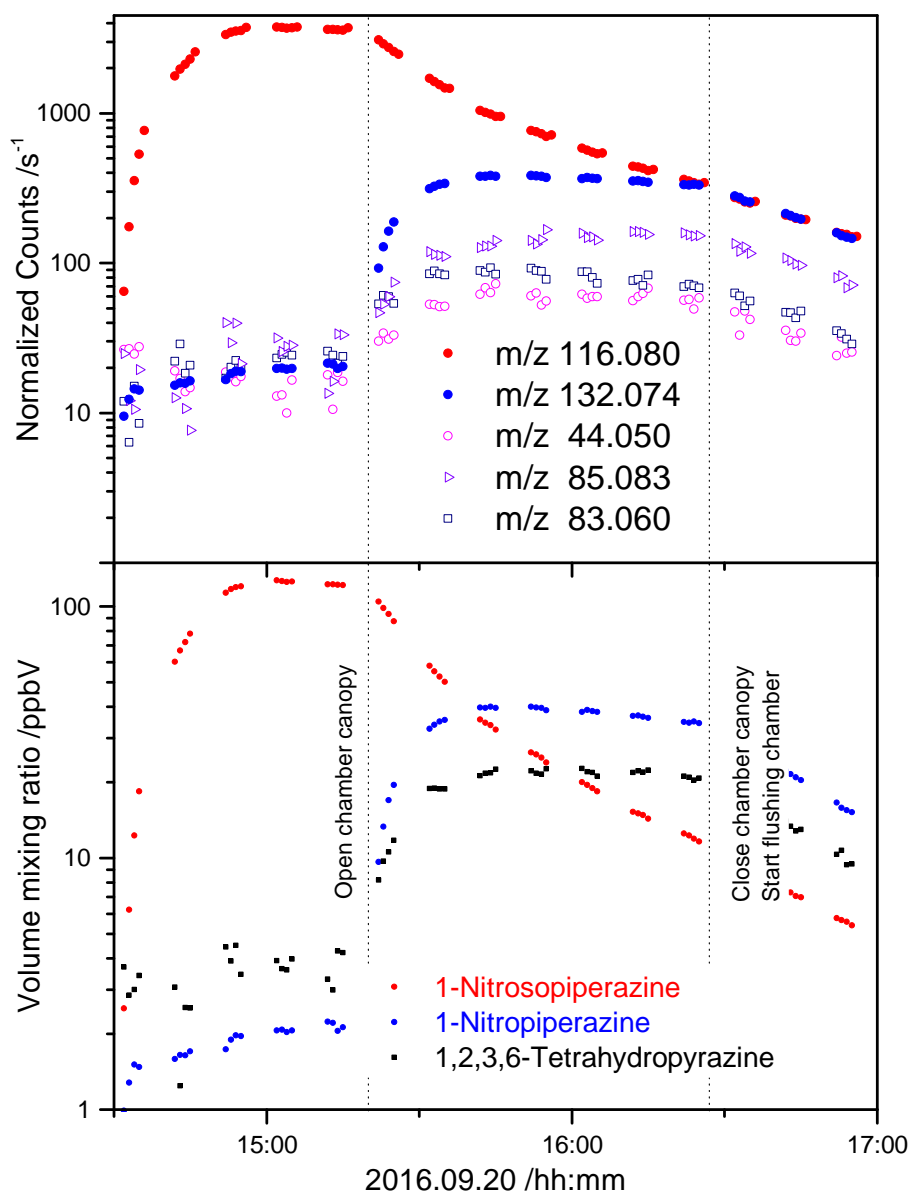


Figure 2. Top: Fragmentation-corrected normalized ion signals during the 1-nitrosopiperazine photolysis experiment on 2016.09.20. Ion signals increasing by less than 2% of the m/z 116.080 decrease have been omitted for the sake of clarity. Bottom: Derived volume mixing ratios (ppbV) of 1-nitroso-, 1-nitropiperazine and 1,2,3,6-tetrahydropyrazine during the experiment.

3.2.4 Piperazine photo-oxidation studies

Previous PZ photo-oxidation experiments in the EUPHORE chamber were severely affected by wall adsorption/desorption and particle formation,⁷ and a new series of piperazine photo-oxidation experiments was carried out under different, warmer conditions reducing wall adsorption (see Table S8 in the Supporting Information). Piperazine, dissolved in water, was injected into the chamber in a stream of N_2 . The detected ion signals can be separated according to their time evolution: (1) signals that appear upon injection of piperazine along with that of protonated piperazine, (2) signals that grow and decrease again

during the photo-oxidation experiment (intermediate products), and (3) signals that grow steadily after opening the chamber canopy. Figure 3 exemplifies the observed time evolution of PZ and the major products during a photo-oxidation experiment; the ion signals observed in the new as well as in the previous experiments are collected in Table S9.

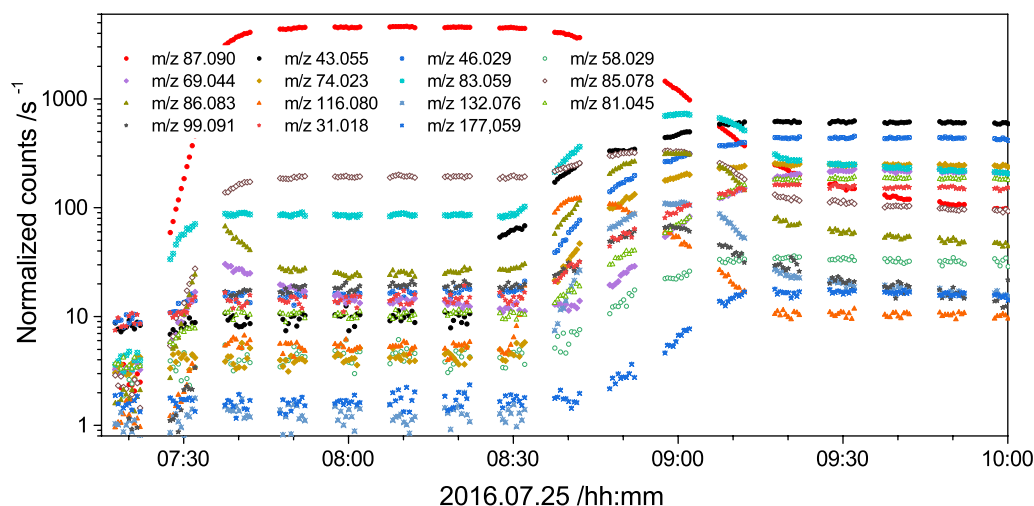


Figure 3. Time evolution of ion signals during the piperazine photo-oxidation experiment on 2016.07.25. Ion signals increasing by less than 2% of the m/z 87.090 decrease have been omitted for the sake of clarity.

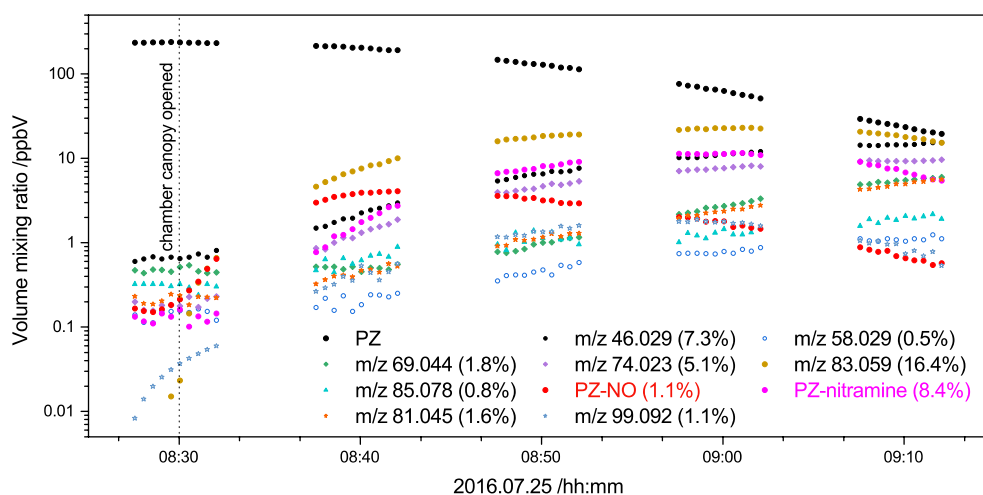


Figure 4. Time evolution of ion signals during the piperazine photo-oxidation experiment on 2016.07.25. Derived volume mixing ratios (ppbV) of piperazine and **xxxxxx** during the experiment. Product yields, calculated after the 30 mins of reaction, are shown in parentheses.

Other major ions in the spectrum include m/z 83.059, m/z 74.023 and m/z 46.029. We assigned the peak at m/z 83.059 to $C_4H_7N_2^+$ whose profile as well as that of m/z 81.045 highly correlates with the one of the imine (m/z 85.078). This is consistent with the observation in

aforementioned Nitrosopiperazine photolysis section. The peak at m/z 74.023 was assigned to *N*-formylformamide (CHONHCHO), a product from the reaction initiated by hydrogen-abstraction from one of the $-CH_2-$ groups in piperazine (Scheme 2). The yield of m/z 74.023 was estimated to be 5.1%. The signal at m/z 46.029 is identified as formamide (NH₂CHO). Piperazine is expected to condense with formaldehyde, forming C₅H₁₁N₂⁺ at m/z 99.092.

, whereas Table 2 summarises the major PTR-TOF-MS ion signals observed. The [M-H₂]⁺ fragment (4.1%) of piperazine upon protonation was taken into consideration

Table 2. Major PTR-TOF-MS ion signals observed during OH initiated piperazine photo-oxidation experiments.

m/z	Ion sum formula	Interpretation
43.0551	C ₃ H ₇ ⁺	Fragment of organic species
46.0301	CH ₄ NO ⁺	[NH ₂ CHO]H ⁺
59.0490	C ₃ H ₇ O ⁺	[CHONHCH ₂ OH]H ⁺ - NO ₂
69.0447	C ₃ H ₅ N ₂ ⁺	Imidazole ?
74.0243	C ₂ H ₄ NO ₂ ⁺	[CHONHCHO]H ⁺
81.045	C ₄ H ₅ N ₂ ⁺	[Pyrazine]H ⁺
83.0604	C ₄ H ₇ N ₂ ⁺	PZ, H ₂ elimination from [PZ-Im]H ⁺
85.0756	C ₄ H ₉ N ₂ ⁺	From piperazine ^a and PZ-Im, Fragment of [PZ-Im]H ⁺ , [PZ-NO]H ⁺ , [PZ-NO ₂]H ⁺
87.0900	C ₄ H ₁₁ N ₂ ⁺	PZ
99.0561	C ₄ H ₇ N ₂ O ⁺	?
99.0923	C ₅ H ₁₁ N ₂ ⁺	PZ + CH ₂ O condensation product
115.0862	C ₅ H ₁₁ N ₂ O ⁺	PZ-Im + CH ₂ O condensation product
116.0787	C ₄ H ₁₀ N ₃ O ⁺	PZ-NO
119.0855	C ₄ H ₁₁ N ₂ O ₂ ⁺	?
132.0750	C ₄ H ₁₀ N ₃ O ₂ ⁺	PZ-NO ₂

The predicted major products originated from one of the $-NH-$ groups, PZ-NO₂ (m/z 132.077), PZ-NO (m/z 116.084) and the imine, PZ-Im, (m/z 85.078) showed lower presence in spectrum. The former two fragments extensively upon protonation, and the co-existing photolysis caused further signal drop. Upon calibration, the PZ-NO and PZ-NO₂ yield was estimated to be 1.1% and 8.4% in this experiment. It should be note that the numbers present lower limits due to the underestimated wall losses.

The ion pair m/z 85.078 and 83.059 partially rationalized the low signal of protonated imine signal at m/z 85.078. Taking all factors into consideration, the yield of imine and its derivatives are approximately 17.2%.

One uncertainty factor in data analysis is the peak at m/z 43.055 (C₃H₇⁺), potentially a fragment of higher order of organic species. We decided to leave it out of yield estimation due

to 1), lab calibrations and literature suggest major products such as piperazine-imine, PZ-nitramine, *N*-formylformamide and formamide do not produce such a fragment; 2), the profile of $C_3H_7^+$ peak shows a high correlation with one of acetone, therefore it is suspected the origin of $C_3H_7^+$ peak to be chamber artefacts.

Other minor products are m/z 31.018, m/z 58.029 and m/z 69.044. We assigned this peak to formaldehyde. The m/z 69.044 is tentatively assigned to imidazole. The m/z 58.029 ($C_2H_4NO^+$) is believed to origin in $CHONHCH_2OH$ eliminating H_2O upon protonation. A detailed mass spectrum and list of the trace products and its possible interpretation is presented in [Figure S18](#) and [Table S8](#).

The present piperazine experiments are hampered by wall adsorption/desorption and particle formation, and the data are insufficient to allow and unambiguously determine the neutral molecules corresponding to the detected ion signals. However, none of the molecular formulae can be related to alkyl nitramines.

3.2.5 Piperazine + OH particle analysis

The aerosol formed in the PZ+OH photo-oxidation was simultaneously detected by three instruments: (1) a PTR-ToF 8000 instrument coupled with a Chemical Analysis of aeRosol ON-line (CHARON) inlet;³¹ (2) a compact time-of-flight Aerosol Mass Spectrometer (C-ToF-AMS); and (3) a Scanning Mobility Particle Sizer (SMPS). A detailed description of experimental and instrumental settings is available elsewhere¹¹. A relatively good agreement was achieved amongst the results obtained by these three instruments ([Figure S19](#)). The aerosol formation proceeded fast and heavily ($>10^5$ #/cm³ during all the experiments), as depicted in [Figure S20](#). The C-ToF-AMS measurements indicated that ammonium salt is the main constituent in particle phase. In addition to the PZ-nitrate salt, there exists a large fraction of organic species in the aerosol phase. This is in consistence with C-ToF-AMS observations.

To better understand the aerosol composition, we scrutinized the molecular information by CAHRON-PTR-ToF measurement of the piperazine photo-oxidation experiments. The experiments carried out on 26 July 2016, with the highest aerosol yield, was taken as an example. [Figure 4](#) illustrates the quantified mass spectrum of particle-phase products. The most abundant peaks are at m/z 87.092 (piperazine) and m/z 45.993 (NO_2^+), suggesting piperazine nitrate salt as the main component in aerosols, which accounts for 52.6% in total aerosol mass loading.

The other major peaks are m/z 72.042 ($C_3H_6NO^+$), m/z 99.078 ($C_5H_{11}N_2^+$, PZ-formaldehyde condensation product), m/z 101.071 ($C_4H_9N_2O^+$), m/z 115.082 ($C_5H_{11}N_2O^+$), m/z 117.067 ($C_4H_9N_2O_2^+$, $CHONHCH_2CH_2NHCHO$), and m/z 142.092 (tentatively assigned to $PZ \cdot (H_2O)_2H_3O^+$). In addition, nitramine were detected in particles, m/z 132.076 and m/z 177.059, assigned to PZ- NO_2 and di-PZ- NO_2 , respectively. It is worth to note that the PZ- NO_2 and di-PZ- NO_2 account for 1.7% and 0.88% in total aerosol mass, after fragmentation corrections.

The PZ portioning in aerosol varied from 0.2% (slow IPN, no NO_x) to 6% (quick IPN, 90 ppbv NO_2) in PZ+OH photooxidation experiment (see [Table S9](#)). Under the assumption that all organic carbon species in principle should originate from piperazine, this would further increase the portioning of PZ into aerosols to 15.5%. This is justified due to the expected heterogeneous reactions involving primary amines, imines, aldehydes, ketones and carboxylic acids happening on aerosol surfaces.

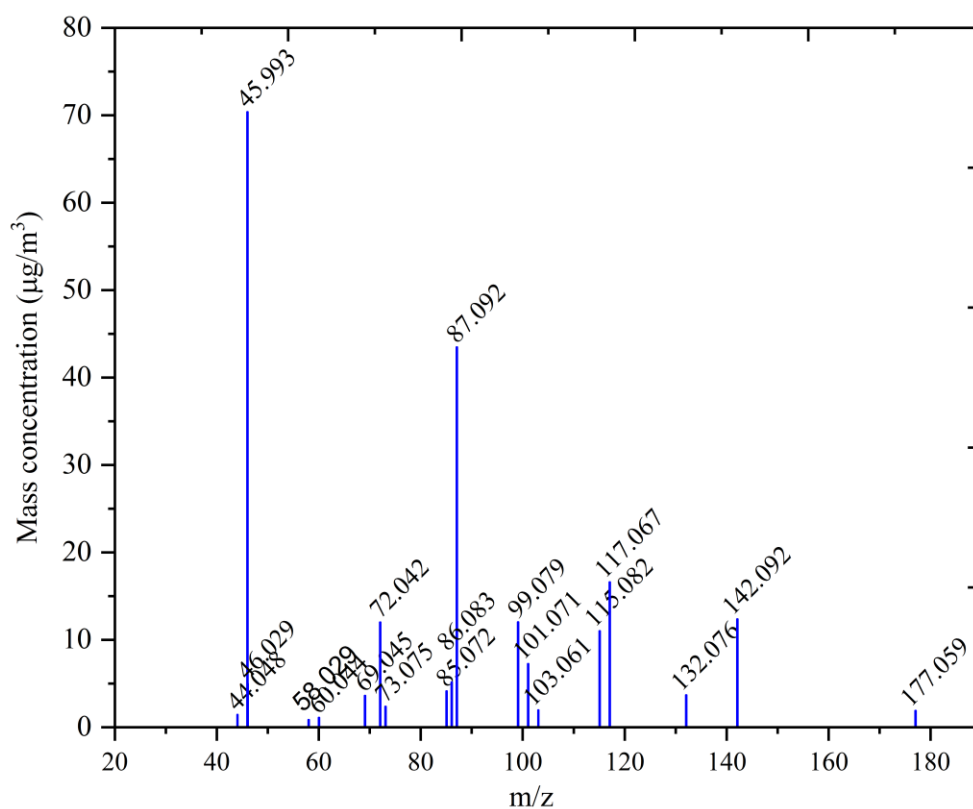


Figure 4. A typical spectrum of particle composition obtained at the highest aerosol loading of piperazine photooxidation under high NO_x initial condition, the mass concentration of each m/z is calculated by CHARON-PTR-ToF-MS

4. CONCLUSIONS

H-law

5. ATMOSPHERIC IMPLICATIONS

We should mention that there is no experimental value for the corresponding NO₃ rate coefficient, but the empirical correlation between OH and NO₃ rate coefficients for reaction with amines suggests a PZ + NO₃ rate coefficient around $4.6 \times 10^{-11} \text{ cm}^3 \text{ molecules}^{-1} \text{ s}^{-1}$ at 298 K.⁵

Other

■ Associated content

●, Supporting Information

The Supported Information is available free of charge on the [ACS Publications website](#) at DOI: [10.1021/acs/jPCA.xxxxxxxx](https://doi.org/10.1021/acs/jPCA.xxxxxxxx).

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Notes

The authors declare no competing financial interest

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