The elusive cyclotriphosphazene molecule and its Dewar benzene–type valence isomer (P$_3$N$_3$)

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Although the chemistry of phosphorus and nitrogen has fascinated chemists for more than 350 years, the Hückel aromatic cyclotriphosphazene (P$_3$N$_3$, 2) molecule—a key molecular building block in phosphorus chemistry—has remained elusive. Here, we report a facile, versatile pathway producing cyclotriphosphazene and its Dewar benzene–type isomer (P$_3$N$_3$, 5) in ammonia-phosphine ices at 5 K exposed to ionizing radiation. Both isomers were detected in the gas phase upon sublimation via photoionization reflectron time-of-flight mass spectrometry and discriminated via isomer-selective photochemistry. Our findings provide a fundamental framework to explore the preparation of inorganic, isovalent species of benzene (C$_6$H$_6$) by formally replacing the C–H moieties alternatingly through phosphorus and nitrogen atoms, thus advancing our perception of the chemical bonding of phosphorus systems.

INTRODUCTION

Ever since the pioneering discovery of the p-block element phosphorus by the alchemist Hennig Brand 350 years ago (1), phosphorus allotropes such as white, red, and black phosphorus along with phosphorus heterocycles such as the diphenphiphiazoiate anion (P$_2$N$_3^-$, 1) (2) and cyclotriphosphazene (P$_3$N$_3$, 2; Fig. 1) (3, 4) have captivated the interest of the theoretical, physical (in)organic, preparative, and organometallic chemistry communities (1) from the fundamental viewpoints of electronic structure (5, 6) and chemical bonding (7, 8), with both heterocycles representing benchmarks of a class of exotic inorganic (4n + 2)π aromatic molecules. Whereas 1 is isovalent to the cyclopentadienyl anion (3), 2 can be linked to the prototype 6π-Hückel aromatic (9) benzene molecule (C$_6$H$_6$, 4) by formally replacing the C–H moieties alternatingly by isovalent phosphorus (P) and nitrogen (N) atoms. This leads to a planar, D$_6$h symmetric 2 molecule compared to a D$_6$h point group of benzene (C$_6$H$_6$, 4), with both molecules holding 1A$_1$ electronic ground states (5, 10).

Cyclotriphosphazene (2) has received particular attention as the prototype of a hydrogen-deficient, inorganic Hückel aromatic system. The distinct Pauling electronegativities of nitrogen (3.0) and phosphorus (2.2) result in a less efficient delocalization of the π electrons and hence diminished aromaticity compared to benzene (C$_6$H$_6$, 4). This is supported by nucleus-independent chemical shifts (NICSs) of −3.9 parts per million (ppm) (h = 1.0 Å) and −10.2 ppm (h = 1.0 Å) for 2 and benzene (C$_6$H$_6$, 4), respectively (6, 7), along with the charge population analysis suggesting positive and negative partial charges on the phosphorus and nitrogen atoms, respectively (5). Similarities in the isovalent 4–2 system are evident from the molecular structures of the thermodynamically less stable bicycle [2.2.0]hexa-2,5-diene (“Dewar benzene”) (6; +354 kJ mol$^{-1}$) (11) and 1,3,5-triphosphas-2,4,6-triazabicyclo [2.2.0]hexa-2,5-diene (5; +155 kJ mol$^{-1}$) valence isomers (Fig. 1), with the latter only predicted theoretically to exist (8). Although 2 core-based dendrimers (12) and bulky moieties such as diphenyl (13) are stable at room temperature, the parent compound 2 has not been undeniably prepared. Matrix isolation studies by Atkins and Timms (3) at 10 K tentatively assigned a cyclic [phosphorusmononitride (PN)$_3$] isomer via infrared spectroscopy through absorptions at 1137 and 718 cm$^{-1}$, with Schnöckel and co-workers (4) validating the aforementioned analysis. Therefore, despite tentative evidence for the existence of 2, the preparation and isolation of 2 in the gas phase have eluded synthetic chemists. Considering the difficulties in preparation, short lifetimes above 50 K, and the tendency for polymerization, free cyclotriphosphazenes are one of the least explored classes of inorganic molecules.

RESULTS

Here, we report on the first preparation of 2 along with its Dewar benzene–type valence isomer 5 by exposing low-temperature (5 K) ammonia (NH$_3$) + phosphine (PH$_3$) ices to ionizing radiation in the form of energetic electrons. By combining our experiments with electronic structure computations on the P$_3$N$_3$ potential energy surface, both isomers were unambiguously identified via photoionization reflectron time-of-flight mass spectrometry (PI-ReTOF-MS) during the temperature-programmed desorption (TPD) phase of the irradiated ices based on their adiabatic ionization energies (IEs), mass shifts upon $^{15}$N substitution, and their distinct photoelectron spectra at λ = 452 nm and λ = 388 nm, respectively. By identifying 2 and 5, this study provides a deeper understanding of the molecular structures and (photo)reactivities of previously obscure heteroaromatic, inorganic molecules (P$_3$N$_3$) along with their valence isomers, thus advancing our perception how we think about chemical bonding of phosphorus in aromatic compound.

Fourier transform infrared spectroscopy

The ammonia-phosphine ices were monitored during the irradiation at 5 K via Fourier transform infrared (FTIR), which excels in identifying
small, individual molecules and functional groups of complex molecules (fig. S1 and table S2). Radiation exposure produced seven new absorptions: 3039 cm\(^{-1}\) (\(v_{\text{N} \rightarrow \text{H}}\)), 2782 cm\(^{-1}\) [\(v_{\text{N}=\text{NH}_2}\) (\(v_5\))], 2235 cm\(^{-1}\) (\(v_{\text{P} \rightarrow \text{H}}\) (14)), 2085 cm\(^{-1}\) (\(v_{\text{N} \equiv \text{N}}\)) [\(v_{\text{N} \equiv \text{N}}\)], 1506 cm\(^{-1}\) [\(v_{\text{NH}_2}\) (\(v_2\))], 1153 cm\(^{-1}\) [\(v_{\text{P} \equiv \text{N}}\)], and 720 cm\(^{-1}\) [\(\text{NH}_2\) wagging] (15–18). These assignments were upheld by performing experiments with \(^{15}\text{N}\)-labeled ice precursors (table S2), which induced red shifts for the nitrogen-bearing functional groups. The \(\text{P}=\text{N}\) stretching at 1153 cm\(^{-1}\) could be associated to multiple \(\text{P}_3\text{N}_3\) isomers carrying the \(\text{PN}\) moiety (table S3). However, infrared spectroscopy alone cannot identify individual \(\text{P}_3\text{N}_3\) isomers as their absorptions fall in a similar range. Hence, an alternative analytical technique is necessary to probe discrete isomers selectively.

**Photoionization reflectron time-of-flight mass spectrometry**

Therefore, we exploited PI-ReTOF-MS during the TPD phase of the irradiated ices to 300 K (19). This method represents a unique approach to detecting gas-phase molecules isomer-selectively via soft photoionization based on their distinct IEs by systematically tuning the photon energies above and below the IE of the isomer(s) to be identified. This results in the identification of the parent ions at well-defined mass/charge ratio (\(m/z\)). Considering the computed IEs of the \(\text{P}_3\text{N}_3\) isomers (Fig. 2 and table S3), a photoionization energy (PE) of 10.49 eV (\(\lambda = 118.22\) nm) was first chosen to ionize all isomers except 9 (IE = 11.10 eV) and 10 (IE = 10.54 eV); thereafter, we selected a photon energy of 9.10 eV (\(\lambda = 136.25\) nm), which can only ionize isomers having IEs less than 9.10 eV (7, 8, 11, 12, 13, and 14) but not the target isomers 2 (IE = 9.28 eV) and 5 (IE = 9.32 eV) (Fig. 3). By comparing the TPD profiles at \(m/z = 135\) (\(\text{P}_3\text{N}_3^+\)) at 10.49 and 9.10 eV, isomers 2 and/or 5 could be identified. The mass spectra of the subliming molecules are compiled in detail in Fig. 3 as a function of temperature. Focusing on the \(\text{P}_3\text{N}_3\) isomers, we extracted the TPD profiles of ions at \(m/z = 135\) (\(\text{P}_3\text{N}_3^+\)) (Fig. 4). At 10.49 eV, three sublimation peaks centered at 215, 237, and 256 K are observed (Fig. 4A). Upon tuning to 9.10 eV, below the IEs of 2 and 5, the first sublimation event is still present, while the latter two peaks vanish (Fig. 4B). These findings suggest that the first sublimation events can be linked to 7, 8, 11, 12, 13, and/or 14, whereas the two latter ones correlate with 2 and/or 5. In separate experiments exploiting \(^{15}\text{N}\)-substituted ammonia (\(^{15}\text{NH}_3\)–\(\text{PH}_3\) system) at 10.49 eV, signal shifted from \(m/z = 135\) by 3 atomic mass unit to \(m/z = 138\) (\(\text{P}_3\text{N}_3^+\)) (fig. S2). The corresponding TPD profile also reveals a triplet with maxima at 216, 238, and 259 K; this finding mirrors the sublimation events at \(m/z = 135\) in nonspectroscopically labeled experiments confirming the aforementioned assignments. In blank studies, experiments were carried out under identical conditions, but without electron processing of the ices, no ion counts at these \(m/z\) were detected at all (Fig. 4), demonstrating that the identified species are due to the irradiation of the ices but not from ion–molecule reactions in the gas phase. Therefore provided compelling evidence for the detection of \(\text{P}_3\text{N}_3\) isomers 2 and/or 5. Since their IEs are of 9.28 and 9.32 eV and are too close to each other, additional investigations are required to discriminate which isomer can be assigned to these sublimation events.

To discriminate between 2 and 5, we conducted isomer-selective ultraviolet-visible (UV-vis) photolysis experiments (20, 21). Here, the ices were first processed by energetic electrons to produce the \(\text{P}_3\text{N}_3\) isomers and then to photolyze selectively to degrade and/or isomerize these isomers. This requires the knowledge of the UV-vis absorption spectrum of 2 and 5. Time-dependent density functional theory (DFT) computations reveal that absorption bands at \(\lambda = 452\) nm [highest occupied molecular orbital (HOMO) (E\(^{-}\)) → lowest unoccupied molecular orbital (LUMO) (E\(^{+}\))] and \(\lambda = 388\) nm [HOMO (A\(^{-}\)) → LUMO +1 (A\(^{+}\)) and HOMO −1 (A\(^{-}\)) → LUMO +1 (A\(^{+}\))] are exclusive to 2 and 5, respectively (table S4 and fig. S3). Therefore, photolysis at \(\lambda = 452\) nm and \(\lambda = 388\) nm for 1 hour with 30 mW selectively isomerizes and/or photolyzes isomers 2 and 5, respectively. A comparison of the TPD graph at \(m/z = 135\) of the electron irradiation system (Fig. 4A) with the photolyzed sample (452 nm; absorption band of 2; Fig. 4C) reveals that the 256-K sublimation event vanishes; the 215- and 237-K peaks are still present. On the other hand, 388-nm photolysis (absorption of 5) leads to the disappearance of the 237-K sublimation event, but the peaks at 215
and 256 K still remain (Fig. 4D). The phenomena are not due to laser-induced thermal decomposition, as the temperature of the sample was kept at 5 K during the photolysis. It is highly unlikely that the two experiments could generate new species, which would selectively react with the carriers of the 237- and 256-K peaks. Therefore, our results demonstrate that the 237- and 256-K peaks can be linked to two distinct P3N3 isomers 5 and 2, respectively. Note that the 452-nm irradiation also results in a new peak at 200 K, which can be assigned to P3N3 isomers holding IEs less than 10.49 eV [7 (IE = 7.58 eV), 8 (IE = 7.38 eV), 11 (IE = 8.94 eV), 12 (IE = 7.76 eV), 13 (IE = 8.08 eV), or 14 (IE = 8.64 eV), except 5 as it sublimes at 256 K]. These new species may be generated by isomerization of 2 or reactions of other molecules within the ice induced by blue light irradiation. No new peak is observed in the TPD profile of m/z = 135 recorded in the 388-nm photolysis, suggesting that 5 either rearranges to isomers having IEs higher than 10.49 eV [9 (IE = 11.10 eV) and 10 (IE = 10.54 eV)] or fragments to smaller molecules, e.g., PN.

**Structure and aromaticity**

Having established the formation of the P3N3 isomers 2 and 5, we are shifting our attention now to their electronic and geometric structures. For isomer 5, the length of both P=N bonds is 1.62 Å at the CCSD(T)/cc-pVTZ level of theory, while the remaining P–N single bonds are 0.11 to 0.25 Å longer (table S3); this is similar to the C–C bond lengths in Dewar benzene (22). In 2, all six P–N bonds have a length of 1.64 Å (table S3); this bond length is shorter than P–N bonds in saturated phosphazanes (1.77 Å) (23), which indicates partial double bond character. Conjugation-induced shortening of the P–N bond from saturated species (−0.13 Å) is similar to that of the C–C bond of isovalent benzene (C6H6) (−0.14 Å). The aromatic character of 2 can be evaluated from NICS values (vide supra) and homodesmotic equations (Fig. 5). We computed the NICS(0) (h = 0.0 Å) and NICS(1) (h = 1.0 Å) values for benzene (C6H6) (4), borazine (B3N3H6) (15), and 2 (table S5). The NICS(0) values for benzene (C6H6) (4) and borazine (B3N3H6) (15) are negative (−8.2 and −1.7 ppm), while for 2, a positive value ensues (+2.8 ppm). All computed NICS(1) values are negative (−10.4, −3.0, and −3.5 ppm), indicating the deshielding and \(\pi\)-aromatic character (to view the \(\pi\)
orbits of 2, see fig. S4) for all compounds. NICS values for 2 become negative at \( h = 0.5 \) Å and reach a maximum at \( h = 1.2 \) Å (−3.8 ppm), which is still remarkably lower than in benzene (\( C_6H_6 \), 4) (maximum at \( h = 0.8 \) Å, −10.7 ppm; table S5). An explanation might be the larger atomic radii of phosphorus in comparison to carbon. According to the NICS aromaticity criteria, 2 would be almost three times less aromatic than 4 and shows similar aromaticity as borazine (\( B_3N_3H_6 \), 15). We computed the aromatic stabilizations of 2, benzene (\( C_6H_6 \), 4), and borazine (\( B_3N_3H_6 \), 15). With the help of the homodesmotic equations (24) depicted in Fig. 5, cyclohexene (16) reacts in a hypothetical reaction with cyclohexadiene (17) to form cyclohexane (18) and benzene (\( C_6H_6 \), 4). The reaction enthalpy \( \Delta H^0_r \) equals the aromatic and conjugative stabilization of benzene (\( C_6H_6 \), 4) and is computed to be \( \Delta H^0_r = -151 \) kJ mol\(^{-1}\). The reaction enthalpies for borazine (\( B_3N_3H_6 \), 15) and 2 are computed with similar equations (Fig. 5, II and III) to be \( \Delta H^0_r = -233 \) kJ mol\(^{-1}\) and \( \Delta H^0_r = -45 \) kJ mol\(^{-1}\). Benzene is almost three times more stable than 2, in line with our NICS(1) computations. However, borazine (\( B_3N_3H_6 \), 15) seems to be the most stable one. These numbers might explain why benzene (\( C_6H_6 \), 4) and borazine (\( B_3N_3H_6 \), 15) are easy to synthesize and bench stable, while the formation and isolation of 2 are highly challenging.

DISCUSSION

We prepared the highly reactive cyclotriphosphazene (\( P_3N_3 \), 2) along with its Dewar benzene–type isomer 1,3,5-triphasa-2,4,6-triazabicyclo[2.2.0]hexa-2,5-diene (\( P_3N_3 \), 5) in ammonia-phosphine ices exposed to energetic electrons and also explored their photoreactions at \( \lambda = 452 \) nm and \( \lambda = 388 \) nm. Both isomers were detected in the gas phase upon sublimation of the newly formed molecules exploiting tunable, single PI-ReTOF-MS. Considering the average velocity of \( 200 \) m s\(^{-1}\) of 2 and 5 subliming in the 230- to 300-K range and the 2-mm distance between the ice and the photoionization laser, the lifetimes of both neutral isomers in the gas phase have to be at least 8 \( \mu \)s to survive the flight time from sublimation and photoionization. Structure 2 is the parent to a broad variety of phosphacyclic compounds that have been extensively applied in various areas, such as biomedicine, nanotechnology, and catalysis (12, 13, 25, 26). This work

![Diagram](http://advances.sciencemag.org/22/2020/)

Fig. 5. Homodesmotic equations (I to III) for the determination of the aromatic stabilization of benzene (4), borazine (15), and cyclotriphosphazene (2) and computed NICS values (given in parts per million, ppm) at the B3LYP/cc-pVTZ level of theory.
represents a versatile benchmark to prepare and identify highly reactive molecules isomer selectively by coupling distinct photochemistry schemes with soft photoionization.

**MATERIALS AND METHODS**

**Experimental**

The experiments were performed at the W. M. Keck Research Laboratory in Astrochemistry (19). The experimental setup consists of a contamination-free stainless steel ultrahigh vacuum chamber (UHV) evacuated to a base pressure of a few $10^{-11}$ torr by magnetically levitated turbo molecular pumps coupled to oil-free scroll backing pumps. Within the chamber, a silver mirror substrate is interfaced to a cold finger, which is connected to a closed cycle helium compressor (Sumitomo Heavy Industries, RDK-415E). Using a doubly differentially pumped rotational feedthrough (Thermionics Vacuum pumps. Within the chamber, a silver mirror substrate is interfaced to a multichannel plate is based on

The ice composition was determined via a modified Beer-Lambert solution. The FTIR spectra of the pristine ice are shown in fig. S1. Resonant and nonresonant processes ($2\nu_1$) were calculated to be 360 ± 40 nm and 830 ± 90 nm, respectively (table S1). The experiments were performed with $10.49$-eV photoionization energy and repeated at $9.10$ eV to distinguish between the P$_3$N$_3$ isomers. The $10.49$-eV (118.222 nm) light was generated via frequency tripling ($\omega_{\text{TPD}} = 3\omega_0$) of the third harmonic (355 nm) of the fundamental of a Nd:YAG laser (YAG A) in pulsed gas jets of Xe. To produce $9.10$ eV, the third harmonic (355 nm) of a Nd:YAG laser was used to pump a coumarin 450 dye [ethanol (0.20 g liter$^{-1}$)] to obtain 445.132 nm (2.72 eV; Sirah, Cobra-Stamp), which underwent a frequency doubling process to achieve $\omega_0 = 222.566$ nm (5.57 eV) ($\beta$-BaB$_2$O$_4$ crystals, 57.4°). A second Nd:YAG laser (second harmonic at 532 nm) pumped rhodamine 610/640 dye mixture [ethanol (0.17/0.04 g liter$^{-1}$)] to obtain $\omega_0 = 607$ nm (2.04 eV), which then combined with $2\omega_0$, using xenon as a nonlinear medium and generated $\omega_{\text{TPD}} = 136.246$ nm (9.10 eV) at $10^{12}$ photons per pulse. The VUV light was spatially separated from other wavelengths [due to multiple resonant and nonresonant processes ($2\omega_0 + \omega_0; 3\omega_0; 3\omega_0$)] using a lithium fluoride biconvex lens (ISP Optics) and directed 2 mm above the sample to ionize subliming molecules. The ionized molecules were mass analyzed with the ReTOF-MS, where the arrival time to a multichannel plate is based on $m/z$, and the signal was amplified with a fast preamplifier (Ortec 9305) and recorded with a personal computer multichannel scalar (FAST ComTec, P7888-1E), which is triggered via a pulse delay generator at $30$ Hz. Here, the ReTOF signal is the average of 3600 sweeps of the mass spectrum in 4-nm bin widths, which corresponds to an increase in the substrate temperature of 2 K.

All computations were carried out with Gaussian 16 Revision A.03 (30) (Figs. 2 and 5, figs. S3 and S4, and tables S3 to S5). For geometry optimizations and frequency computations, the DFT B3LYP functional (31–33) was used with the Dunning correlation-consistent split valence basis set cc-pVTZ (34). On the basis of these geometries, the corresponding frozen core–coupled cluster (35–38) CCSD(T)/cc-pVQZ single-point energies were computed and extrapolated to complete basis set limit (39) CCSD(T)/CBS with B3LYP/cc-pVTZ zero-point vibrational energy (ZPVE) corrections. The adiabatic IEs were computed by

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taking the ZPVE-corrected energy difference between the neutral and ionic species that correspond to similar conformations. The UV-vis spectra for cyclotriphosphazene (P₃N₃ isomer, 2) and 1,3,5-triphosha-2,4,6-triazacyclob[2.2.0]hexa-2,5-diene (P₃N₅ isomer, 5) were computed using TD-B3LYP method with a cc-pV(TZ) basis set. For all TD-B3LYP computations, 16 transitions into the first excited singlet state were computed. The geometries and frequencies of 2 and 5 were also computed at the CCSD(T) cc-pV(TZ) level of theory. NICSs were computed at the B3LYP/cc-pVTZ level of theory. Natural bond orbitals were computed with the NBO6 program (40).

SUPPLEMENTARY MATERIALS
Supplementary material for this article is available at http://advances.sciencemag.org/cgi/content/full/6/30/eaba6934/DC1

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