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# A combined experimental and DFT mechanistic study for the unexpected nitrosolysis of *N*-hydroxymethyldialkylamines in fuming nitric acid†

Yu Zhang, Po Zou, Yingbin Han, Yongliang Geng, Jun Luo \* and Baojing Zhou\*

The reaction of dimorpholinomethane in fuming HNO<sub>3</sub> was investigated. Interestingly, the major product was identified as *N*-nitrosomorpholine and a key intermediate *N*-hydroxymethylmorpholine was detected during the reaction by <sup>1</sup>H-NMR tracking which indicates that the reaction proceeds *via* an unexpected nitrosolysis process. A plausible nitrosolysis mechanism for *N*-hydroxymethyldialkylamine in fuming nitric acid involving a HNO<sub>3</sub> redox reaction is proposed, which is supported by both experimental results and density functional theory (DFT) calculations. The effects of ammonium nitrate and water on the nitrosolysis were studied using different ammonium salts as additives and varying water content, respectively. Observations show the key role of ammonium ions and a small amount of water in promoting the nitrosolysis reaction. Furthermore, DFT calculations reveal an essential point that ammonia, merged from the decomposition of the ammonium salts, acts as a Lewis base catalyst, and the hydroxymethyl group of the substrate participates in a hydrogen-bonding interaction with the NH<sub>3</sub> and H<sub>2</sub>O molecules.

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## Introduction

*N*-Nitramines and *N*-nitrosamines are present in a wide range of drug molecules, functional organic chemicals and energetic materials.<sup>1</sup> For example, some *N*-nitrosamines have biological activity and can be used in various treatments for illnesses including cancer, cardiovascular diseases, central nervous, and diseases related to immunity and physiological disorders.<sup>2</sup> *N*-Nitrosamines are valuable intermediates in organic synthesis<sup>3</sup> such as in application for the preparation of biologically important  $\alpha$ -disubstituted hydrazines<sup>4</sup> and mesoionic-heterocyclic compounds like sydnones.<sup>5</sup> *N*-Nitrosamines can also be applied as a nitroso source to prepare aryl *C*-nitroso compounds in good yield through a Fischer-Hepp rearrangement.<sup>6</sup> Recently, *N*-nitroso functionality has emerged as a traceless directing group for the activation of inert C–H bonds in aryl rings by associating with transition metals.<sup>7</sup> Furthermore, some widely used explosives with excellent performances such as 1,3,5-trinitro-1,3,5-triazacyclohexane (RDX), 1,3,5,7-tetranitro-1,3,5,7-tetraazacyclooctane (HMX) and 2,4,6,8,10,12-hexanitro-2,4,6,8,10,12-hexaazaisowurtzitane (CL-20) belong to the *N*-nitramine group, which can be prepared from acetamides and *N*-nitrosamines through nitrolysis or oxidation.<sup>8</sup>

Interestingly, when nitril chloride, nitrogen pentoxide, nitril fluoride, nitronium fluoroborate and tetranitromethane were used for the *N*-nitration of secondary or tertiary amines at low temperatures, nitrosamine products were obtained with low yields.<sup>9</sup> Our group made efforts to synthesize HMX by nitrolyzing 3,7-dinitro-1,3,5,7-tetraazabicyclo[3.3.1]nonane (DPT) with traditional nitration systems such as fuming HNO<sub>3</sub>, HNO<sub>3</sub>–H<sub>2</sub>SO<sub>4</sub>, HNO<sub>3</sub>–N<sub>2</sub>O<sub>5</sub>, and HNO<sub>3</sub>–Ac<sub>2</sub>O below 0 °C, but 1-nitroso-3,5,7-trinitro-1,3,5,7-tetraazacyclooctane (MNX) was achieved as the main product. Furthermore, NH<sub>4</sub>NO<sub>3</sub> can inevitably improve the yield of MNX.<sup>8f</sup> Recently, the so called “small-molecule route” using urea as the starting material to prepare HMX *via* DPT has been thought to be a promising method for scale-up.<sup>10</sup> Although the development of various techniques has enhanced the yield of HMX by the nitrolysis of DPT, no significant progress in practical potential has been achieved.<sup>11</sup> The main reason is attributed to the ambiguity of the nitrolysis mechanism.<sup>8e,8f</sup> It's widely considered that the bridging methylene of the DPT is cleaved to produce an extremely active intermediate 1-hydroxymethyl-3,5,7-trinitro-1,3,5,7-tetraazacyclooctane.<sup>12</sup> The addition of NH<sub>4</sub>NO<sub>3</sub> can hinder esterification (*O*-acetylation in HNO<sub>3</sub>–Ac<sub>2</sub>O and *N*-nitration in fuming HNO<sub>3</sub>) but promote dehydroxylation to improve the HMX yield.<sup>8e,8f,11,12</sup> However, there is no sufficient or direct experimental evidence to support the above mechanism because of the rapid reaction characteristics of the nitrolysis process, and the production of a complex by-product, and a short-lived intermediate.<sup>12</sup>

In this work, we selected a bridging methylene diamine dimorpholinomethane as a model substance and investigated

School of Chemical Engineering, Nanjing University of Science and Technology, Nanjing 210094, China. E-mail: [luojun@njust.edu.cn](mailto:luojun@njust.edu.cn); [bzhou@njust.edu.cn](mailto:bzhou@njust.edu.cn)

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in detail its reaction in fuming nitric acid to mimic the nitrolysis of DPT. Interestingly, nitrosomorpholine was obtained as the main product and the key intermediate *N*-hydroxymethylmorpholine was detected by  $^1\text{H-NMR}$  tracking during the reaction. Based on experimental observations, a plausible mechanism for the nitrosolysis of *N*-hydroxymethylalkylamines in fuming nitric acid is proposed. We further provide computational results for the possible reaction mechanism at the molecular level using the density functional theory (DFT), which is one of the popular routes to understand the course of an organic reaction.<sup>13</sup>

## Results and discussion



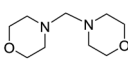
Similar to DPT, dimorpholinomethane has the *N,N'*-methylene motif. Therefore, we first examined the reaction of the dimorpholinomethane with fuming nitric acid. To our surprise, *N*-nitrosomorpholine was obtained rather than *N*-nitromorpholine when the reaction was carried out at room temperature (Table 1, entry 1). Prolonging the reaction time increased the yield of *N*-nitrosomorpholine (entries 2, 3). A small amount of *N*-nitromorpholine was isolated after reacting for 12 hours (entry 3). When the reaction temperature was elevated from 25 °C to 40 °C, both of them were obtained with higher yields (entry 4). A further increase of the reaction temperature to 70 °C afforded a lower yield of *N*-nitrosomorpholine but a higher yield of *N*-nitromorpholine (entry 5). It is well known that  $\text{NH}_4\text{NO}_3$  plays a key role in the nitrolysis of DPT.<sup>11</sup> Here, similarly, the yield of *N*-nitrosomorpholine increased dramatically when  $\text{NH}_4\text{NO}_3$  was used as an additive to the reaction mixture (entry 6).

It is generally accepted that 1-hydroxymethyl-3,5,7-trinitro-1,3,5,7-tetraazacyclooctane is a key intermediate during the nitrolysis of DPT in fuming nitric acid. Accordingly, we hypothesized that *N*-hydroxymethylmorpholine is the intermediate in the reaction of dimorpholinomethane in fuming nitric

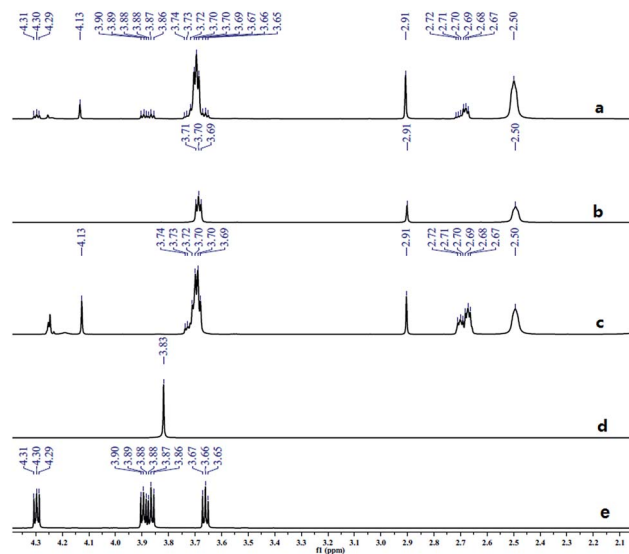
acid. To verify this, the reaction was performed at 0 °C for 10 minutes, then the reaction mixture was poured into ice water, extracted with cold dichloromethane, and washed with ice water to neutralise it. A yellow oily liquid was obtained by rotary evaporation at room temperature. According to the  $^1\text{H-NMR}$  spectrum, three compounds can be identified (Fig. 1a). As shown in Fig. 1b, the chemical shifts of the dimorpholinomethane occur at 3.70–3.69, 2.91, 2.50 ppm.<sup>14</sup> Clearly, the mixture mainly consists of dimorpholinomethane. As shown in Fig. 1c, hydrogen atoms observed at 4.13, 3.74–3.69 and 2.72–2.67 ppm are assigned to *N*-hydroxymethylmorpholine.<sup>14</sup> Comparing Fig. 1a with 1c, we can distinguish *N*-hydroxymethylmorpholine from the mixture. However, *N*-nitromorpholine, whose chemical shift is at 3.83 ppm (Fig. 1d), is not found in the mixture. Moreover, comparing Fig. 1a with 1e, we found that the chemical shifts at 4.31–4.29, 3.91–3.86 and 3.68–3.66 ppm in the former spectrum belong to *N*-nitrosomorpholine.<sup>14a</sup> Thus, *N*-hydroxymethylmorpholine is formed during the reaction and nitrosolysis is predominant at room temperature in classical nitration systems.

Although *N*-hydroxymethylmorpholine was observed in the reaction, it is not clear whether it is an intermediate or a by-product. To shed some light on this, we prepared a mixture of *N*-hydroxymethylmorpholine and dimorpholinomethane as the substrate for the reaction. As shown in Table 2, *N*-nitrosomorpholine is still the major product in fuming nitric acid, and only 3% of *N*-nitromorpholine is obtained (entry 1). The yields of *N*-nitrosomorpholine and *N*-nitromorpholine increase with the elongation of the reaction time (entries 1–3). In contrast, with an elevation in the reaction temperature, the yield of *N*-nitrosomorpholine first increases and then decreases (entries 1, 4, 5). Once again,  $\text{NH}_4\text{NO}_3$  significantly promotes nitrosolysis (entry 6). These observations are similar to those

**Table 1** The results for the reaction of dimorpholinomethane in fuming  $\text{HNO}_3$ <sup>a</sup>

Substrate	Entry	Reactant system	$T/^\circ\text{C}$	$t/\text{h}$	Yield <sup>c</sup> /%	
						
	1	$\text{HNO}_3$	25	1	10	0
	2	$\text{HNO}_3$	25	4	18	0
	3	$\text{HNO}_3$	25	12	25	5
	4	$\text{HNO}_3$	40	1	29	10
	5	$\text{HNO}_3$	70	1	15	16
	6	$\text{HNO}_3/\text{NH}_4\text{NO}_3$ <sup>b</sup>	25	1	31	0

<sup>a</sup> Unless otherwise noted, dimorpholinomethane (5 mmol), fuming  $\text{HNO}_3$  (200 mmol). <sup>b</sup>  $\text{NH}_4\text{NO}_3$  (12.5 mmol). <sup>c</sup> Isolated yield.



**Fig. 1**  $^1\text{H-NMR}$  spectrum of: (a) the product from the reaction of dimorpholinomethane in fuming  $\text{HNO}_3$  at 0 °C for 10 minutes; (b) dimorpholinomethane; (c) the mixture of *N*-hydroxymethylmorpholine and dimorpholinomethane; (d) *N*-nitromorpholine; (e) *N*-nitrosomorpholine.



Table 2 The results for the reaction of *N*-hydroxymethyldialkylamines in fuming HNO<sub>3</sub><sup>a</sup>

Substrate <sup>b</sup>	Entry	Reactant system	Nitrosamine product	Nitrosamine yield <sup>h</sup> /%	Nitramine product	Nitramine yield <sup>h</sup> /%
	1	HNO <sub>3</sub>		21		3
	2 <sup>c</sup>	HNO <sub>3</sub>		30		6
	3 <sup>d</sup>	HNO <sub>3</sub>		35		12
	4 <sup>e</sup>	HNO <sub>3</sub>		33		9
	5 <sup>f</sup>	HNO <sub>3</sub>		25		18
	6	HNO <sub>3</sub> /NH <sub>4</sub> NO <sub>3</sub> <sup>g</sup>		51		2
	7	HNO <sub>3</sub> /(NH <sub>4</sub> ) <sub>2</sub> SO <sub>4</sub> <sup>g</sup>		46		3
	8	HNO <sub>3</sub> /NH <sub>4</sub> Cl <sup>g</sup>		49		2
	9	HNO <sub>3</sub> /CH <sub>3</sub> COONH <sub>4</sub> <sup>g</sup>		47		2
	10	HNO <sub>3</sub>		21		0
	11	HNO <sub>3</sub> /NH <sub>4</sub> NO <sub>3</sub> <sup>g</sup>		39		0
	12	HNO <sub>3</sub>		19		4
	13	HNO <sub>3</sub> /NH <sub>4</sub> NO <sub>3</sub> <sup>g</sup>		38		4

<sup>a</sup> Unless otherwise noted, fuming HNO<sub>3</sub> (200 mmol),  $T = 25\text{ }^{\circ}\text{C}$ ,  $t = 1\text{ h}$ . <sup>b</sup> Substrate (5 mmol),  $n(N\text{-hydroxymethylmorpholine}) : n(\text{dimorpholinomethane}) = 1 : 1.1$ ,  $n(N\text{-hydroxymethylpiperidine}) : n(\text{dipiperidinomethane}) = 1 : 1.25$ ,  $n(N\text{-hydroxymethyldibutylamine}) : n(\text{dibutylamine}) = 1 : 1.17$ . <sup>c</sup>  $t = 4\text{ h}$ . <sup>d</sup>  $t = 12\text{ h}$ . <sup>e</sup>  $T = 40\text{ }^{\circ}\text{C}$ . <sup>f</sup>  $T = 70\text{ }^{\circ}\text{C}$ . <sup>g</sup> NH<sub>4</sub>NO<sub>3</sub> (12.5 mmol); NH<sub>4</sub>Cl (12.5 mmol); CH<sub>3</sub>COONH<sub>4</sub> (12.5 mmol); (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub> (6.25 mmol). <sup>h</sup> Isolated yield.

made in the nitrosolysis of dimorpholinomethane. Thus, *N*-hydroxymethylmorpholine should be an intermediate in the reaction, rather than a by-product. The nitrosolysis of the mixture of *N*-hydroxymethylmorpholine and dimorpholinomethane was further carried out in fuming nitric acid/ammonium salts containing (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>, NH<sub>4</sub>Cl, and CH<sub>3</sub>COONH<sub>4</sub>. Similar to NH<sub>4</sub>NO<sub>3</sub>, these ammonium salts can also improve the yield of *N*-nitrosomorpholine (entries 7–9).

We further examined the nitrosolysis of the mixture of *N*-hydroxymethylpiperidine with dipiperidinomethane and that of *N*-hydroxymethyldibutylamine with dibutylamine in the fuming HNO<sub>3</sub>, respectively. As shown in Table 2, the main products are identified to be *N*-nitrosamines<sup>15</sup> (entries 10, 12). To estimate the interference of dibutylamine, we calculated its amount in the mixture according to the proton NMR test. Then an equal amount of dibutylamine was used as the substrate for a reaction under common reaction conditions, which was monitored by gas chromatography. Only less than 4% *N*-nitrosodibutylamine (gas chromatography yield) was detected over 2 h. So we conclude that the *N*-nitrosodibutylamine is formed predominantly by the nitrosolysis of *N*-hydroxymethyldibutylamine. The achieved yield of *N*-nitrosopiperidine and *N*-nitrosodibutylamine in the fuming HNO<sub>3</sub>/NH<sub>4</sub>NO<sub>3</sub> system was 39% and 38% respectively, much higher than for the reactions without NH<sub>4</sub>NO<sub>3</sub> (entries 10–13).

The influence of water on the nitrosolysis of the mixture of *N*-hydroxymethylmorpholine and dimorpholinomethane was evaluated by performing the reaction in water-containing nitric acid, which was prepared by the addition of different amounts

of water into freshly prepared 100% HNO<sub>3</sub>. As shown in Fig. 2, the yield of *N*-nitrosomorpholine first increases and then decreases with the increase of water content. The highest yield of 21% was obtained when *ca.* 5% water was added. Meanwhile, a small amount of *N*-nitromorpholine was observed in all cases. These results suggest that *ca.* 5% water is optimal for the nitrosolysis of the mixture of *N*-hydroxymethylmorpholine and dimorpholinomethane.

Considering the fact that only very little HNO<sub>2</sub> and its analogues are present in fuming nitric acid, it is puzzling that the product is mostly *N*-nitrosamine in the above reactions. It is natural to surmise that a redox reaction has occurred. It is plausible that *N*-hydroxymethyldialkylamines react with NO<sup>+</sup> to

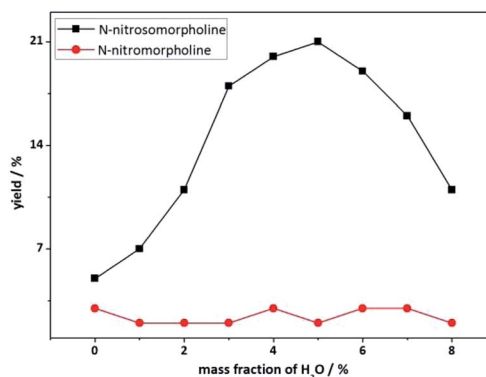


Fig. 2 The effect of water on the nitrosolysis of *N*-hydroxymethylmorpholine.



afford the *N*-nitrosoamines and release formaldehyde, which reduces nitric acid to nitrous acid. Then nitrous acid can be viewed as a catalyst. To further verify this viewpoint, three secondary amines were reacted in the fuming nitric acid with or without paraformaldehyde (Table 3). It turned out that *N*-nitrosamines were obtained in less than 6% yield in the absence of paraformaldehyde (entries 1, 3, 5). The addition of paraformaldehyde greatly increases the yield (entries 2, 4, 6). These results suggest that *N*-hydroxymethylalkylamines are formed as the more active intermediates under the acidic conditions when paraformaldehyde is present.<sup>13</sup>

In the above reaction, some *N*-nitromorpholine was observed and its amount increased with the reaction time (Table 2, entries 1–3). To investigate whether it was formed from the nitrolysis of *N*-nitrosomorpholine, we monitored the reaction of *N*-nitrosomorpholine in fuming nitric acid at 40 °C by <sup>1</sup>H-NMR spectroscopy. As shown in Fig. 3a, no *N*-nitromorpholine was observed at the end of the feeding process (Fig. 3a). However, *N*-nitromorpholine gradually formed with an increase in the reaction time as indicated by the chemical shift of 3.83 ppm in Fig. 3b and c<sup>15</sup> and its proportion increased with the disappearance of *N*-nitrosomorpholine. After 6 hours, almost all of the *N*-nitrosomorpholine had been converted to *N*-nitromorpholine (Fig. 3d). On the other hand, some *N*-nitroamines may have been generated by the oxidation of the *N*-nitrosamines.<sup>8</sup> To examine this possibility, the reactions of nitrosomorpholine in 90% HNO<sub>3</sub>, 80% HNO<sub>3</sub>, 70% HNO<sub>3</sub>, 60% HNO<sub>3</sub>, and 50% HNO<sub>3</sub> were investigated. Almost no product, however, was detected even when the reaction temperature changed from 20 °C to 80 °C. Thus, we conclude that *N*-nitromorpholine is mainly formed through the nitrolysis of *N*-nitrosomorpholine in fuming nitric acid.

We further performed a computational study to investigate the reaction mechanism at the atomic level. Two reaction models were constructed. The first reactant model contained one *N*-hydroxymethylalkylamine, one NH<sub>3</sub> and three H<sub>2</sub>O

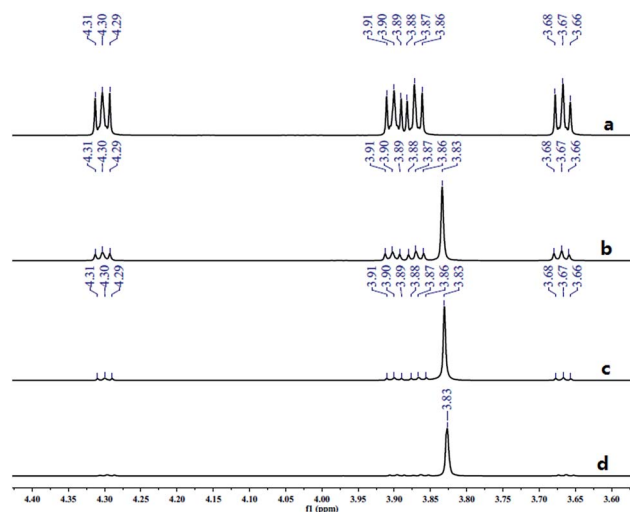


Fig. 3 <sup>1</sup>H-NMR spectra of samples taken from the reaction mixture at: (a) the end of the feeding process; (b) 2 hours; (c) 4 hours (d) 6 hours.







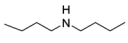
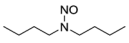
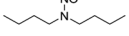
molecules. In another reaction model, the NH<sub>3</sub> molecule was replaced by a H<sub>2</sub>O molecule. Interestingly, we found that the free ammonia, formed from the unfavourable balance with NH<sub>4</sub><sup>+</sup> in an acidic environment, is a Lewis base catalyst, while water molecules are essential for the formation of the hydrogen bond (HB) adduct. A bicyclic ring was inferred from the bond angle of H<sub>2</sub>O molecules. The optimized structures of the reactants, transition states (TSs), and products from the two reaction models in the presence (model a) or absence (model b) of NH<sub>3</sub> are compared in Fig. 4.

For both models, we found that NO<sup>+</sup> is not involved in the bicyclic ring formed from the HB interactions and it approaches the N atom in the substrate from the other side. In the TS, the N–N bond forms, and the C–N bond is breaking, while the H is transferring from the hydroxyl group to the NH<sub>3</sub> (model a) or H<sub>2</sub>O molecule (model b), which suggests a synergistic mechanism. The computed relative Gibbs free energy profiles for models a and b are summarized in Fig. 5. For model a, the predicted Gibbs free energy of activation is 3.3 kcal mol<sup>−1</sup>. For model b, the reaction barrier is 13.7 kcal mol<sup>−1</sup>, which is 10.4 kcal mol<sup>−1</sup> higher than that of model a. These results indicate that the NH<sub>3</sub> molecule plays the role of a catalyst. Therefore, the addition of NH<sub>4</sub><sup>+</sup> can significantly promote the nitrolysis of *N*-hydroxymethylalkylamines.

The gradual changes of the four chemical bonds along the reaction coordinate are illustrated in Fig. 6. First, the hydroxyl H is extracted by the N of ammonia, which leaves the negative charge on the *N*-hydroxymethylalkylamine substrate. Then the electrophilic attack is initiated by NO<sup>+</sup> as indicated by its decreasing separation from the N atom of the substrate. When the N–N bond is half formed, the C–N and O–H bonds break, which leads to the formation of the products, *i.e.* nitrosamine, formaldehyde, and ammonium. The highly synchronous variations further indicate the concerted reaction mechanism for the nitrolysis of *N*-hydroxymethylmorpholine.

Based on our experimental and theoretical results and the related reports,<sup>8c,8f,11,12</sup> we propose a plausible mechanism for

Table 3 The nitrosation of secondary amines is promoted by paraformaldehyde<sup>a</sup>

Substrate	Entry	$n(\text{substrate}) : n(\text{CH}_2\text{O})$	Product	Yield <sup>b</sup> /%
	1	0		6
	2	1 : 1		33
	3	0		4
	4	1 : 1		51
	5	0		3
	6	1 : 1		65

<sup>a</sup> Substrate (5 mmol), fuming HNO<sub>3</sub> (200 mmol)  $T = 25$  °C,  $t = 1$  h.

<sup>b</sup> Isolated yield.



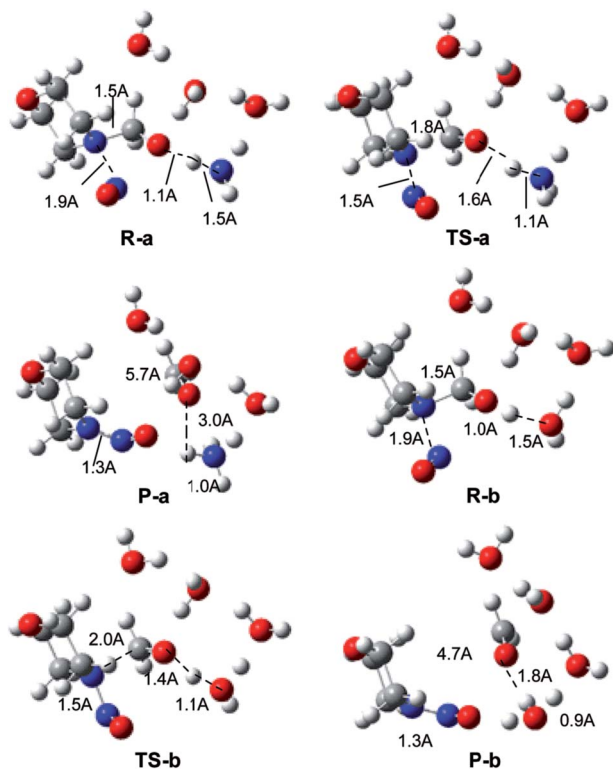


Fig. 4 Optimized structures for the two reaction models representing the nitrosolysis of *N*-hydroxymethylmorpholine. Model a contains an  $\text{NH}_3$  molecule, while in model b, the  $\text{NH}_3$  molecule is replaced by a  $\text{H}_2\text{O}$  molecule. O: red, C: dark grey, H: light grey, N: blue.

the formation of *N*-nitrodialkylamines from *N*-hydroxymethylalkylamines in fuming nitric acid and an ammonium nitrate system (Scheme 1). First, a small amount of  $\text{HNO}_2$  in fuming  $\text{HNO}_3$  is converted to  $\text{NO}^+$  and some free  $\text{NH}_3$  emerges through the equilibrium reaction with  $\text{NH}_4\text{NO}_3$ . The substrate is conjoined to one ammonia and three water molecules by HB interactions to form the pre-reactive complex *in situ*. Then, the  $\text{NH}_3$  abstracts the hydrogen from the hydroxyl group to increase the electron density of the tertiary N atom, which facilitates the

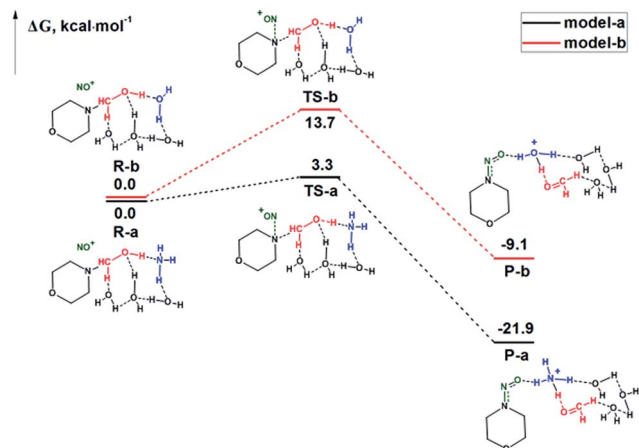


Fig. 5 Energy profile for the nitrosolysis of *N*-hydroxymethylmorpholine in the presence or absence of  $\text{NH}_3$ .

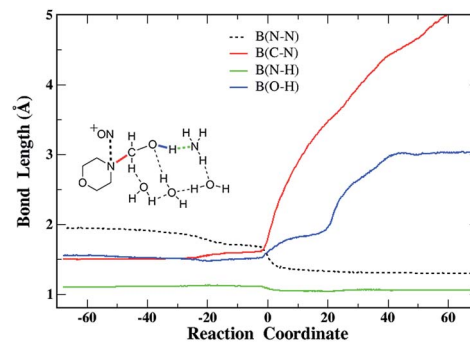
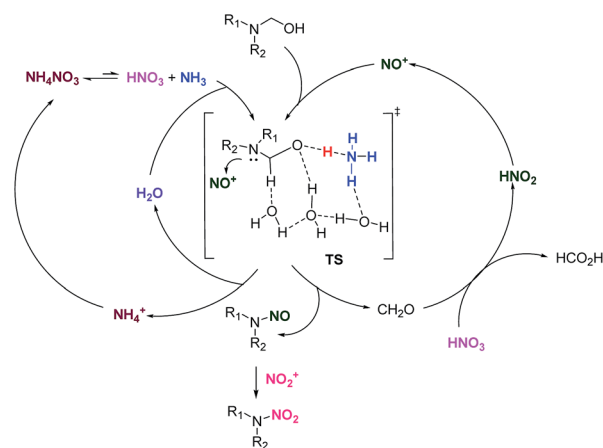


Fig. 6 Bond cleavages and formations along the IRC path of the nitrosolysis of *N*-hydroxymethylmorpholine.



Scheme 1 Possible reaction mechanism for the nitrosolysis of *N*-hydroxymethylalkylamine.

attack of  $\text{NO}^+$ . Then *N*-nitrosamine forms along with the departure of formaldehyde. Subsequently, formaldehyde is oxidized by  $\text{HNO}_3$  to form  $\text{HCOOH}$ .<sup>16</sup> Simultaneously,  $\text{HNO}_3$  is reduced to  $\text{HNO}_2$  to complete one nitrosolysis cycle. *N*-Nitrosamine is afforded from *N*-nitrosamine by the replacement of the nitroso group with a nitro group *via* a nitrolysis process.

## Conclusions

We synthesized *N*-nitrosodialkylamines from *N*-hydroxymethylalkylamines in fuming  $\text{HNO}_3$  *via* nitrosolysis. The reaction mechanism at the molecular level was established based on experimental observations and DFT calculations and involves a redox reaction of  $\text{HNO}_3$  to afford  $\text{HNO}_2$ . The DFT calculations indicate that the nitrosolysis reaction proceeds very smoothly with a rigid bicyclic transition state consisting of *N*-hydroxymethylalkylamine, ammonia and water.

## Experimental section

### General procedure for the nitrosolysis of the mixture of *N*-hydroxymethylalkylamines in fuming nitric acid

The mixture of *N*-hydroxymethylalkylamines (5 mmol) was added in portions to fuming nitric acid (200 mmol) under



vigorous stirring at 0 °C, and the reaction mixture was then warmed to a certain temperature. The reaction was quenched by being poured into ice water (20 mL). The resulting mixture was extracted with dichloromethane (3 × 30 mL). The combined organic layer was washed with saturated NaHCO<sub>3</sub> and dried with Na<sub>2</sub>SO<sub>4</sub>. After filtration, the mixture was concentrated under reduced pressure to afford the crude product, and the desired product was obtained by column chromatography.

The same procedure as described above was used for the nitrosolysis of dimorpholinomethane.

## Conflicts of interest

There are no conflicts to declare.

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