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Probing the Reaction Mechanisms Involved in the Decomposition of Solid 1,3,5-Trinitro-1,3,5-Triazinane (RDX) by Energetic Electrons

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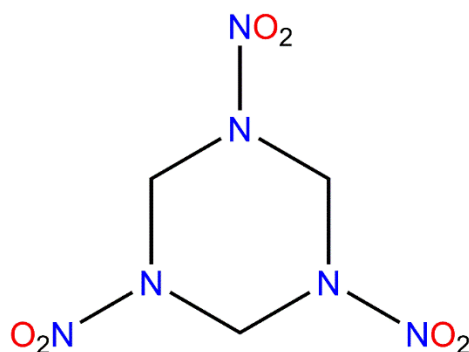
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Abstract

The decomposition mechanisms of RDX have been explored over the past decades, but as of now, a complete picture on these pathways has not yet emerged as evident from the discrepancies in proposed reaction mechanisms and the critical lack of products and intermediates observed experimentally. This study exploited a surface science machine to investigate the decomposition of solid-phase RDX by energetic electrons at a temperature of 5 K. The products formed during the irradiation were monitored online and *in situ* via infrared and UV-VIS spectroscopy, and products subliming in the temperature programmed desorption phase were probed with a reflectron time-of-flight mass spectrometer coupled with a soft photoionization at 10.49 eV (ReTOF-MS-PI). The infrared spectroscopy revealed the formation of water (H₂O), carbon dioxide (CO₂), dinitrogen oxide (N₂O), nitrogen monoxide (NO), formaldehyde (H₂CO), nitrous acid (HONO) and nitrogen dioxide (NO₂). ReTOF-MS-PI identified 39 cyclic and acyclic products arranged into, e.g., dinitro, mononitro, mononitroso, nitro-nitroso, and amines species. Among these molecules, 21 products such as N-methylnitrous amide (CH₄N₂O), 1,3,5-triazinane (C₃H₉N₃) and N-(aminomethyl)methanediamine (C₂H₉N₃) were detected for the first time in laboratory experiments; mechanism based on gas phase and condensed phase calculations were exploited to rationalize the formation of the observed products. The present studies reveal a rich, unprecedented chemistry in the condensed phase decomposition of RDX, which is significantly more complex than the unimolecular gas phase decomposition of RDX thus leading us closer to an understanding of the decomposition chemistry of nitramine-based explosives.

1. Introduction

Nitramine-based (N-NO₂) energetic materials such as 1,3,5-trinitro-1,3,5-triazinane (RDX) (Scheme 1) have been widely exploited as explosives and propellants.¹⁻³ Therefore, over the past decades, extensive studies have been performed to understand the fundamental decomposition mechanisms and the combustion behavior of RDX.¹⁻⁴¹ These investigations are very much required by the energetic material community to unravel the mechanism along with initial and higher order bond breaking processes, which trigger the decomposition of energetic molecules such as nitramines. Our current understanding of the reaction mechanisms involved in the decomposition of RDX is based on extensive experimental as well as computational studies performed in the gas-phase and in the condensed-phase (Table 1).⁴⁻³⁷ Gas-phase studies predicted the N-NO₂ bond fission as the initial step in the unimolecular decomposition of RDX.^{6-8, 10} For instance, the photodissociation of supersonic jet-cooled RDX at 226 nm revealed the formation of nitrogen monoxide (NO) as the decomposition product via rapid predissociation of nitrogen dioxide (NO₂) with the latter being released by the initial N-NO₂ bond rupture.^{6, 10} However, infrared multiphoton dissociation (IRMPD) studies of RDX in a jet-cooled molecular beam proposed a different mechanism.⁹ This study elucidated a concerted, symmetric triple dissociation of the RDX-ring producing three molecules of methylene nitramine (CH₂NNO₂) as the dominant decomposition pathway in contrast to the N-NO₂ bond fission.⁹ On the other hand, IRMPD experiments of RDX performed by Zuckermann et al. in the gas-phase revealed yet a third decomposition route involving a five-membered ring intermediate, which eventually formed hydroxyl (OH) radicals either directly or via dissociation of nitrous acid (HONO).¹¹



Scheme 1. Molecular structure of 1,3,5-trinitro-1,3,5-triazinane (RDX)

Table 1. Molecular formula, mass, and ionization energies of decomposition products of RDX previously reported in gas-phase and condensed-phase studies.

Method	Products observed				References
	Formula	Name	Mass (amu)	IE (eV)	
Photodissociation of RDX at 225 nm in supersonic jet	NO	Nitrogen monoxide	30	9.26	Im, H.-S. <i>et al.</i> ⁸
Photodissociation at 226 nm in supersonic jet	NO	Nitrogen monoxide	30	9.26	Guo, Y. Q. <i>et al.</i> ⁷
Photodissociation at 226 nm in supersonic jet	NO	Nitrogen monoxide	30	9.26	Lemire, G. W. <i>et al.</i> ¹⁰
	NO ₂	Nitrogen dioxide	46	8.80	
IR multiphoton dissociation in molecular beam	HCN	Hydrogen cyanide	27	13.59	Zhao, X. <i>et al.</i> ⁹
	CO	Carbon monoxide	28	14.01	
	NO	Nitrogen monoxide	30	9.26	
	H ₂ CO	Formaldehyde	30	10.88	
	CH ₂ NN	Diazomethane	42	8.99	
	N ₂ O	Nitrous oxide	44	12.77	
	CO ₂	Carbon dioxide	44	13.77	
	NO ₂	Nitrogen dioxide	46	8.80	
	HONO	Nitrous acid	47	-	
	C ₂ H ₄ N ₂	-	56	-	
	C ₃ H ₃ N ₃	-	74	-	
	CH ₂ N ₂ O ₂	s-triazene	81	10.00	
	C ₃ H ₄ N ₃	-	82	-	
	C ₃ H ₅ N ₃	-	83	-	
	C ₂ H ₄ N ₃ O ₂	-	102	-	
	CH ₂ N ₃ O ₄	-	120	-	
	C ₂ H ₄ N ₄ O ₃	-	132	-	
	C ₂ H ₃ N ₄ O ₄	-	148	-	
Shockwave Decomposition	NO ₂	Nitrogen dioxide	46	8.80	Miao, M. <i>et al.</i> ²⁵
	HONO	Nitrous acid	47		

Thermal	H ₂ O	Water	18	12.80	Behrens, R. <i>et al.</i> ⁴
Decomposition	NO	Nitrogen monoxide	30	9.26	
	CH ₂ O	Formaldehyde	30	10.88	
	N ₂ O	Nitrous oxide	44	12.77	
	NO ₂	Nitrogen dioxide	46	8.80	
	(CH ₃)NHCHO	N-Methylformamide	59	9.50	
	C ₃ H ₃ N ₃ OH	Hydroxytriazine	98	-	
Thermal	H ₂ O	Water	18	12.80	Maharrey, S. <i>et al.</i> ²⁴
decomposition	HCN	Hydrogen cyanide	27	13.59	
	CO	Carbon monoxide	28	14.01	
	CH ₂ O	Formaldehyde	30	10.88	
	NO	Nitrogen monoxide	30	9.26	
	N ₂ O	Nitrous oxide	44	12.77	
	H ₂ NCHO	N-Formamide	45	10.20	
	NO ₂	Nitrogen dioxide	46	8.80	
	HONO	Nitrous acid	47	-	
	(CH ₃) ₃ N	Trimethyl amine	58	7.80	
	(CH ₃)NHCHO	N-methyl formamide	59	9.05	
	C ₂ H ₂ N ₂ O	-	70	-	
	(CH ₃) ₂ NCHO	N,N-dimethyl formamide	73	9.55	
	(CH ₃) ₂ NNO	N-nitroso dimethylamine,	74	8.69	
	C ₃ H ₃ N ₃	s-triazene	81	10.00	
	C ₃ H ₃ N ₃ O	oxo-s-triazine	97		
	C ₃ H ₄ N ₄ O ₂	-	128	-	
	C ₃ H ₆ N ₆ O ₅	ONDNTA	206	-	
CO ₂ Laser Pyrolysis	HCN	Hydrogen cyanide	27	13.59	Botcher, T. R. <i>et al.</i> ²⁷
	NO	Nitrogen monoxide	30	8.80	
	N ₂ O	Nitrous oxide	44	12.77	
	CO ₂	Carbon dioxide	44	13.77	
	N ₂ O ₄	Nitrogen dioxide dimer	92	-	
CO ₂ laser pyrolysis and combustion	H ₂	Hydrogen molecule	2	15.44	Lee, Y. <i>et al.</i> ²⁸
	H ₂ O	Water	18	12.80	

	HCN	Hydrogen cyanide	27	13.59	
	CO	Carbon monoxide	28	14.01	
	H ₂ CNH	Methyleneimine	29	9.90	
	NO	Nitrogen monoxide	30	9.26	
	H ₂ CO	Formaldehyde	30	10.88	
	HNCO	Isocyanic acid	43	-	
	N ₂ O	Nitrous oxide	44	12.77	
	NO ₂	Nitrogen dioxide	46	8.80	
	C ₂ H ₄ N ₂	Diazaethane	46	-	
	HONO	Nitrous acid,	47	-	
	C ₂ H ₂ N ₂	Iminoacetonitrile	54	11.60	
	C ₂ H ₂ N ₂ O	-	70	-	
	C ₃ H ₃ N ₃	s-triazene	81	10.00	
	C ₃ H ₃ N ₃ O	oxo-triazine	97	-	
Thermal decomposition:	HNCO	Isocyanic acid	43	11.60	Gongwer, P. E. <i>et al.</i> ²⁹
	HONO	Nitrous acid	47	-	
Flash –heating	C ₃ H ₇ NO ₂	C-hydroxy-N-methylformamide,	58	-	
	C ₃ H ₃ N ₃	s-triazene	81	10.0	
Photodissociation at 248 nm	OH	Hydroxyl radical	17	-	Capellos, C. <i>et al.</i> ³⁰
	NO	Nitrogen monoxide	30	9.26	
	NO ₂	Nitrogen dioxide	46	8.80	
Photodissociation at 248 nm	H ₂	Hydrogen	2	15.44	Dickinson, J. T. <i>et al.</i> ³⁴
	OH	Hydroxyl radical	17	13.6	
	H ₂ O	Water	18	12.80	
	HCN	Hydrogen cyanide	27	13.59	
	H ₂ CO	Formaldehyde	30	10.88	
	NO	Nitrogen monoxide	30	9.26	
	CH ₂ NN	Diazomethane	42	9.00	
	C ₂ H ₄ N	Ethanimitrilium	42	-	
	N ₂ O	Nitrous oxide	44	12.77	
	C ₂ H ₄ N	Ethanimitrilium	45	-	
	NO ₂	Nitrogen dioxide	46	8.80	
	CH ₂ CH ₂ N ₂	Diazaoethane	46	-	
	CH ₂ NNO ₂	Methylene nitramine	74	-	
	C ₃ H ₃ N ₃	s-triazene	81	10.0	

Photolysis at 266 nm	O	Nascent Oxygen	16		Tang, T. B. <i>et al.</i> ³¹
	OH	Hydroxyl radical	17	13.6	
	CN	Cyanide radical	26		
	HCN	Hydrogen cyanide	27	13.59	
	NCN	Cyanonitrene	40	-	
	CHNN	Diazomethane radical	41	-	
	CH ₂ NN	Diazomethane	42	9.00	
	NO ₂	Nitrogen dioxide	46	8.80	
Photolysis at 229 nm	NO ₂	Nitrogen dioxide	46	8.80	Gares, K. L. <i>et al.</i> ³³
Thermal decomposition at high temperature and pressure	CO ₂	Carbondioxide	44	-	Connor, L. E. <i>et al.</i> ³⁸
	N ₂ O	Nitrous oxide	44	12.77	
Radiation-induced decomposition in solution	NO ₂	Nitrogen dioxide	46	8.80	Markarov, I. E. <i>et al.</i> ²⁶
	H ₂ O ₂	Hydrogen peroxide	34	12.10	
Catalytic thermal decomposition	CO	Carbon monoxide	28	14.01	Song, N. –M. <i>et al.</i> ³⁹
	N ₂	Nitrogen	28		
	N ₂ O	Nitrous oxide	44	12.77	
	NO ₂	Nitrogen dioxide	46	8.80	
Photodissociation at 236 nm	NO fragments	Nitrogen monoxide	30	9.26	Wynn, C. M. <i>et al.</i> ⁴⁰

Several theoretical calculations have been performed to resolve the aforementioned discrepancies for the gas-phase decomposition of RDX.^{12-16, 18-21} The majority of these calculations favored the homolytic N-NO₂ bond fission over the concerted triple C-N scission mechanism.¹³⁻¹⁸ The energy required for the N-NO₂ homolysis is lower by about 84 kJ mol⁻¹ relative to the concerted mechanism.¹⁴ In addition to the most favorable N-NO₂ bond fission, Goddard et al. proposed a new decomposition pathway, which involved three successive molecular eliminations of nitrous acid (HONO) forming eventually the 1,3,5-triazine molecule (C₃H₃N₃).¹⁴ Quantum calculations performed by Swadley et al. further confirmed the N-NO₂ bond rupture and nitrous acid (HONO) elimination reactions as initial steps in the decomposition of RDX.¹⁵ Lammertsma and co-workers proposed that the N-NO₂ homolysis could be followed by hydrogen migration during

the decomposition of RDX.¹⁸ In contrast to these three pathways (N-NO₂ bond fission, HONO elimination, concerted triple C-N scission), Bernstein et al. extracted a fourth pathway where isomerization of one of the nitro (-NO₂) to a nitroso (-ONO) group was considered as an initial route for the unimolecular decomposition of RDX.⁸ Bartlett et al. revealed that the nitrous acid (HONO) elimination reaction represents the most preferred route at 298 K and 1 atm pressure compared to N-NO₂ homolysis, concerted C-N scission, and nitroso (-ONO) isomerization.¹³ However, *ab initio* molecular dynamics calculation by Schweigert proposed the dominance of N-NO₂ bond rupture over nitrous acid (HONO) elimination and triple C-N scission mechanism over a temperature range from 1,000 to 2,000 K.¹⁶

Besides the gas phase, the decomposition of RDX in the solid as well as liquid phase was explored exploiting thermal energy, shock waves, and laser irradiation.^{4, 24-34} For example, thermal decomposition of RDX was investigated by Behrens and co-workers using simultaneous thermogravimetric modulated beam mass spectrometry at a temperature above and below the melting point of RDX (478 K).⁴⁻⁵ These authors observed several products such as water (H₂O), hydrogen cyanide (HCN), carbon monoxide (CO), formaldehyde (H₂CO), nitrogen monoxide (NO), dinitrogen oxide (N₂O), formamide (NH₂CHO), nitrogen dioxide (NO₂), nitrous acid (HONO), N-methylformamide ((CH₃)NHCHO), oxy-s-triazine (C₃H₃N₃O; OST), and 1-nitroso-3,5-dinitrohexahydro-s-triazine (ONDNTA; C₃H₆N₆O₅).^{4, 24} The authors proposed that ONDNTA as the only product formed during the early stage of the decomposition; this molecule subsequently decomposed to dinitrogen monoxide (N₂O) and formaldehyde (H₂CO).⁴ However, the authors did not observe any primary intermediates in the experiments. Further, shock wave decomposition of RDX crystals at pressures lower than 10 GPa revealed that the decomposition pathways leading to the formation of nitrogen dioxide (NO₂) and nitrous acid (HONO) were dominant. However, stresses higher than 10 GPa resulted in dominance of successive between the pristine radicals generated and unreacted RDX.²⁵ Wight and Botcher studied the laser pyrolysis of RDX at 77 K.²⁷ Their observation of dinitrogen tetroxide (N₂O₄) during the initial stage of the decomposition proposed that the N-NO₂ bond fission was likely the primary pathway. Laser pyrolysis of RDX was also studied at 293 K; the authors observed hitherto higher molecular weight species with mass-to-charge (m/z) values of 47 (HONO), 54 (C₂H₂N₂), 56 (C₂H₂N₂), 70 (C₂H₂N₂O), 81 (C₃H₃N₃) and 97 (C₃H₃N₃O).²⁸ UV-photolysis of RDX crystals were explored at distinct wavelengths.^{30-31, 33-34} Photolysis at 248, 236 nm and 229 nm

predominantly produced nitrogen dioxide (NO_2), which further supports the N- NO_2 bond fission to be the primary route for the decomposition of RDX.^{30, 34} Exposure to 266 nm resulted in successive NO_2 losses, ring opening forming nitrogen-bearing species along with highly unsaturated molecules carrying C=N moieties.³¹ Molecular dynamics simulations revealed the formation of nitrogen dioxide (NO_2) via N- NO_2 bond rupture, nitrous acid (HONO) elimination, and fragmentation to methylene nitramine (CH_2NNO_2).³⁵⁻³⁷

In summary, condensed phase studies exposed a significantly richer and more complex chemistry as evidenced in higher molecular weight products, which were not observed in the gas phase conditions. It is important to mention that the majority of the thermal decomposition studies of RDX were carried out in a temperature range from 443 K to 573 K.^{4, 24} At these temperatures, the reactive intermediates formed during the decomposition may undergo rapid decomposition to secondary products; likewise, the initially formed carbon-, oxygen-, and nitrogen-centered radicals can react further.²⁷ Therefore, a complete picture and a consistent decomposition mechanism of condensed phase RDX has not yet emerged. Here, we employed a surface science machine to explore the decomposition of RDX in the condensed phase (solid) via energetic electrons. The sample is kept at 5 K during the exposure thus eliminating the escape of any volatile products into the gas phase. The products are identified via Fourier transform infrared (FTIR) and ultraviolet-visible (UVVIS) spectroscopy (condensed phase) and by a reflectron time-of-flight mass spectrometer coupled with vacuum ultraviolet photoionization (PI-ReTOF-MS) during the temperature programmed desorption (TPD). This technique has been used previously to study the decomposition mechanism of nitromethane (CH_3NO_2), which is considered as the simplest model compound of nitrohydrocarbon-based energetic materials.⁴²⁻⁴⁵ Our study on the decomposition of RDX revealed 21 previously unobserved intermediates. Most significantly, the observation of mononitroso intermediates propose the existence of ONDNATA decomposition products. Our results also show evidence of reactive radical species previously only predicted computationally, but not observed in experiments as reactive intermediates. Based on our observations, we propose comprehensive decomposition pathways of RDX in the condensed phase.

2. Experimental Methods

2.1. Experimental

The experiments were carried out in an ultrahigh vacuum (UHV) chamber evacuated to a pressure of typically 2×10^{-10} torr exploiting oil-free turbomolecular pumps backed by dry scroll pumps.⁴⁶⁻⁴⁸ A polished silver wafer (substrate) coated with a film of RDX at a thickness of $16.8 \pm 1.0 \mu\text{m}$ is sandwiched with indium foil and interfaced to a cold finger manufactured from oxygen-free high conductivity copper (OFHC). This assembly is connected to a UHV compatible closed cycle helium refrigerator (Sumitomo Heavy Industries, RDK-415E) which cools down the temperature of the substrate to 5.0 ± 0.1 K. The temperature of the substrate can be controlled with the help of a cartridge heater. This system is freely rotatable horizontally and translatable vertically exploiting a differently pumped rotary feedthrough (Thermoionics Vacuum Products, RNN-600/FA/MCO). Infrared (IR) spectra of the RDX film were collected *in situ* at 5 K in the $4000\text{-}500 \text{ cm}^{-1}$ region using a Fourier Transform Infrared spectrometer (Nicolet 6700) operated at a resolution of 4 cm^{-1} . All FTIR measurements were performed in an absorption-reflection-absorption geometry at an angle of 45° to the normal of the substrate (Figs. 1 - 2, Table 2). The thickness of RDX film was determined based on interference patterns observed in the infrared spectrum due to fringing effect via equation (1)

$$d = \frac{N}{2\sqrt{n^2 - \sin^2\theta} (\nu_1 - \nu_2)} \quad (1)$$

where d is the thickness of the film, N is the number of fringes in the spectral region from ν_1 to $\nu_2 \text{ cm}^{-1}$, n is the refractive index of RDX ($n = 1.49$)⁴⁹, θ is the angle of incidence, and ν_1 and ν_2 are the start and end points of the spectrum in cm^{-1} covering the fringes. The UV-VIS spectra of the RDX films were also recorded *in situ* at 5 K covering the range from 190-800 nm using a modified UV-VIS spectrophotometer (Evolution 600). All UV-VIS measurements were performed at an angle of 30° relative to the normal of the substrate. After the reference spectra of the unirradiated RDX samples were taken, each RDX film was exposed to energetic electrons (5 keV) at an angle of 0° relative to the normal of sample surface, at two different electron currents of $20 \pm 2 \text{ nA}$ for 30 min (low dose) and $150 \pm 2 \text{ nA}$ for 60 min (high dose) by scanning the electron beam over an area of $1.0 \pm 0.1 \text{ cm}^2$. Infrared spectra of the RDX films were collected *in situ* before and after the irradiation to monitor the changes induced by the ionizing radiation inside the films. Monte Carlo simulations via the CASINO 2.42 software⁵⁰ were performed to

determine electron penetration depth and the average energy deposited on the RDX-film (Table 3). The average penetration depth of the electrons is calculated to be 267 ± 80 nm with averages doses of 8.1 ± 1.0 and 100 ± 16 eV per molecule for the low dose and high dose experiments, respectively. The penetration depth of the electron is less than the thickness of the RDX film guaranteeing that electron beam does not interact with the silver substrate.

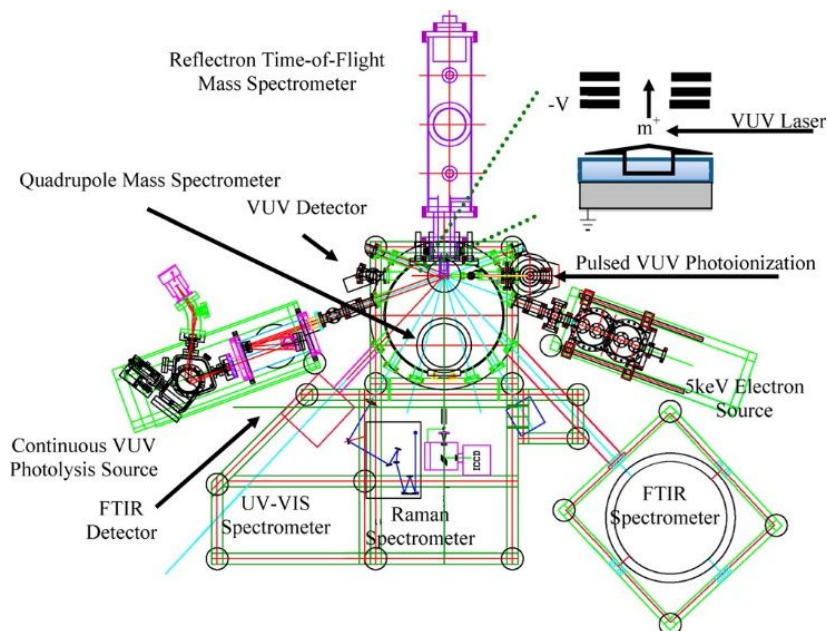


Figure 1. Schematic top view of the ultra-high vacuum chamber including the radiation sources (electron source), analytical instruments (FTIR, UV-VIS, ReTOF), and cryogenic target (point of convergence lines).⁴⁶⁻⁴⁸

After the irradiation, the exposed samples were annealed from 5 K to 320 K at a rate of 1 K min^{-1} (temperature programmed desorption; TPD). During the TPD, the products subliming from the substrate were analyzed exploiting photoionization reflectron time-of-flight mass spectrometer (PI-ReTOF-MS). Details of the PI-ReTOF-MS setup have been described previously.^{47-48, 51-53} In brief, we exploited pulsed vacuum ultraviolet (VUV) light at 10.49 eV to softly photoionize the subliming molecules. These ions are extracted and eventually detected in the reflectron time-of-flight tube based on their mass-to-charge (m/z) ratios prior to reaching the microchannel plate (MCP) detector (Jordan TOF Products Inc.). The signal generated by MCP detector is amplified using a pre-amplifier (Ortec 9305) and shaped with a 100 MHz discriminator (Advanced Research Instruments Corporation; F-100TD). A computer based multichannel scaler (FAST ComTec, P7888-1 E) receives the signal from the discriminator and

records it in 4 ns bins triggered at 30 Hz by a pulse delay generator (Quantum Composers 9518). 3600 sweeps are collected per mass spectrum per 1 K increase during the TPD phase.

Table 3. Data applied to calculate the average dose per RDX molecule for the low and high dose experiments.

	(a)	(b)
Initial Kinetic energy of the electrons	5keV	5keV
Irradiation current (I)	20±2 nA	150±2 nA
Irradiation time (t)	1800 s	3600 s
Average penetration depth, l	267±80 nm	267±80 nm
Average kinetic energy of backscattered electrons, E_{bs}^a	3.6±0.3 keV	3.6±0.3 keV
Fraction of backscattered electrons, f_{bs}^a	0.6±0.1	0.6±0.1
Average kinetic energy of transmitted electrons, E_{trans}^a	0.0 keV	0.0 keV
Fraction of transmitted electrons, f_{trans}^a	0	0
Density of the ice, ρ	1.82 gcm ⁻³	1.82 gcm ⁻³
Irradiated area, A	1.0±0.1 cm ²	1.0±0.1 cm ²
total number of molecules processed	(1.1±0.2)×10 ¹⁸	(1.6±0.2)×10 ¹⁹
dose per molecule, D	8.1 ± 1.0 eV	100 ± 16 eV
Total number of electrons	(2.3±1.0)×10 ¹⁴	(3.4±1.0)×10 ¹⁵

^a Values from CASINO simulations.

2.2. Sample Preparation

RDX samples were received from BAE Systems, Inc. prepared according to the MIL-DTL-398 D (Military Specification, Detail Specification RDX, 12 DEC 1996); this specifies an HMX (1,3,5,7-tetranitro-1,3,5,7-tetrazoctane; C₄H₈N₈O₈) impurity in RDX at levels from 4 % to 17 %. The maximum permissible quantity of other impurities if present is 0.08 %. In order to remove the impurities, production grade RDX was recrystallized from acetone (Fisher Scientific, Inc.). Nuclear magnetic resonance (NMR) spectroscopy was exploited to characterize the recrystallized RDX. Figure S1 of the supporting information shows the ¹³C NMR spectra of the crude and recrystallized RDX measured using a Spinsolve 60 Carbon benchtop 60 MHz NMR spectrometer after dissolving in dimethylsulfoxide-d₆ (DMSO-d₆) solvent. The disappearance of the peaks associated with the chemical shift of HMX indicates that RDX has been sufficiently purified to a

level of at least 99.9 %.⁵⁴ A thin film of RDX onto the silver substrate was prepared using a drop-casting method. For the drop-casting, about 2 mg of recrystallized (purified) RDX is dissolved in 3 g of methyl ethyl ketone (MEK; Acros Organics) inside a vial. Sonication and mild heating at 305 K ensure complete dissolution of the RDX in MEK. Using a glass pipette, a drop of RDX/MEK solution was deposited onto the silver substrate and then left for drying. A uniform layer of RDX is observed after the evaporation of solvent and characterized via infrared spectroscopy (3.1.).

3. Results

3.1 Infrared spectroscopy

3.1.1 Qualitative analysis

Figure 2(a, b) displays the infrared (IR) spectra of RDX before and after the irradiation in the region from 4000 to 600 cm^{-1} . Detailed assignments of the observed infrared bands are provided in Table 2. The infrared spectrum of RDX reveals prominent NO_2 -stretch, N-N stretch, CH_2 -bending and ring vibrational bands in the region 1600-500 cm^{-1} along with CH_2 -stretches in the 3100-3000 cm^{-1} region; these findings are in excellent agreement with the infrared spectrum of RDX reported in the literature.⁵⁵ The thin film of RDX can exist in crystalline or amorphous phase. The crystalline form of RDX shows sharp IR bands with narrow band width while amorphous phase of RDX exhibit broad and diffuse bands.²⁷ The IR spectra of RDX film measured in the present study and that of crystalline phase recorded by Botcher et al. are compared in Figure S2 of the supporting information. Figure S2 unveils that the thin film of RDX used in this work is amorphous. The fundamentals at 3068 and 3006 cm^{-1} correspond to CH_2 asymmetric and symmetric stretching modes, respectively. The CH_2 bending modes (in-plane and out-of-plane) are observed in the spectral range from 1450-1380 cm^{-1} . A group of three bands 1596, 1579 and 1531 cm^{-1} can be attributed to the antisymmetric stretching modes of $-\text{NO}_2$ group. The characteristic N- NO_2 stretching vibration of RDX appears at 1352 and 1322 cm^{-1} . The spectral regions 1240-1210 cm^{-1} , 1050-950 cm^{-1} and 880-600 cm^{-1} are dominated by ring vibrations. Bands observed in the region 1240-1210 cm^{-1} are linked to the N-C-N stretching modes of the ring. Symmetric vibration modes of the ring - also known as ring breathing - appear at 879 cm^{-1} ; bands at 840 and 784 cm^{-1} are attributed to C-N-C stretching vibrations.

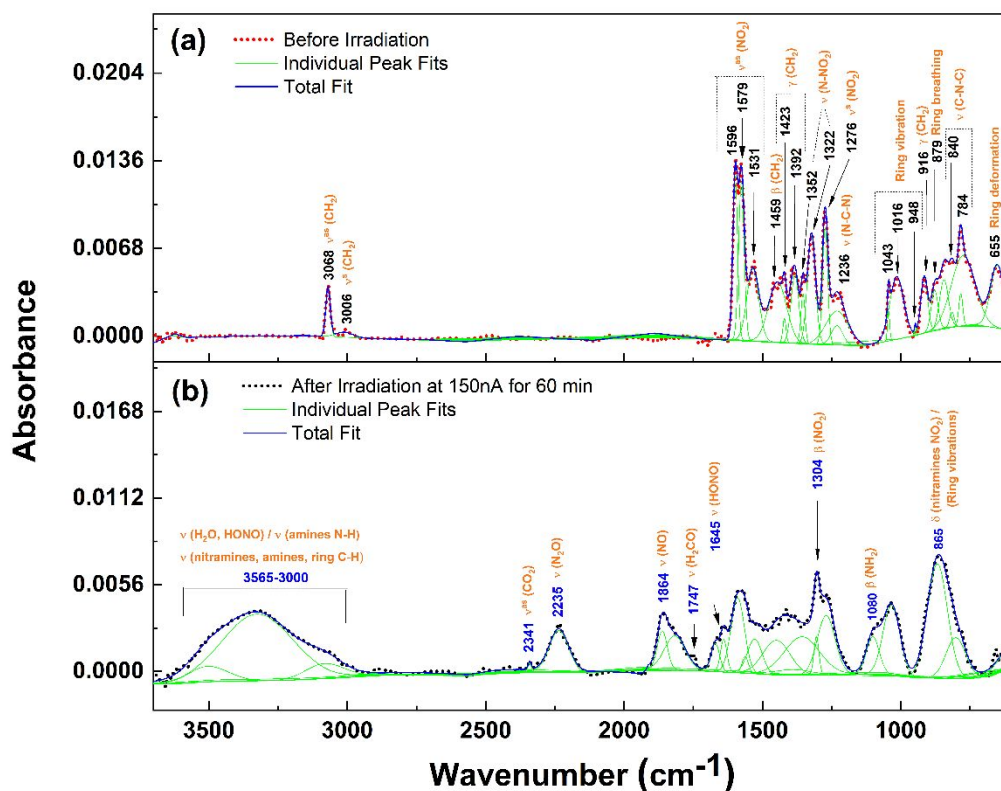


Figure 2. Infrared spectra of RDX collected at 5 K (a) before and (b) after the irradiation of the high dose experiment.

The infrared spectrum after the irradiation is distinct from the data obtained for the pure RDX sample (Figure 2b); the fundamentals of RDX decrease in column density by $40 \pm 10 \%$ (low dose) and $72 \pm 14 \%$ (high dose); new bands appear in the region 3565-3000, 2341, 2235, 1864, 1747, 1645, 1304, 1080 and 865 cm^{-1} . These new features can be assigned to primary and secondary decomposition products of RDX. Bands at 2341, 2235, 1864 and 1747 cm^{-1} can be associated with stretching vibrations of carbon dioxide (CO_2), dinitrogen oxide (N_2O), nitrogen monoxide (NO), and formaldehyde (H_2CO), respectively. Absorptions corresponding to secondary products (CO_2 , N_2O , NO , and H_2CO) have been observed by Botcher et al. and Alix et al. during decomposition of RDX via laser pyrolysis and photolysis.^{27, 56} The band observed at 1645 cm^{-1} can be allocated to the fundamental N=O stretch of nitrous acid (HONO) which is in good agreement with the 1640 cm^{-1} assigned previously to the cis conformer of HONO .⁵⁷

Table 2. (a) Infrared features of RDX before the irradiation along with (b) new bands observed in the spectrum in the high dose experiment at 5 K.

(a) Before Irradiation			
Wavenumber Observed (cm ⁻¹)	Wavenumber Literature (cm ⁻¹) ⁵⁵	Vibrational Assignments	Vibrational Modes
3068	3068	ν ^{as} (CH ₂)	C-H asymm. stretch
3006	3004	ν ^s (CH ₂)	C-H symm. stretch
1596	1593	ν ^{as} (NO ₂)	NO ₂ asymm. stretch
1579	1576	ν ^{as} (NO ₂)	NO ₂ asymm. stretch
1531	1535	ν ^{as} (NO ₂)	NO ₂ asymm. stretch
1459	1460	β(CH ₂)	CH ₂ Bending in plane
1435	1435	β(CH ₂)	CH ₂ Bending in plane
1423	1424	γ(CH ₂)	CH ₂ Bending out of plane
1392	1391	γ(CH ₂)	CH ₂ Bending out of plane
1352	1352	ν ^s (N-NO ₂)	N-N symm. stretch
1322	1322	ν ^s (N-NO ₂)	N-N symm. stretch
1276	1275	ν ^s (NO ₂)	NO ₂ symm. stretch
1236	1232	ν(N-C-N)	Ring skeletal vibrations
1218	1219	ν(N-C-N)	Ring skeletal vibrations
1043	1040	ν ^{as} (Ring)	Ring symm. vibrations
1016	1020	ν ^{as} (Ring)	Ring symm. vibrations
948	947	ν ^{as} (Ring)	Ring symm. vibrations
916	917	γ(CH ₂)	CH ₂ Bending out of plane
879	882	ν ^s (Ring)	Ring symm. vibration
840	844	ν(C-N-C)	Ring skeletal vibrations
784	790	ν(C-N-C)	Ring skeletal vibrations
655	670	δ(Ring)	Ring deformation
(b) After Irradiation			
3565-3000		ν(H ₂ O)	O-H stretch of H ₂ O
		ν(HONO)	O-H stretch of HONO
		ν(NH ₂)	N-H stretch of amines
		ν(CH ₂ NNO ₂)	C-H stretch of nitramines
		ν(NH ₂)	C-H stretch of amines
		ring	C-H stretch of ring
2341	2342 ²⁷	ν(CO ₂)	C=O stretch of CO ₂
2235	2237 ²⁷	ν(N ₂ O)	N=N stretch of N ₂ O
1864	1864 ²⁷	ν(NO)	Free NO stretch
1747	1742 ⁵⁶	ν(H ₂ CO)	C=O stretch of formaldehyde
1645	1640 ⁵⁷	ν(HONO)	N=O stretch of HONO
1304	1304 ⁵⁸	β(NO ₂)	NO ₂ deformation
1080		β(NH ₂)	NH ₂ bending modes of amine
865	871 ²⁷	δ(NNO ₂)	Deformation modes of nitramines / ring vibrations

Further, the characteristic bending vibration of nitrogen dioxide (NO₂) is observed at 1304 cm⁻¹, which matches nicely the reported literature value of 1304 cm⁻¹.⁵⁸ The stretching vibration of NO₂ (1613 cm⁻¹) is masked by the broad absorption feature of RDX-NO₂ (stretching vibrations; 1600-1520 cm⁻¹). We also observe a feature at 1080 cm⁻¹, merged with the absorption band of

RDX; this can be tentatively assigned to bending mode of amines ($-\text{NH}_2$).⁵⁹⁻⁶¹ The absorption band observed at 865 cm^{-1} could be linked to decomposition product methylene nitramine (CH_2NNO_2) based on its calculated vibrational frequencies reported by Mowrey et al.⁶² This assignment is tentative as it is merged with the ring vibrations of RDX. In fact, probing the formation of methylene nitramine (CH_2NNO_2) during RDX decomposition is difficult via infrared spectroscopy as most of its vibrational bands are obscured by the absorption bands of RDX.²⁷ It is also important to highlight that evidence of formation of nitrous acid (HONO), nitrogen dioxide (NO_2) and amines ($-\text{NH}_2$) from the decomposition of RDX have not been reported previously through IR spectroscopy. Their signatures have been reported only through mass spectrometry.^{4, 9, 24} Figure 2(b) also shows broad absorption band covering the spectral range from $3565\text{--}3000\text{ cm}^{-1}$. This broad feature is due to multiple vibrations in the region $3565\text{--}3000\text{ cm}^{-1}$ which include O-H stretch (H_2O , HONO), N-H stretch (amines) and C-H stretch (amines, nitramines, ring).^{57, 59, 61-63}

3.1.2 Quantitative analysis: Mass balance

The column densities were calculated using a modified Lambert-Beer law to determine the amount of RDX molecules destroyed and that of products formed.⁶⁴ The integrated absorption co-efficient of the reactant and products used for the calculation of column densities are taken from literature. The absorption coefficient of RDX band at 1596 cm^{-1} is $1.0 \times 10^{-18}\text{ cm molecule}^{-1}$. The integrated band strength of bending mode of NO_2 (1304 cm^{-1}) is $6.2 \times 10^{-18}\text{ cm molecule}^{-1}$.⁶⁵ The N=O stretch of NO (1864 cm^{-1}) and HONO (1645 cm^{-1}) hold integrated band strengths of 6.8×10^{-18} and $7.2 \times 10^{-18}\text{ cm molecule}^{-1}$ respectively, while the N=N stretch of N_2O (2235 cm^{-1}) has an absorption coefficient of $5.7 \times 10^{-17}\text{ cm molecule}^{-1}$.⁶⁴⁻⁶⁶ Carbon dioxide (CO_2) was quantified via the stretching mode (2341 cm^{-1}) which has an absorption coefficient of $7.6 \times 10^{-17}\text{ cm molecule}^{-1}$.⁴⁵ The column density of H_2O (water) is measured in the region of $3500\text{--}3100\text{ cm}^{-1}$ using an integrated band strength of $2.0 \times 10^{-16}\text{ cm molecule}^{-1}$.⁶⁷ It is important to note here that the broad absorption in the region $3567\text{--}3000\text{ cm}^{-1}$ includes O-H stretch of HONO, amines N-H stretch and ring C-H stretch along with O-H stretch of H_2O . Therefore, to determine the column density of water only region $3500\text{--}3100\text{ cm}^{-1}$ is considered where O-H stretch of H_2O has maximum absorption. The amount of RDX molecules destroyed via the radiolysis (high dose) is $1.3 \pm 0.3 \times 10^{16}\text{ molecules cm}^{-2}$, i.e. 72 % of the initial column density ($1.8 \pm 0.3 \times 10^{16}\text{ molecules cm}^{-2}$). On the other hand, the total number of molecules of NO_2 ($4.5 \pm 0.4 \times 10^{14}\text{ molecules cm}^{-2}$),

HONO ($4.4 \pm 0.4 \times 10^{14}$ molecules cm^{-2}), NO ($1.5 \pm 0.2 \times 10^{14}$ molecules cm^{-2}), CO₂ ($2.6 \pm 0.3 \times 10^{13}$ molecules cm^{-2}), N₂O ($7.1 \pm 0.7 \times 10^{13}$ molecules cm^{-2}) and H₂O ($6.0 \pm 0.6 \times 10^{14}$ molecules cm^{-2}) accounts for only 9 ± 1 % of the decomposed RDX molecules. Therefore, we can conclude that nearly 90% of RDX molecules decomposes into products, which remained unobserved via infrared spectroscopy due to overlapping fundamentals with the RDX reactant.

In summary, our infrared spectroscopic studies provide clear evidence of formation of critical decomposition products such as water (H₂O), carbon dioxide (CO₂), dinitrogen oxide (N₂O), nitrogen monoxide (NO), formaldehyde (H₂CO), nitrous acid (HONO) and nitrogen dioxide (NO₂) after the irradiation of RDX (Figure 3). Although, presence of primary products such as methylene nitramine (CH₂NNO₂), amines, and cyclic as well as acyclic intermediates cannot be unraveled with absolute certainty through FTIR spectroscopy due to functional groups overlapping with RDX, their existence is inferred indirectly based on the appearance of broad absorption bands in the region 3565-3000 cm^{-1} , 1100-1000 cm^{-1} and 950-850 cm^{-1} , where vibrational frequencies of amines, nitramines and ring vibrations are observable.

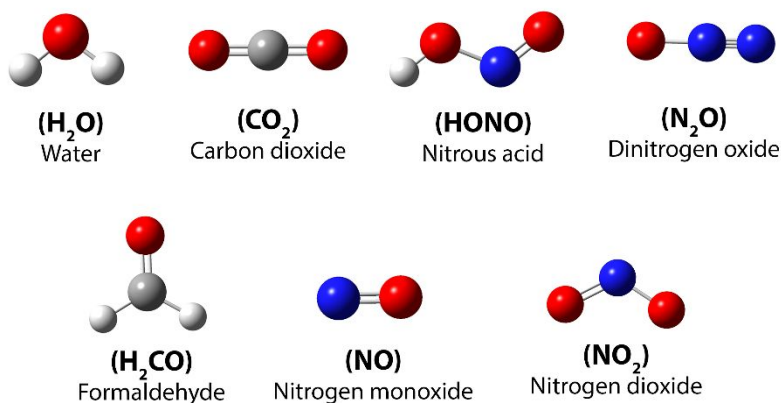


Figure 3. Structures of the decomposition products detected through FTIR spectroscopy.

3.2 UV-VIS spectroscopy

Figure 4 shows the UV-VIS spectra of RDX film measured before and after the irradiation in the spectral range of 190-800 nm. UV-VIS spectrum of RDX prior to the irradiation displays an absorption maximum at around 207 nm and a shoulder at around 236 nm. The absorption at 207 and 237 nm is primarily due to the $\pi \rightarrow \pi^*$ transition from the nitro group; the broad absorption around 236 nm has contributions from the weak $n \rightarrow \pi^*$ transition.⁶⁸⁻⁶⁹ After the irradiation, a

decrease in the intensity of the absorption at 236 nm and an increase in the intensity of absorption maxima at 207 nm are observed. Furthermore, the maximum at around 207 nm is slightly blue shifted after the irradiation. Our UV spectra measured before and after the irradiation of RDX is in good agreement with that of reported by Gares et al. and Makarov et al.^{26, 33} The increase in the intensity of the absorption at 207 nm after the irradiation is due to accumulation of decomposition products of RDX such as dinitrogen oxide (N_2O), methylene nitramine (CH_2NNO_2), diazomethane (CH_2N_2), nitrogen dioxide (NO_2) which contribute through their $\pi \rightarrow \pi^*$ transitions to the 207 nm band.³³

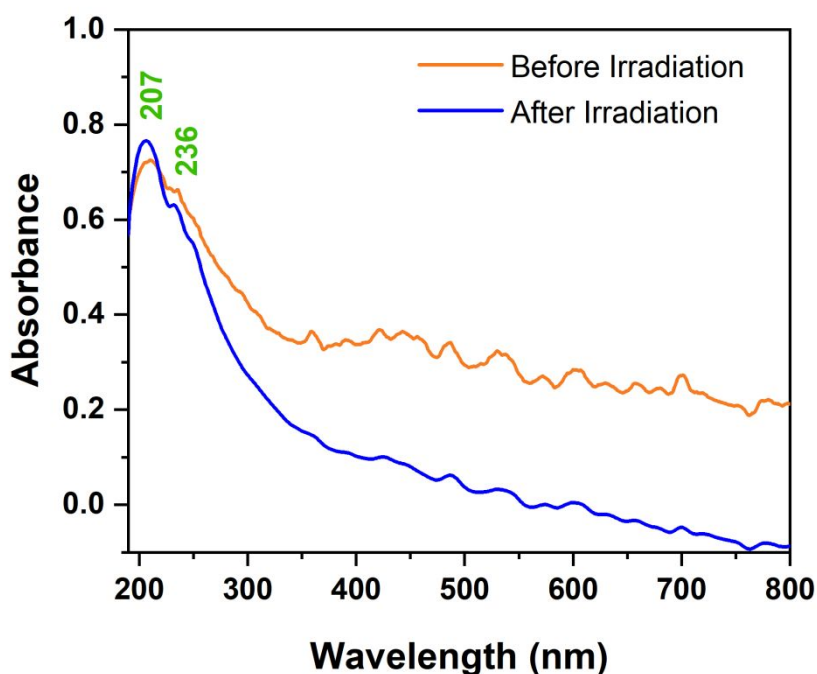


Figure 4. UV-VIS spectra of RDX collected at 5 K before and after the high dose.

3.3 Single Photon Ionization Reflectron Time-of-Flight Mass Spectrometry

To identify individual molecules formed in the decomposition of RDX, single photon ionization reflectron time-of-flight mass spectrometry (PI-ReTOF-MS) was exploited;⁷⁰⁻⁷⁷ this approach allows the detection of a broad range of products based on their mass-to-charge ratio (m/z) and sublimation temperature. Figure 5 (a) and (b) depicts the temperature dependent mass spectra of the ionized decomposition products of the irradiated RDX samples at the high dose and low dose experiments, respectively. It is apparent that the ion counts of the decomposition

products formed after the irradiation are enhanced with the dose, i.e. higher ion counts as the dose increases. Further, the ion counts of the lower molecular weight products are enhanced compared to the higher molecular weight species in the high dose experiments proposing that

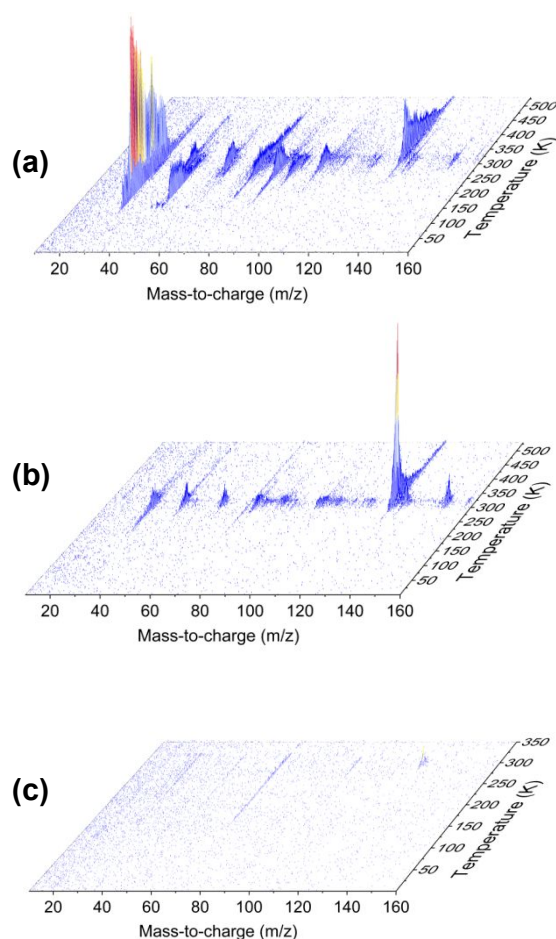
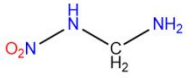
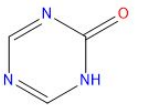
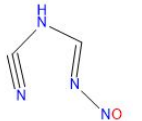
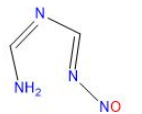
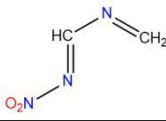
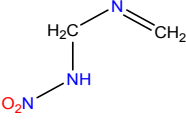
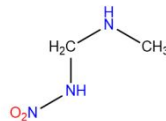
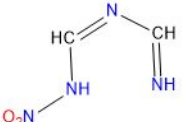
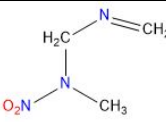
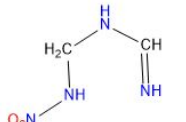


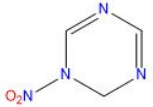
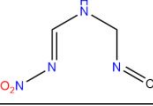
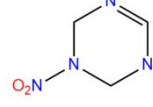
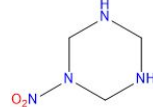
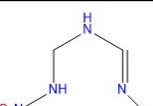
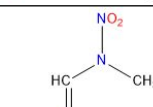
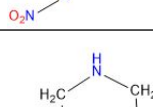
Figure 5. Temperature dependent ReTOF data of the decomposition products of RDX for the high dose (a), low dose (b), and blank (c) experiments.

initially formed high molecular (possibly primary decomposition products) subsequently decompose to lower molecular products as the dose increases. Overall, we observed species holding mass-to-charge (m/z) values of 30, 31, 42, 43, 45, 46, 56, 58, 60, 70, 71, 72, 73, 74, 75, 81, 82, 83, 85, 87, 89, 91, 97, 98, 100, 101, 103, 105, 116, 117, 118, 128, 130, 132, 148, 149, 150, and 157 in the ReTOF after the high dose irradiation. All masses are also observed at the low dose experiment with the exception of $m/z = 31, 42, 45, 60, 70$, and 117. The assigned molecular formulae and potential structures of all these species are shown in Table 4.

Table 4. Molecular formula, structure and ionization energies of the masses observed in ReTOF mass spectrometer via photoionization at 10.49 eV. Masses marked with green color are reported previously by both experiment and theoretical calculations, while the masses marked with blue color are observed only in previous experimental studies and masses marked with red color are supported only through theoretical calculations reported in the literature

m/z	Molecular formula	Structure	I.E. (eV)	Low Dose	High Dose
30	NO		9.26	+	+
31	HNO		10.1		+
	CH ₃ NH ₂		8.90		
42	CH ₂ N ₂		8.99		+
43	C ₂ H ₃ N		9.30		+
			9.20		
45	CH ₃ NO		10.16		+
			9.30		
			10.11		
46	NO ₂		9.58	+	+
56	C ₂ H ₄ N ₂		8.95	+	+
58	CH ₂ N ₂ O		-		+
	C ₂ H ₆ N ₂			+	
60	CH ₄ N ₂ O		-		+
70	C ₃ H ₆ N ₂		-		+
71	C ₂ H ₅ N ₃			+	+
72	C ₃ H ₈ N ₂		-	+	+
73	C ₂ H ₇ N ₃		-	+	+
74	CH ₂ N ₂ O ₂			+	+
75	C ₂ H ₉ N ₃			+	+
81	C ₃ H ₃ N ₃		9.80	+	+
82	C ₃ H ₄ N ₃			+	+
83	C ₃ H ₅ N ₃			+	+
85	C ₃ H ₇ N ₃			+	+
87	C ₃ H ₉ N ₃			+	+
89	CH ₃ N ₃ O ₂			+	+

m/z	Molecular formula	Structure	I.E. (eV)	Low Dose	High Dose
91	CH ₅ N ₃ O ₂			+	+
97	C ₃ H ₃ N ₃ O			+	+
98	C ₂ H ₂ N ₄ O			+	+
100	C ₂ H ₄ N ₄ O			+	+
101	C ₂ H ₃ N ₃ O ₂			+	+
103	C ₂ H ₅ N ₃ O ₂			+	+
105	C ₂ H ₇ N ₃ O ₂			+	+
116	C ₂ H ₄ N ₄ O ₂			+	+
117	C ₃ H ₇ N ₃ O ₂				+
118	C ₂ H ₆ N ₄ O ₂			+	+

m/z	Molecular formula	Structure	I.E. (eV)	Low Dose	High Dose
128	C ₃ H ₄ N ₄ O ₂			+	+
130	C ₃ H ₆ N ₄ O ₂			+	+
132	C ₃ H ₈ N ₄ O ₂			+	+
148	C ₂ H ₄ N ₄ O ₄			+	+
149	C ₂ H ₇ N ₅ O ₃			+	+
150	C ₂ H ₆ N ₄ O ₄			+	+
157	C ₃ H ₃ N ₅ O ₃			+	+

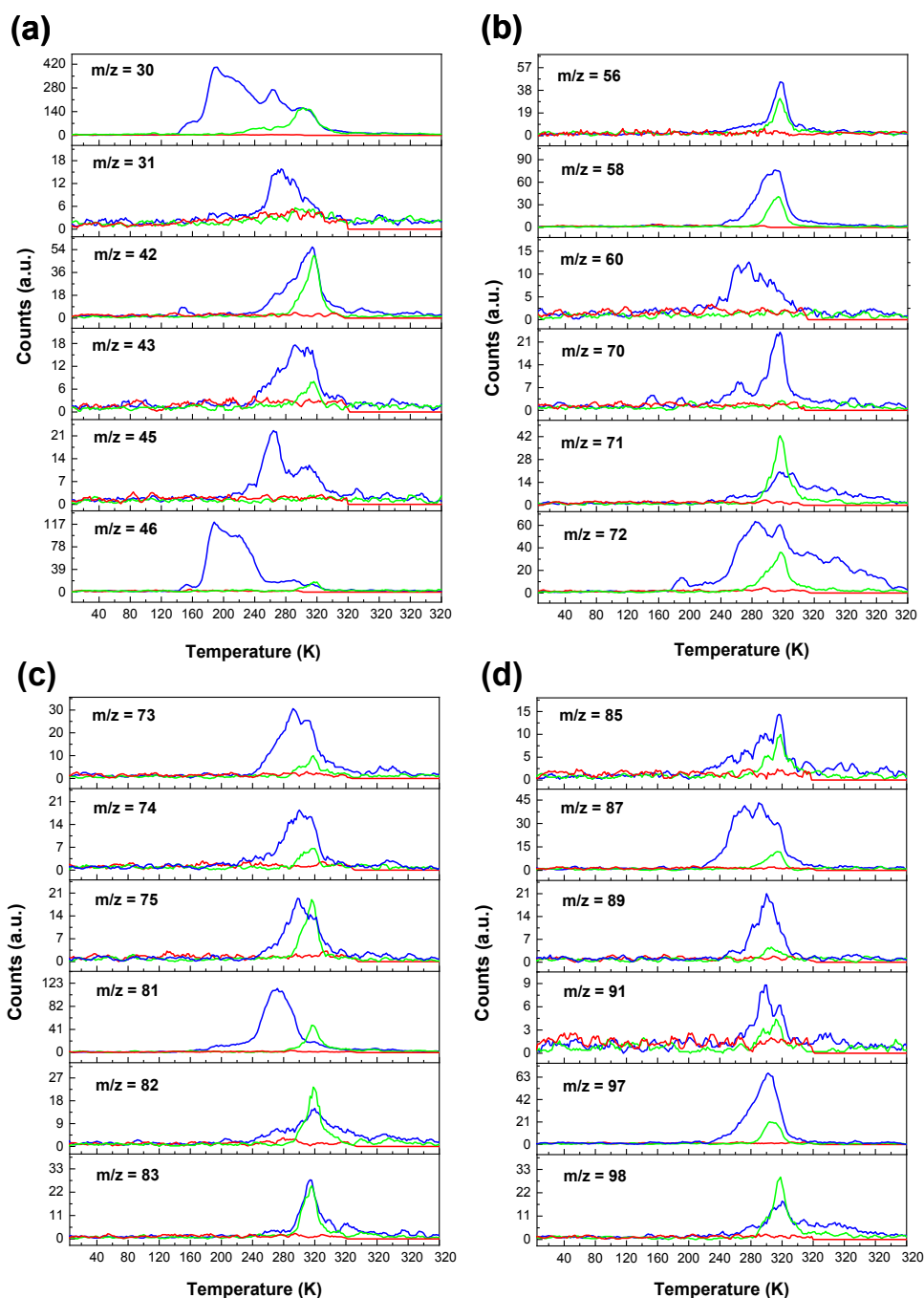


Figure 6. TPD profiles of the ion counts recorded at mass-to-charge ratios of (a) 30, 31, 42, 43, 45, 46, (b) 56, 58, 60, 70, 71, 72, (c) 73, 74, 75, 81, 82, 83, and (d) 85, 87, 89, 91, 97 and 98 at a photoionization energy of 10.49 eV. Blue: high dose; green: low dose; red: blank.

It is essential to mention that the ReTOF data collected in blank experiments - experiments conducted in the similar fashion, but without exposing the samples to energetic electrons (Figure

5c) - did not show any signal which ensures that the decomposition products observed in the irradiation experiments are solely due to the impinging ionizing radiation. The corresponding TPD traces are compiled in Figure 6.

Figure 6(a) shows the TPD profiles of $m/z = 30, 31, 42, 43, 45$ and 46 . The ion signal intensities at $m/z = 30, 43$, and 46 are lower by a factor of about three in the low dose experiments. Signal at $m/z = 30$ can be assigned to nitrogen monoxide (NO ; IE = 9.26 eV), which has been previously observed in the decomposition of RDX.^{4, 7, 9} Ion signal at $m/z = 31$ can be either assigned to nitrosyl hydride (HNO ; IE = 10.1 eV) and/or methylamine (CH_3NH_2 ; IE = 8.90 eV). The maximum of the ion count profile at $m/z = 31$ at 280 K correlates with the second sublimation event recorded at $m/z = 30$ at 280 K. Therefore, ion counts at $m/z = 30$ at around 280 K likely represent fragments of $m/z = 31$ (CH_5N^+ ; HNO^+), i.e. CH_4N^+ and/or NO^+ . Lossing *et al.* demonstrated an appearance energy (AE) of AE = 10.18 eV for CH_4N^+ from CH_5N^+ .⁷⁸ Therefore, we may conclude that at least some of the ion counts at $m/z = 31$ and 30 originate from singly ionized methylamine (CH_5N^+) and its CH_4N^+ fragment, respectively. It is important to note that $m/z = 31$ has not been identified in any previous experimental study investigating the decomposition of RDX. Ion signal at $m/z = 42$ can be associated with the molecular formula CH_2NN (diazomethane; IE = 8.99 eV), which has been observed in previous experimental studies.^{4, 9, 28, 31} Two molecules with the molecular formula $\text{C}_2\text{H}_5\text{N}$ can contribute to $m/z = 43$: ethylenimine ($\text{c-C}_2\text{H}_4\text{NH}$) and N-methylmethanimine (CH_3NCH_2). The maximum of the ion counts measured at $m/z = 43$ and $m/z = 42$ at 320 K correlates very well, suggesting that some of the ion counts measured at $m/z = 42$ could be due to fragmentation of $\text{C}_2\text{H}_5\text{N}^+$. Photofragmentation of ethylenimine ($\text{c-C}_2\text{H}_4\text{NH}$) into $\text{C}_2\text{H}_4\text{N}^+$ has been observed by Scala *et al.* at 8.4 eV. On the other hand, photofragmentation of N-methylmethanimine (CH_3NCH_2) at < 9.8 eV mostly leads to CH_3NH^+ .⁷⁹ Therefore, it is highly likely that some of the ion counts observed at $m/z = 42$ are due to fragmentation of $\text{c-C}_2\text{H}_4\text{NH}^+$ ($m/z = 43$) to $\text{c-C}_2\text{H}_4\text{N}^+$. Ion counts at $m/z = 45$ can be assigned to isomers of molecular formula CH_3NO : nitrosomethane (CH_3NO), formamide (HCONH_2), and formaldehyde oxime (H_2CNOH). The TPD profile measured at $m/z = 45$ shows two maxima, which correlate well with the maxima observed in the TPD profile of $m/z = 30$ at around 280 K and 320 K. This implies that ion signal at $m/z = 30$ at around 280 K and 320 K could be due to NO^+ ions generated as a result of photofragmentation of nitrosomethane.⁸⁰ Therefore, we can conclude that some of the ion counts at $m/z = 45$ and m/z

=30 are due to the nitrosomethane ion (CH_3NO^+) and its fragment NO^+ , respectively. Finally, ion signal at $m/z = 46$ can be assigned to nitrogen dioxide (NO_2 ; IE = 8.80 eV). Fragmentation of $m/z = 46$ (NO_2^+) to $m/z = 30$ (NO^+) can be easily presumed based on the fact that the TPD profile measured at $m/z = 46$ in the temperature range of 160-240 K is very similar to that of measured at $m/z = 30$. Dibeler et al. have demonstrated photofragmentation of nitrogen dioxide into NO^+ at 12.34 eV.⁸¹ All three m/z values of 43, 45 and 46 have been reported previously in various experimental studies investigating the decomposition of RDX.^{4, 9, 28, 34}

The TPD profiles of $m/z = 56, 58, 60, 70, 71$ and 72 are shown in Figure 6(b). $m/z = 56$ was identified by Dickinson et al. and Zhao et al. while investigating the decomposition of RDX in the solid state and in the gas phase, respectively.^{9, 34} $m/z = 56$ can be associated with the molecular formula $\text{C}_2\text{H}_4\text{N}_2$ (Table 4). Ion signal at $m/z = 58$ can be either allocated to the mononitroso form of methylenenitramine (H_2CNNO_2) holding the molecular structure H_2CNNO or to a primary amine moiety with the molecular formula $\text{C}_2\text{H}_6\text{N}_2$ (Table 4). Evidence of mass 58 has been reported earlier by Behrens and co-workers in thermal decomposition of RDX crystals.^{4, 24} Decomposition product at $m/z = 60$ can be accounted for as the hydrogenated form of $m/z = 58$ (H_2CNNO and/or $\text{C}_2\text{H}_6\text{N}_2$) and hence might be assigned to the molecular formula $\text{CH}_4\text{N}_2\text{O}$ and/or $\text{C}_2\text{H}_8\text{N}_2$. Unlike $m/z = 56$ and 58 , evidence of $m/z = 60$ has not been reported in previous studies. Ion signal at $m/z = 70, 71$, and 72 is linked to the molecular formulae $\text{C}_3\text{H}_6\text{N}_2$, $\text{C}_2\text{H}_5\text{N}_3$, and $\text{C}_3\text{H}_8\text{N}_2$, respectively (Table 4). The peak of the sublimation event of $m/z = 72$ at 320 K matches the maximum of the TPD profile at $m/z = 71$. Therefore, at least some ion counts at $m/z = 71$ likely represent fragments of $\text{C}_3\text{H}_8\text{N}_2^+$ ($m/z = 72$): $\text{C}_3\text{H}_7\text{N}_2^+$ ($m/z = 71$).

Figure 6(c) shows TPD profiles recorded at $m/z = 73, 74, 75, 81, 82$, and 83 . Like $m/z = 71$ ($\text{C}_2\text{H}_5\text{N}_3$), ion signal at $m/z = 73$ can be associated to an amine moiety. In fact, $m/z = 73$ can be designated to the hydrogenated counterpart of $\text{C}_2\text{H}_5\text{N}_3$ ($m/z = 71$) and, therefore, might be assigned to the molecular formula $\text{C}_2\text{H}_7\text{N}_3$ (Table 4). Similarly, $m/z = 75$ is attributed to the molecular formula $\text{C}_2\text{H}_9\text{N}_3$ since it might represent the hydrogenated analog of $m/z = 73$ ($\text{C}_2\text{H}_7\text{N}_3$) and/or 71 ($\text{C}_2\text{H}_5\text{N}_3$). It is important to mention that formation of higher molecular weight amine-based species have been predicted by *ab initio* molecular dynamics simulations, but these molecules have never observed in experiments so far.¹⁶ In this study, both mass and FTIR spectra (section 3.1) reveal evidence of formation of amines in the decomposition of RDX.

Ion signal at $m/z = 74$ has been observed in several experimental studies and can be allocated to methylene nitramine (H_2CNNO_2), formally the monomeric unit of RDX.⁹ Interestingly, TPD profiles measured at $m/z = 73$, 74, and 75 show peak maximum at the same temperature at 300 K. This could be either due to co-sublimation of the products associated with $m/z = 73$, 74, and 75 or fragmentation of one species into another. Photofragmentation of $\text{C}_2\text{H}_9\text{N}_3$ ($m/z = 75$) to $\text{C}_2\text{H}_7\text{N}_3^+$ ($m/z = 73$) is highly viable. Similarly, the fragment of $\text{C}_2\text{H}_7\text{N}_3^+$ ($m/z = 73$) - $\text{C}_2\text{H}_6\text{N}_3^+$ - could contribute to the ion counts at $m/z = 72$.

Further, ion signals at $m/z = 81$, 82, and 83 are assigned to the molecular formulae $\text{C}_3\text{H}_3\text{N}_3$, $\text{C}_3\text{H}_4\text{N}_3$, and $\text{C}_3\text{H}_5\text{N}_3$, respectively. These intermediates carry ring moieties of RDX and have been detected in gas phase as well as condensed phase decomposition of RDX.^{9, 34} The fragment of $\text{C}_3\text{H}_5\text{N}_3$ ($m/z = 83$) - $\text{C}_3\text{H}_4\text{N}_3^+$ - could account for the ion counts measured at $m/z = 82$. Higher masses carrying the ring moiety such as $m/z = 85$ and 87 are also observed in our experiments, and their TPD profiles are shown in Figure 6(d). It is intriguing to note here that $m/z = 85$ and 87 have not been identified in previous studies. Figure 6(d) further shows TPD profiles of $m/z = 89$, 91, and 97. Ion signal at $m/z = 89$ and 91 can be assigned to the molecular formulae $\text{CH}_3\text{N}_3\text{O}_2$ (aminomethylenenitramine) and $\text{CH}_5\text{N}_3\text{O}_2$ (aminomethylnitramine) respectively. Both molecules with likely structures presented in Table 4 have not been detected before.

The maximum of the ion counts measured at $m/z = 89$ and 91 appears at the same temperature of 300 K suggesting that some of the ion counts at $m/z = 89$ arise from fragmentation of $\text{CH}_5\text{N}_3\text{O}_2^+$ ($m/z = 91$) into $\text{CH}_3\text{N}_3\text{O}_2^+$ ($m/z = 89$). Signal at $m/z = 97$ is associated to oxy-s-triazine (OST) with molecular formula $\text{C}_3\text{H}_3\text{N}_3\text{O}$. Until now, evidence of OST has been observed only during thermal decomposition of RDX in the liquid phase.^{4, 24}

The plots of ion signal intensities as a function of temperature at higher mass-to-charge ratios of $m/z = 100$, 101, 103, 105, 116, 117, and 118 are compiled in Figure 6(e). Interestingly, the ion signal intensities of these TPD profiles are nearly identical in low dose and high dose irradiation experiments. Ion signal at $m/z = 100$ has the same m/z as the mononitroso intermediate holding the molecular formula $\text{C}_2\text{H}_4\text{N}_4\text{O}$. The TPD profile at $m/z = 100$ and $m/z = 98$ are very similar suggesting that some of the ion counts at $m/z = 98$ are fragments of $m/z = 100$ ($\text{C}_2\text{H}_4\text{N}_4\text{O}^+$) - $\text{C}_2\text{H}_2\text{N}_4\text{O}^+$. Ion counts at m/z 101, 103, and 105 are likely associated to mononitro species with the molecular formulae $\text{C}_2\text{H}_3\text{N}_3\text{O}_2$, $\text{C}_2\text{H}_5\text{N}_3\text{O}_2$, and $\text{C}_2\text{H}_7\text{N}_3\text{O}_2$, respectively. It is intriguing to

note here that fragmentation of $\text{C}_2\text{H}_5\text{N}_3\text{O}_2^+$ ($m/z = 103$) into $\text{C}_2\text{H}_3\text{N}_3\text{O}_2^+$ ($m/z = 101$) is highly feasible. Although these mononitro species have not been identified in previous experimental studies, formation of the $\text{C}_2\text{H}_5\text{N}_3\text{O}_2$ and $\text{C}_2\text{H}_3\text{N}_3\text{O}_2$ radicals have been predicted by theoretical calculations.¹⁶ Ion signal at $m/z = 116$, 117 , and 118 are proposed to be associated with mononitro intermediates with the possible molecular formulae $\text{C}_2\text{H}_4\text{N}_4\text{O}_2$, $\text{C}_3\text{H}_7\text{N}_3\text{O}_2$, and $\text{C}_2\text{H}_6\text{N}_4\text{O}_2$ respectively. Ion counts at $m/z = 116$ could be due to fragmentation of $\text{C}_2\text{H}_6\text{N}_4\text{O}_2^+$ ($m/z = 118$) into $\text{C}_2\text{H}_4\text{N}_4\text{O}_2^+$ since the TPD profile recorded at $m/z = 116$ peaking at 280 and 320 K correlates well with two of the ion count maxima observed at 280 K and 320 K in the sublimation profile measured at $m/z = 118$. Evidence of these species ($m/z = 116$, 117 , 118) has not been reported through previous experimental or theoretical calculations.

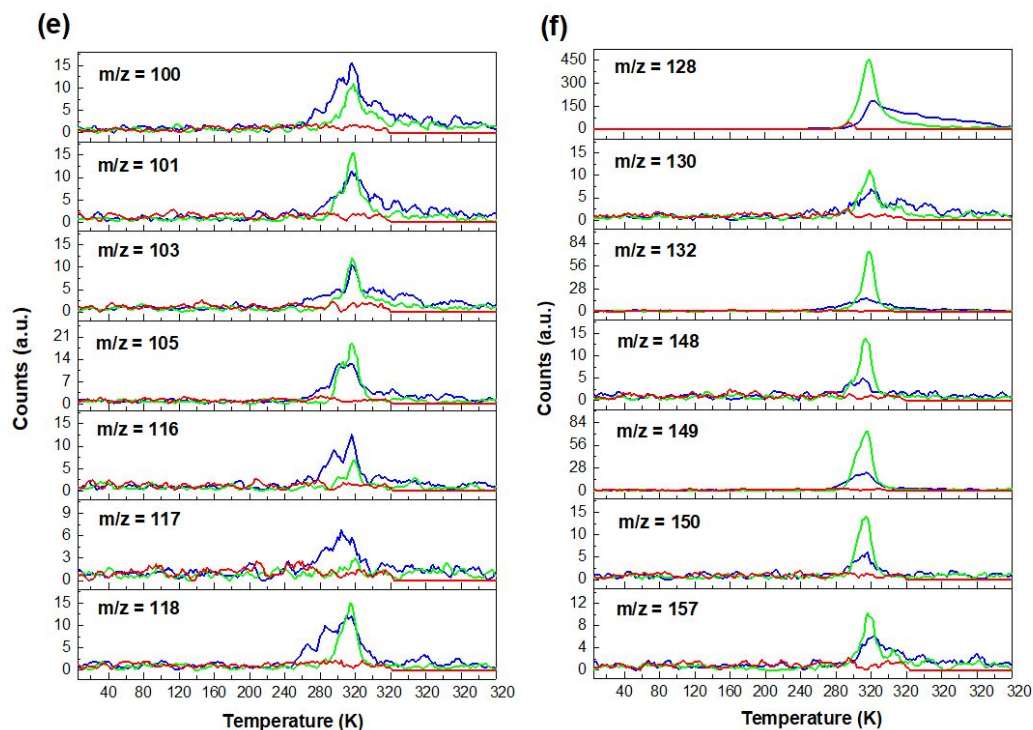


Figure 6. TPD profiles of the ion counts recorded at mass-to-charge ratios of (e) 100, 101, 103, 105, 116, 117, 118, (f) 128, 130, 132, 148, 149, 150 and 157 at a photoionization energy of 10.49 eV. Blue: high dose; green: low dose; red: blank.

Finally, TPD profiles of $m/z = 128$, 130 , 132 , 148 , 149 , 150 , and 157 are shown in Figure 6(f). Ion counts are higher by around 2 times in low dose relative to high dose irradiation experiments. Signal at $m/z = 128$ is allocated to molecular formula $\text{C}_3\text{H}_4\text{N}_4\text{O}_2$ whose structure consists of the

ring moiety of RDX with only one nitro group (Table 4 for structure). $m/z = 128$ has the highest signal intensity compared to all remaining masses in low dose irradiation experiment. Ion signal at $m/z=130$ can be considered as hydrogenated form of $m/z=128$ and hence might be assigned to $C_3H_6N_4O_2$. Evidence of formation of $m/z = 130$ has not been shown in previous experimental studies, but gas phase theoretical calculations predict its existence.¹⁶ Similarly, $m/z = 132$ could be designated as hydrogenated form of $m/z 130$; its formation has been suggested by both experiments as well as theoretical calculations.^{4, 9, 16, 24} Based on the similarity between the TPD profiles measured at $m/z=130$ and 128, it can be envisaged that $C_3H_6N_4O_2^+$ ($m/z = 130$) could fragment into $C_3H_4N_4O_2^+$ ($m/z = 128$). Dinitro intermediates of RDX like $m/z = 148$ and 150 are also observed in our experiment and are allocated to molecular formulae $C_2H_4N_4O_4$ and $C_2H_6N_4O_4$, respectively (Table 4). Signal at $m/z = 150$ can be considered as a hydrogenated form of $m/z = 148$; only the later was detected during thermal decomposition of RDX.^{9, 14, 16} We have also observed intermediates belonging to nitro-nitroso species, i.e. species carrying both a nitro and nitroso group via at $m/z = 149$ and 157. These species are assigned to molecular formulae $C_2H_7N_5O_3$ and $C_3H_3N_5O_3$, respectively. This is the first experimental evidence of formation of intermediates having both nitro and nitroso groups during the decomposition of RDX.

In summary, our ReTOF mass spectroscopic studied revealed several new decomposition products of RDX that have not been reported before. These include HNO ($m/z = 31$), CH_5N_2 ($m/z = 31$), CH_4N_2O ($m/z = 60$), $C_2H_5N_3$ ($m/z = 71$), $C_3H_8N_2$ ($m/z = 72$), $C_2H_9N_3$ ($m/z = 75$), $C_3H_7N_3$ ($m/z = 85$), $C_3H_9N_3$ ($m/z = 87$), $CH_3N_3O_2$ ($m/z = 89$), $CH_5N_3O_2$ ($m/z = 91$), $C_2H_2N_4O$ ($m/z = 98$), $C_2H_4N_4O$ ($m/z = 100$), $C_2H_3N_3O_2$ ($m/z = 101$), $C_2H_5N_3O_2$ ($m/z = 103$), $C_2H_7N_3O_2$ ($m/z = 105$), $C_2H_4N_4O_2$ ($m/z = 116$), $C_3H_7N_3O_2$ ($m/z = 117$), $C_2H_6N_4O_2$ ($m/z = 118$), $C_3H_6N_4O_2$ ($m/z = 130$), $C_2H_7N_5O_3$ ($m/z = 149$), $C_2H_6N_4O_4$ ($m/z = 150$) and $C_3H_3N_5O_3$ ($m/z = 157$). We observed intermediates that belong to amines, mononitro, and mononitroso families as well as intermediate that carry both a nitro and nitroso groups. The underlying decomposition reactions involved in the formation of the observed species are proposed in next section.

3.4 Decomposition Mechanism

Here, we discuss the decomposition pathways of RDX based on the reaction mechanisms proposed through gas-phase and condensed phase calculations accounting for the aforementioned experimental findings. The majority of the gas-phase calculations predict that the decomposition

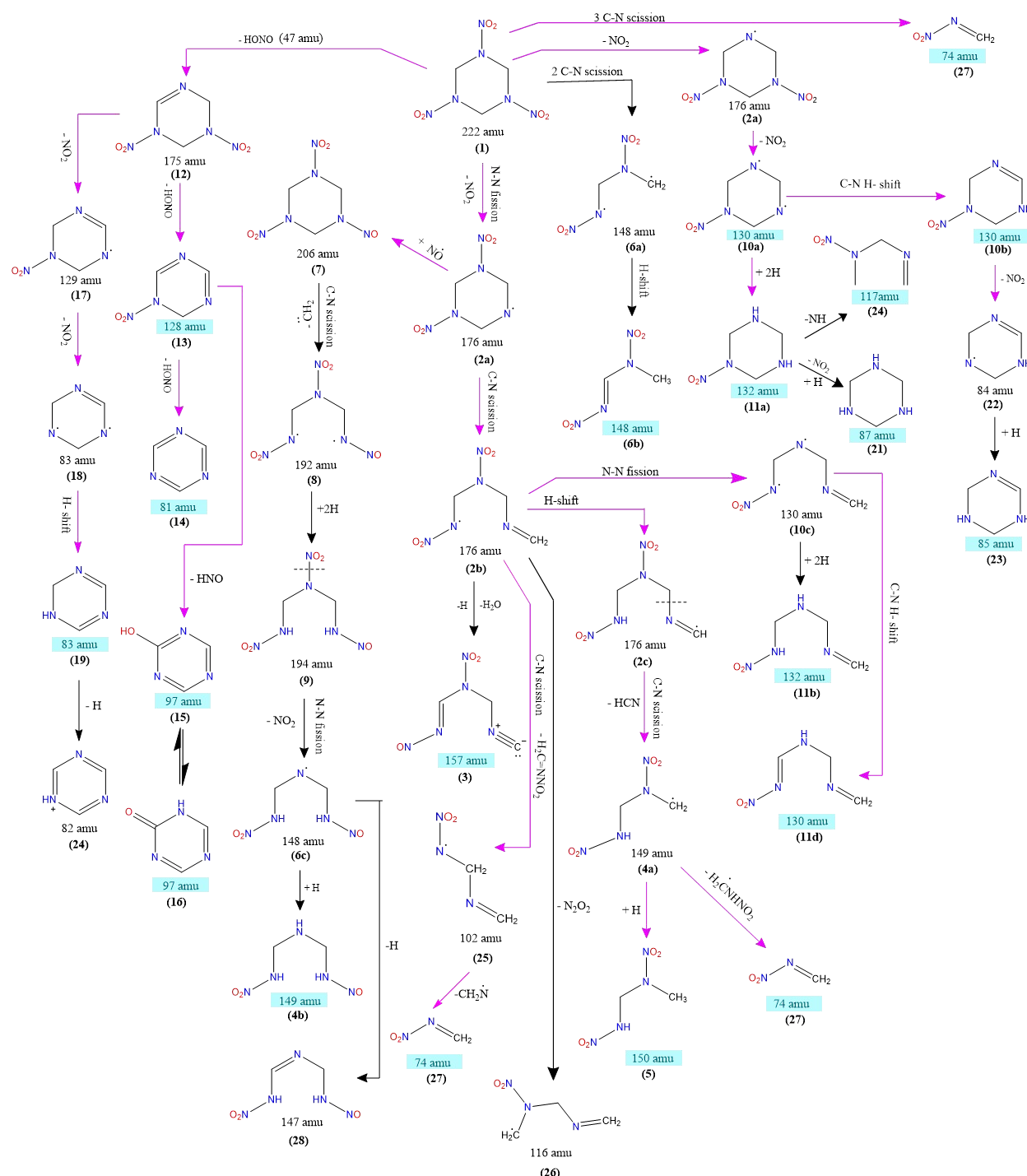


Figure 7a. Proposed decomposition mechanism of RDX into products at 157, 150, 149, 148, 132, 130, 128, 117, 97, 87, 85, 83 and 74 amu. Reactions mechanisms predicted by theoretical calculations are color coded in pink. 14, 16, 18, 22



These products can be broadly categorized into six species: dinitro, mononitro, mononitroso, nitro-nitroso, ring compounds, and smaller molecules.

Formation of dinitro species: At the early stage of RDX decomposition, the major products to be formed are dinitro species. Initially, the N-NO₂ fission of RDX (**1**) leads to a cyclic aminyl radical **2a** (C₃H₆N₅O₄•); this process requires an energy of 163 kJ mol⁻¹.¹⁴ Radical **2a** can undergo ring opening reaction via C-N scission to form an acyclic isomer **2b** (C₃H₆N₅O₄•) via a barrier of 110 kJ mol⁻¹.¹⁴ Intermediate **2b** (C₃H₆N₅O₄•) can further undergo a hydrogen-shift requiring 36 kJ mol⁻¹; this process forms yet another acyclic isomer with a carbon centered radical **2c** (C₃H₆N₅O₄•).¹⁴ The intermediate **2c** decomposes to generate the carbon-centered radical **4a** (C₂H₅N₄O₄•) along with a hydrogen cyanide (HCN) molecule via C-N β-scission. This reaction has a barrier of 68 kJ mol⁻¹.¹⁴ Addition of a hydrogen atom to **4a** results in a stable neutral intermediate **5** (C₂H₆N₄O₄; 150 amu), which has been observed in our experiment. Alternatively, radical **4a** may encounter C-N scission to form methylene nitramine **27** (CH₂NNO₂; 74 amu) and the CH₂NHNO₂• radical.¹⁴ Di-nitro intermediate **6b** (C₂H₄N₄O₄; 148 amu) can form from RDX via 2 C-N scissions followed by a hydrogen atom shift. As mentioned before, the decomposition of RDX could also be initiated by concerted HONO elimination. The first HONO elimination from RDX leads to di-nitro intermediate **12** (C₃H₅N₅O₄; 175 amu). The reaction barrier for HONO elimination (164 kJ mol⁻¹) is very similar to that of N-NO₂ fission.¹⁴

Formation of mono-nitro species: The di-nitro species formed during the early stage of decomposition can subsequently undergo C-N scission, N-N fission or hydrogen atom shift to form mono-nitro species. For example, RDX (**1**) undergoes first a N-NO₂ fission to the dinitro radical **2a** (C₃H₆N₅O₄•), which can subsequently undergo a second N-NO₂ fission to a cyclic biradical intermediate **10a** (C₃H₆N₃O^{••}). This radical **10a** undergoes hydrogen atom shift to form a cyclic mono-nitro **10b** (C₃H₆N₄O₂; 130 amu) product. This process has a barrier of 187 kJ mol⁻¹.¹⁴ Intermediate **10a** can also accept two hydrogen atoms yielding a cyclic product **11a** (C₃H₈N₄O₂; 132 amu). Both products (**10b** and **11a**) have been observed in our experiment. Product **11a** can further decompose to **24** (C₃H₇N₃O₂; 117 amu) via C-N scission and a hydrogen atom shift. Similarly, the acyclic aminyl radical **2b** (C₃H₆N₅O₄•) can form a biradical intermediate **10c** (C₃H₆N₄O₂••) via N-N fission which undergoes 1,2-H-shift to form an open chain isomer **10d** (C₃H₆N₄O₂; 130 amu) of cyclic product **10b**; this requires an energy of 190 kJ mol⁻¹.¹⁶ The

biradical intermediate **10c** can also accept two hydrogen radicals to form an acyclic isomer **11b** ($\text{C}_3\text{H}_8\text{N}_4\text{O}_2$; 132 amu) of product **11a**. The acyclic aminyl radical **2b** ($\text{C}_3\text{H}_6\text{N}_5\text{O}_4^\bullet$) can also experience a C-N scission, which requires 103 kJ mol^{-1} of energy to form a mono-nitro N-centered radical **25** ($\text{C}_2\text{H}_4\text{N}_3\text{O}_2^\bullet$) and methylene nitramine **27** (CH_2NNO_2 ; 74 amu).¹⁴ The radical **25** could engage in C-N scission to form a molecule of methylene nitramine (CH_2NNO_2 ; 74 amu) and H_2CN , this process requires an additional energy of 84 kJ mol^{-1} relative to that of radical **25** ($\text{C}_2\text{H}_4\text{N}_3\text{O}_2^\bullet$). Alternatively, the radical **25** can react further by either abstracting a hydrogen atom to form the mono-nitro product **35** ($\text{C}_2\text{H}_5\text{N}_3\text{O}_2$; 103 amu) or by eliminating a hydrogen atom to yield the product **34** ($\text{C}_2\text{H}_3\text{N}_3\text{O}_2$; 101 amu) (Figure 7b). Furthermore, the addition of three hydrogen atoms to radical **25** can generate the product **36** ($\text{C}_2\text{H}_7\text{N}_3\text{O}_2$; 105 amu; Figure 7b).¹⁴ Interestingly, all the four mono-nitro products (103 amu, 101 amu, 105 amu and 74 amu) have been observed in our experiment. The acyclic aminyl radical **2b** ($\text{C}_3\text{H}_6\text{N}_5\text{O}_4^\bullet$) can also decompose to a mono-nitro carbon centered radical **26** ($\text{C}_3\text{H}_6\text{N}_3\text{O}_2^\bullet$) via elimination of N_2O_2 (Figure 7a). The radical **26** ($\text{C}_3\text{H}_6\text{N}_3\text{O}_2^\bullet$) may involve in the formation of five membered ring species, which will be discussed later. The dinitro intermediate **12** ($\text{C}_3\text{H}_5\text{N}_5\text{O}_4$; 175 amu) formed via concerted HONO elimination from RDX can eliminate yet another HONO molecule to the mono-nitro product **13** ($\text{C}_3\text{H}_4\text{N}_3\text{O}_2$; 128 amu) (Figure 7a) with a barrier of 134 kJ mol^{-1} .¹⁴ Intermediate **13** undergoes decomposition to form various cyclic species which will be discussed later.

Formation of ring compounds: Species carrying the cyclic RDX moiety mainly originate from cyclic dinitro or mono-nitro compounds formed via HONO elimination or N- NO_2 fission(s). The most common cyclic products is triazine **14** ($\text{C}_3\text{H}_3\text{N}_3$; 81 amu), which is formed via HONO elimination from the mono-nitro intermediate **13** ($\text{C}_3\text{H}_4\text{N}_3\text{O}_2$; 128 amu). The calculated energy barrier for this reaction is 84 kJ mol^{-1} .¹⁴ Besides triazine, our experiments also suggest the formation of oxy-sym-triazine (OST) **15** ($\text{C}_3\text{H}_3\text{N}_3\text{O}$; 97 amu). Formation of this molecule has been proposed computationally from mononitro intermediate **13** ($\text{C}_3\text{H}_4\text{N}_3\text{O}_2$; 128 amu) after elimination of a HNO molecule.^{4, 14} Hydrogenated forms of the triazine ring such as $\text{C}_3\text{H}_5\text{N}_3$ (83 amu), $\text{C}_3\text{H}_7\text{N}_3$ (85 amu), and $\text{C}_3\text{H}_9\text{N}_3$ (87 amu) could be generated from mono-nitro intermediates via successive N- NO_2 fissions. For example, the di-nitro intermediate **12** ($\text{C}_3\text{H}_5\text{N}_5\text{O}_4$; 175 amu), which is formed via HONO elimination from RDX, could undergo two successive N- NO_2 fission to form a nitrogen-centered bi-radical **18** ($\text{C}_3\text{H}_5\text{N}_3^{\bullet\bullet}$). The energy required for this process is around 184 kJ mol^{-1} .¹⁶ The radical **18** can further undergo a hydrogen shift yielding **19**

($\text{C}_3\text{H}_5\text{N}_3$; 83 amu). Similarly, **23** ($\text{C}_3\text{H}_7\text{N}_3$; 85 amu) can originate from the mono-nitro species **10b** via N- NO_2 fission followed by a hydrogen atom addition. The fully hydrogenated form of triazine ($\text{C}_3\text{H}_9\text{N}_3$; 87 amu) can be formed from mono-nitro **11a** ($\text{C}_3\text{H}_8\text{N}_4\text{O}_2$; 132 amu) after a N- NO_2 fission followed by hydrogen addition. Five membered ring species such as $\text{C}_3\text{H}_8\text{N}_2$ (72 amu) and $\text{C}_3\text{H}_6\text{N}_2$ (70 amu) are also observed. These products could originate from a mono-nitro radical **26** ($\text{C}_3\text{H}_6\text{N}_3\text{O}_2^\bullet$) via ring closure mechanism (Figure 7b). Cyclization of radical **26** leads to a cyclic nitrogen-centered radical which can undergo hydrogen addition followed by N-N fission to form another nitrogen-centered radical **45a** ($\text{C}_3\text{H}_6\text{N}_2^\bullet$). This radical (**45**) can abstract a hydrogen to form product **46** ($\text{C}_3\text{H}_8\text{N}_2$; 72 amu) or it may release a hydrogen atom to form **45b** ($\text{C}_3\text{H}_6\text{N}_2$; 70 amu).

Formation of nitro-nitroso species: Products carrying both a nitro and nitroso groups have not been observed before with the exception of ONDNTA (7).⁴ Here, we few products that could originate from ONDNTA. It has been suggested that the ONDNTA is formed via recombination of cyclic aminyl radical **2a** ($\text{C}_3\text{H}_6\text{N}_5\text{O}_4^\bullet$) with nitrogen monoxide (NO) at the early stage of decomposition.^{5, 22} ONDNTA (7) eventually undergoes C-N scission to form an acyclic nitrogen-centered biradical **8** ($\text{C}_2\text{H}_4\text{N}_6\text{O}_5^{\bullet\bullet}$). Radical **8** can accept two hydrogens to form **9** ($\text{C}_2\text{H}_6\text{N}_6\text{O}_5$; 194 amu); the latter may undergo N-N fission leading to a nitrogen-centered radical **6c** ($\text{C}_2\text{H}_6\text{N}_5\text{O}_3^\bullet$). Addition of a hydrogen atom to radical **6c** give rise to an intermediate **4b** ($\text{C}_2\text{H}_7\text{N}_5\text{O}_3$; 149 amu). Alternately, a hydrogen atom can be abstracted from **6c** to produce intermediate **28** ($\text{C}_2\text{H}_5\text{N}_5\text{O}_3$; 147 amu). Intermediate **4b** can further dissociate to form the mono-nitro product **29** ($\text{C}_2\text{H}_6\text{N}_4\text{O}_2$; 118 amu) via elimination of a HNO molecule (Figure 7b). Similarly, intermediate **28** may eliminate a HNO molecule to form mononitro product **30** ($\text{C}_2\text{H}_4\text{N}_4\text{O}_2$; 116 amu; Figure 7b). Intermediate **4b** ($\text{C}_2\text{H}_7\text{N}_5\text{O}_3$; 149 amu) can also lead to a nitrogen-centered radical **37** ($\text{CH}_4\text{N}_3\text{O}_2^\bullet$) via elimination of a nitrosoamine radical ($\text{CH}_2\text{NHNO}^\bullet$). The reactive radical **37** eventually either abstracts a hydrogen atom to form **39** ($\text{CH}_5\text{N}_3\text{O}_2$; 91 amu) or releases a hydrogen atom yielding **38** ($\text{CH}_3\text{N}_3\text{O}_2$; 89 amu). The nitro-nitroso intermediates such as **3** ($\text{C}_3\text{H}_3\text{N}_5\text{O}_3$; 157 amu) can be generated from a cyclic aminyl radical **2b** ($\text{C}_3\text{H}_6\text{N}_5\text{O}_4^\bullet$) after eliminating water and a hydrogen atom (Figure 7a).

Formation of mono-nitroso species: Products carrying both nitro and nitroso groups could represent the precursor for mono-nitroso species. For example, mono-nitroso product **32a**

($\text{C}_2\text{H}_4\text{N}_4\text{O}$; 100 amu) may form from intermediate **28** ($\text{C}_2\text{H}_6\text{N}_5\text{O}_3$; 147 amu) after elimination of nitrogen dioxide followed by hydrogen atom shift (Figure 7b). Hydrogen abstraction from this molecule **32a** could lead to product **33** ($\text{C}_2\text{H}_2\text{N}_4\text{O}$; 98 amu).

Formation of amines: Mono-nitro and nitro-nitroso species can decompose into amines via N- NO_2 fission as well as C-N scission mechanisms (Figure 7b). For example, mono-nitro product **35** ($\text{C}_2\text{H}_5\text{N}_3\text{O}_2$; 103 amu) can undergo N- NO_2 fission to form a nitrogen-centered amine radical **48** ($\text{C}_2\text{H}_5\text{N}_2^\bullet$), which can accept a hydrogen atom to generate an amine product **49a** ($\text{C}_2\text{H}_6\text{N}_2$; 58 amu). The barrier to this process is about 189 kJ mol^{-1} . Likewise, intermediate **4b** can decompose into diamine radical **40** via two successive N-N fissions; this radical can then undergo a hydrogen-shift to generate **42** ($\text{C}_2\text{H}_7\text{N}_3$; 73 amu). Alternatively, radical **40** may accept two hydrogen atoms to form the diamine **41** ($\text{C}_2\text{H}_9\text{N}_3$; 75 amu). Species **42** can further release two hydrogen atoms to produce **43** ($\text{C}_2\text{H}_5\text{N}_3$; 71 amu). Mono-nitro **36** can decompose into a molecule of methylene nitramine plus methylamine (CH_3NH_2 ; 31 amu; **56**) via C-N scission. Ring compounds such as **21** ($\text{C}_3\text{H}_9\text{N}_3$; 87 amu) can also decompose to generate a carbon-centered biradical **44a** ($\text{C}_2\text{H}_5\text{N}^{\bullet\bullet}$). This radical **44a** can either undergo ring closure to **44b** ($\text{c-C}_2\text{H}_5\text{N}$; 43 amu) or undergoes a hydrogen-shift to **44c** ($\text{C}_2\text{H}_5\text{N}$; 43 amu). Dinitro species **5** can undergo two successive HONO elimination to generate **47** ($\text{C}_2\text{H}_4\text{N}_2$; 56 amu).

Formation of small molecules: Finally, methylene nitramine **27** (CH_2NNO_2 ; 74 amu) formed from higher molecular weight species decomposes into secondary products such as dinitrogen oxide (N_2O), hydrogen cyanide (HCN), nitrogen monoxide (NO; 30 amu; **53**), formaldehyde (H_2CO ; 30 amu; **52**), and methylene nitrosoamine (CH_2NNO ; 58 amu; **49b**). Several pathways for the decomposition of methylene nitramine exist. i) The N- NO_2 fission of **27** (CH_2NNO_2 ; 74 amu) results in formation of NO_2 (46 amu; **51**) and $\text{CH}_2\text{N}^\bullet$ (28 amu; **50**) radical. ii) Methylene nitramine **27** (CH_2NNO_2 ; 74 amu) can also undergo HONO elimination to form hydrogen cyanide (HCN) (27 amu; **52**). iii) It can also decompose into dinitrogen oxide (N_2O) and formaldehyde (H_2CO ; 30 amu; **57**). iv) Another possibility is the N-O fission, which results in formation of methylene nitrosamine (CH_2NNO ; 58 amu; **49b**). Product **49b** can undergo hydrogen addition to form methyl nitrosamine (CH_3NHNO ; 60 amu; **55**). The $\text{CH}_2\text{N}^\bullet$ radical **50** formed from methylene nitramine via N- NO_2 fission can combine with $^\bullet\text{OH}$ radical to form

formaldoxime (CH_2NOH ; 45 amu; **54a**), which can isomerize to nitroso methane (CH_3NO ; 45 amu; **54b**).

4. Conclusions

In the present study, the decomposition of RDX by energetic electrons was explored using a surface science machine at 5 K. The decomposition products were probed exploiting FTIR, UV-VIS, and Re-TOF-PI techniques. Our FTIR data revealed the formation of simple decomposition products such as water (H_2O), carbon dioxide (CO_2), dinitrogen oxide (N_2O), hydrogen cyanide (HCN), nitrogen monoxide (NO), formaldehyde (H_2CO), nitrous acid (HONO), and nitrogen dioxide (NO_2). 39 species were detected through ReTOF-PI mass spectroscopy at m/z values from 39 to 157. Products observed at m/z values of 31, 60, 71, 72, 75, 85, 87, 89, 91, 98, 100, 101, 103, 105, 116, 117, 118, 130, 149, 150, and 157 are detected for the first time. The new products observed in the present study belong to six classes. These are i) dinitro compounds: $\text{C}_2\text{H}_6\text{N}_4\text{O}_4$ ($m/z=150$), ii) mono-nitro compounds: $\text{CH}_3\text{N}_3\text{O}_2$ ($m/z=89$), $\text{CH}_5\text{N}_3\text{O}_2$ ($m/z=91$), $\text{C}_2\text{H}_3\text{N}_3\text{O}_2$ ($m/z=101$), $\text{C}_2\text{H}_5\text{N}_3\text{O}_2$ ($m/z=103$), $\text{C}_2\text{H}_7\text{N}_3\text{O}_2$ ($m/z=105$), $\text{C}_2\text{H}_6\text{N}_4\text{O}_2$ ($m/z=118$), $\text{C}_3\text{H}_6\text{N}_4\text{O}_2$ ($m/z=130$); iii) mono-nitroso compounds: $\text{CH}_4\text{N}_2\text{O}$ ($m/z=60$), $\text{C}_2\text{H}_2\text{N}_4\text{O}$ ($m/z=98$), $\text{C}_2\text{H}_4\text{N}_4\text{O}$ ($m/z=100$); iv) nitro-nitroso compounds: $\text{C}_3\text{H}_3\text{N}_5\text{O}_3$ ($m/z=157$), $\text{C}_2\text{H}_7\text{N}_5\text{O}_3$ ($m/z=149$), v) cyclic compounds: $\text{C}_3\text{H}_8\text{N}_2$ ($m/z=72$), $\text{C}_2\text{H}_9\text{N}_3$ ($m/z=75$), $\text{C}_3\text{H}_7\text{N}_3$ ($m/z=85$), $\text{C}_3\text{H}_9\text{N}_3$ ($m/z=87$), and VI) amines: $\text{C}_2\text{H}_5\text{N}_3$ ($m/z=71$), $\text{C}_2\text{H}_9\text{N}_3$ ($m/z=75$). We also inferred reactive radical intermediates such as $c\text{-C}_3\text{H}_6\text{N}_4\text{O}_2^\bullet$, $\text{C}_3\text{H}_6\text{N}_4\text{O}_2^\bullet$, $\text{C}_2\text{H}_5\text{N}_4\text{O}_4^\bullet$, $\text{C}_2\text{H}_4\text{N}_3\text{O}_2^\bullet$, $c\text{-C}_3\text{H}_5\text{N}_3^\bullet$ and $\text{C}_2\text{H}_5\text{N}_2^\bullet$; previously these radical species have only been predicted through electronic calculations. Finally, the decomposition mechanisms leading to the observed products have been discussed. Overall, the decomposition of RDX is most likely initiated via N-NO_2 fission that leads to the formation of a cyclic aminyl radical, which can further undergo C-N scission, N-NO_2 fission, or hydrogen-shift to form dinitro and mono-nitro intermediates. Alternatively, RDX decomposes through HONO elimination to form dinitro, mono-nitro and triazine-like cyclic compounds. Dinitro and mononitro products could further experience C-N scission or N-NO_2 fission to form methylene nitramine as well as amines. Methylene nitramine further fragment into secondary products such as dinitrogen oxide (N_2O), hydrogen cyanide (HCN), nitrogen monoxide (NO), formaldehyde (H_2CO), and nitrogen dioxide (NO_2). The mono-nitroso and nitro-nitroso species most likely originates from ONDNTA. The latter could form at an early stage of the decomposition via

radical recombination of cyclic aminyl radical and nitrogen monoxide (NO). Mono-nitroso and nitro-nitroso species were detected, as well.

An investigation of the decomposition of RDX as conducted in the present experiments is very challenging and insights can be provided by merging the experimental results with a computational investigation of decomposition pathways. The present study represents the very first step in an investigation of the decomposition by ionizing radiation. Further experiments should exploit tunable vacuum ultraviolet light to explicitly identify the nature of the decomposition products. This requires a computational determination of the ionization energies of all structural isomers. Finally, additional insights will be obtained by selectively photo exciting RDX under the same experimental conditions of 5 K via distinct transitions with the goal to selectively cleave – at least initially – N-NO₂, C-N, N-O, and C-H bonds.

Associated content

Supporting Information. NMR spectra, IR spectra.

This material is available free of charge via the Internet at <http://pubs.acs.org>.

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Conflict of interest

There are no conflicts to declare.

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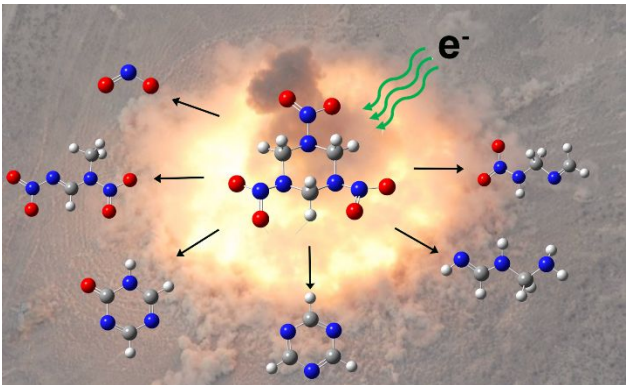
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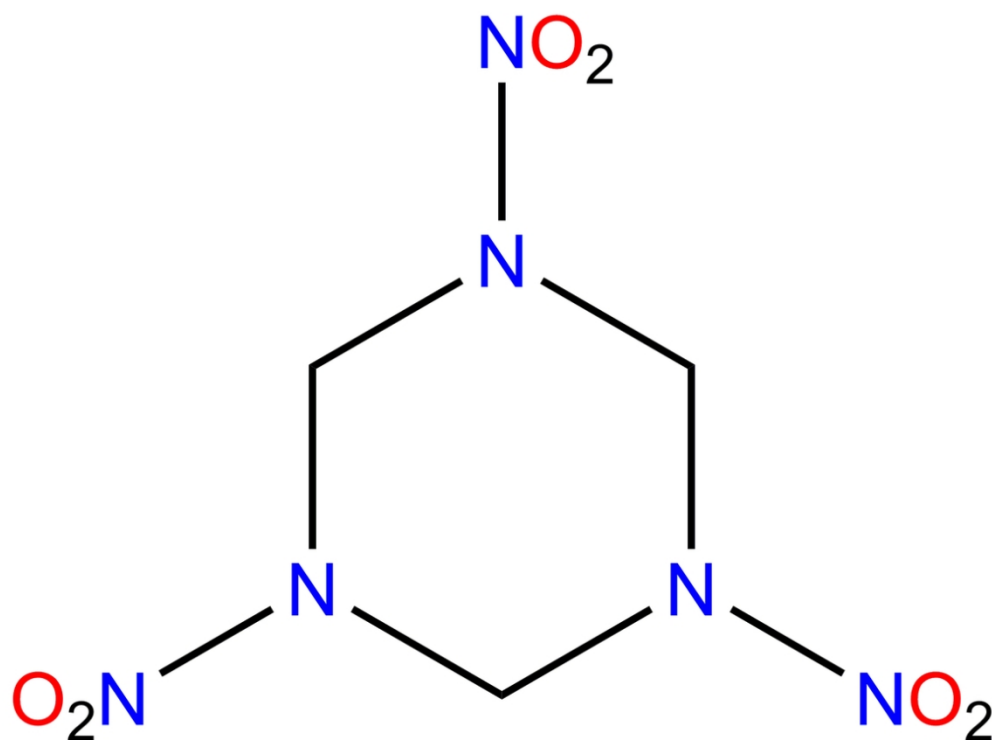
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Table of Content





Scheme 1. Molecular structure of 1,3,5-trinitro-1,3,5-triazinane (RDX).

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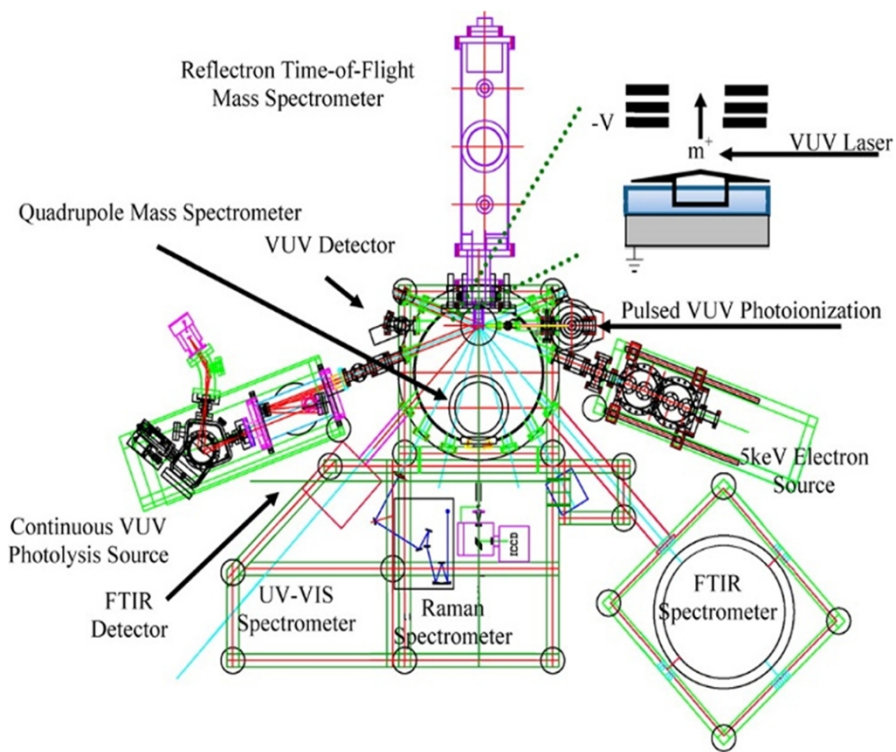


Figure 1. Schematic top view of the ultra-high vacuum chamber including the radiation sources (electron source), analytical instruments (FTIR, UV-VIS, ReTOF), and cryogenic target (point of convergence lines).⁴⁶⁻⁴⁸

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Figure 2. Infrared spectra of RDX collected at 5 K (a) before and (b) after the irradiation of the high dose experiment.

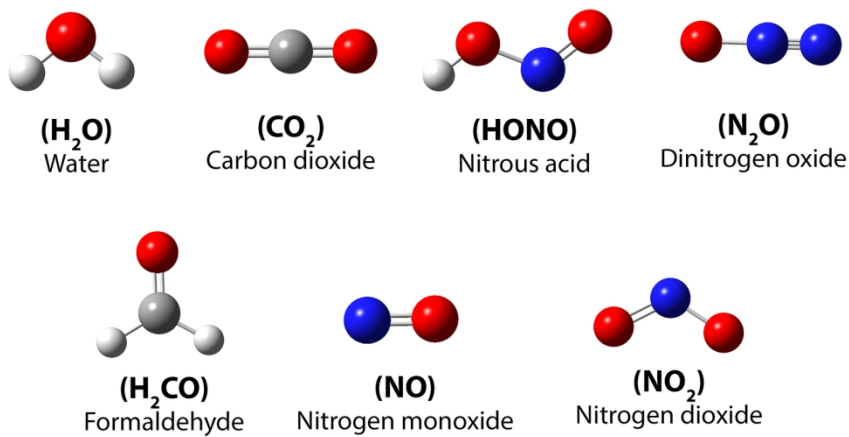


Figure 3. Structures of the decomposition products detected through FTIR spectroscopy.
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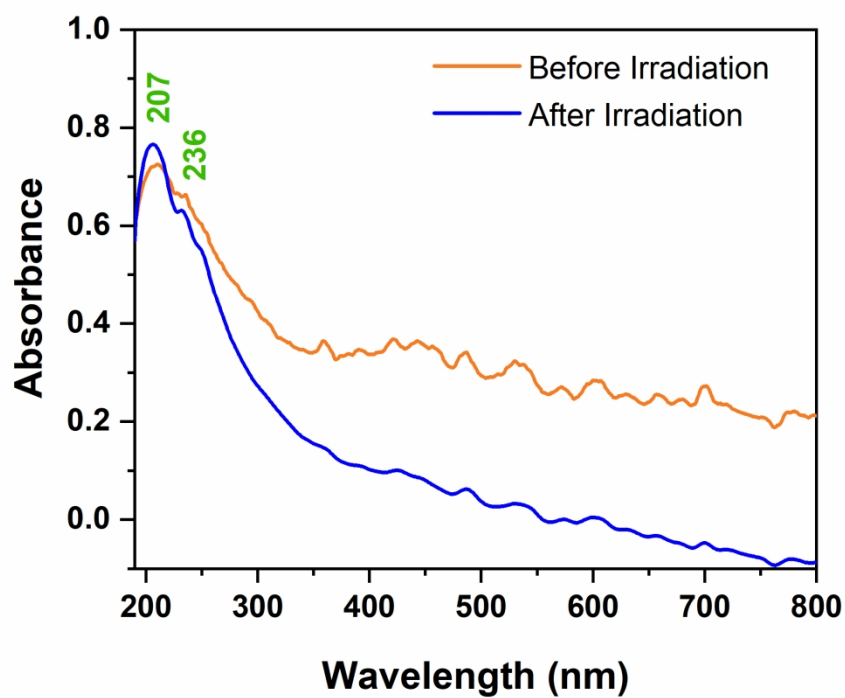


Figure 4. UV-VIS spectra of RDX collected at 5 K before and after the high dose.

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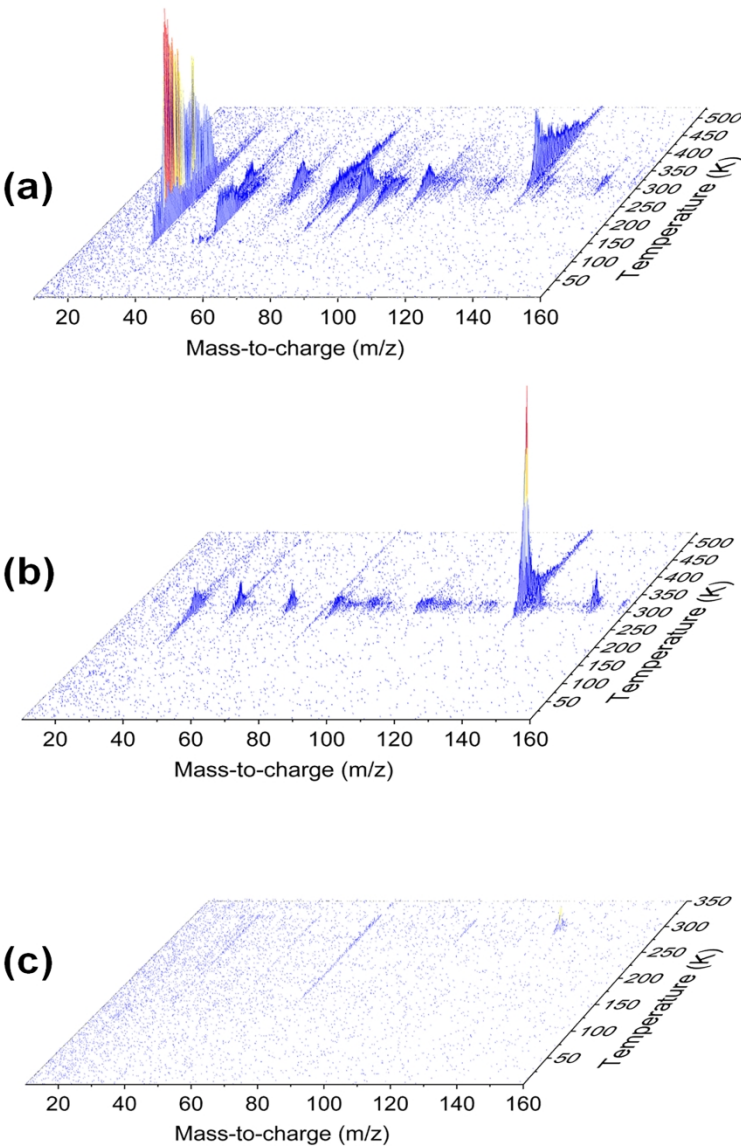


Figure 5. Temperature dependent ReTOF data of the decomposition products of RDX for the high dose (a), low dose (b), and blank (c) experiments.

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Figure 6. TPD profiles of the ion counts recorded at mass-to-charge ratios of (a) 30, 31, 42, 43, 45, 46, (b) 56, 58, 60, 70, 71, 72, (c) 73, 74, 75, 81, 82, 83, and (d) 85, 87, 89, 91, 97 and 98 at a photoionization energy of 10.49 eV. Blue: high doses; green: low dose; red: blank.

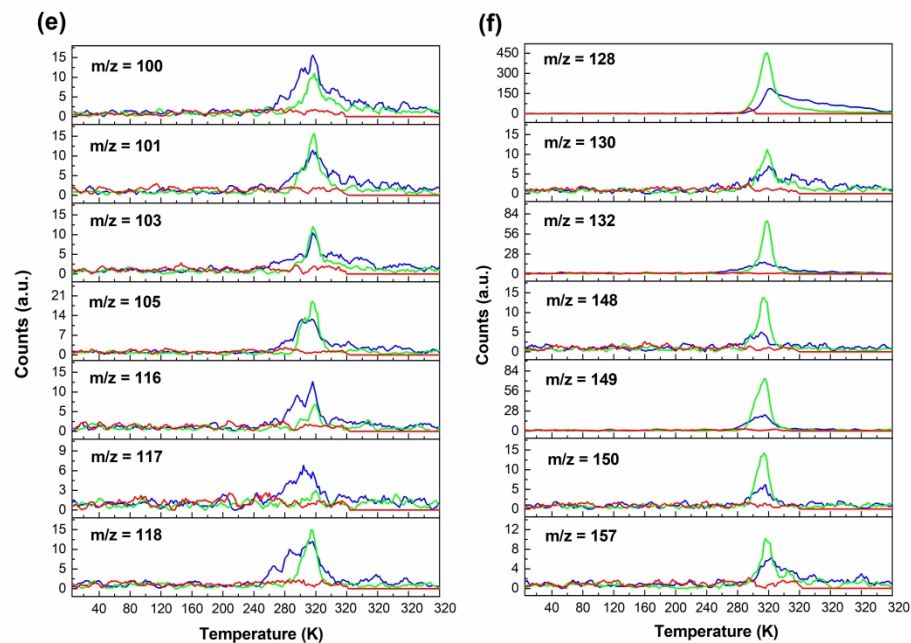


Figure 6. TPD profiles of the ion counts recorded at mass-to-charge ratios of (e) 100, 101, 103, 105, 116, 117, 118, (f) 128, 130, 132, 148, 149, 150 and 157 at a photoionization energy of 10.49 eV. Blue: high dose; green: low dose; red: blank.

518x355mm (300 x 300 DPI)

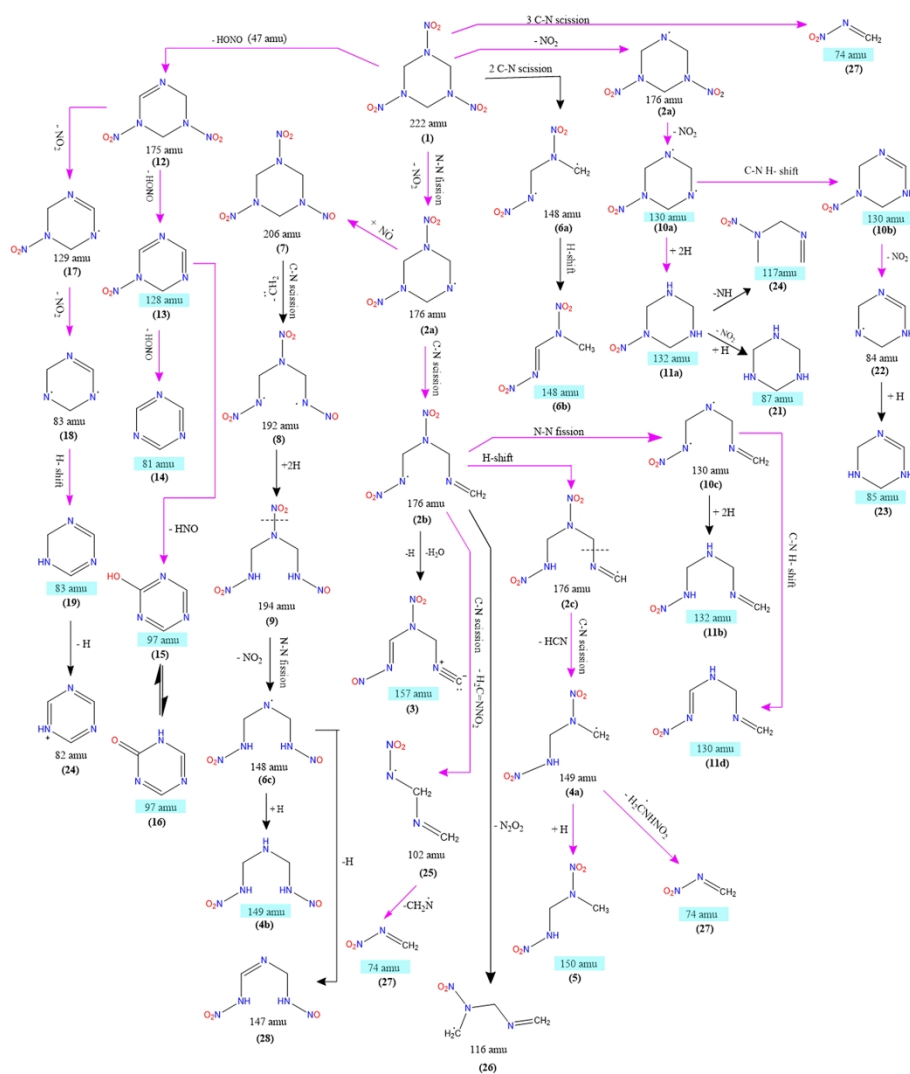


Figure 7a. Proposed decomposition mechanism of RDX into products at 157, 150, 149, 148, 132, 130, 128, 117, 97, 87, 85, 83 and 74 amu. Reactions mechanisms predicted by theoretical calculations are color coded in pink. 14, 16, 18, 22

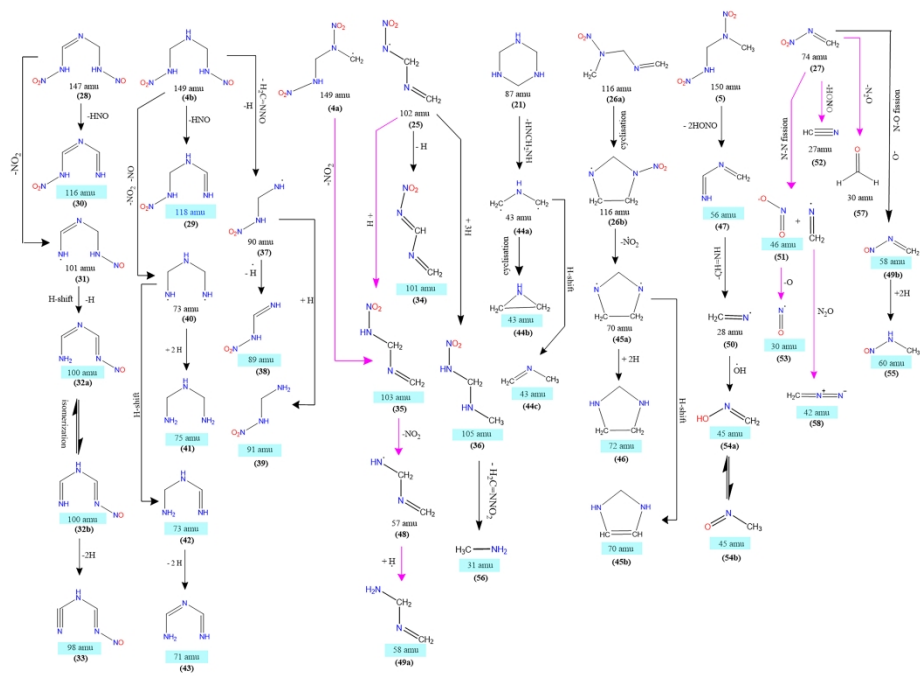


Figure 7b. Proposed decomposition mechanism of RDX into products at 118, 116, 105, 103, 101, 100, 98, 91, 89, 75, 73, 72, 71, 70, 60, 58, 56, 46, 45, 43, 31, and 30 amu. Reactions mechanisms predicted by theoretical calculations are color coded in pink.14, 16, 18

508x361mm (300 x 300 DPI)