Neutral binary chalcogen–nitrogen and ternary S, N,P molecules: new structures, bonding insights and potential applications

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Early theoretical and experimental investigations of inorganic sulfur–nitrogen compounds were dominated by (a) assessments of the purported aromatic character of cyclic, binary S,N molecules and ions, (b) the unpredictable reactions of the fascinating cage compound S₄N₄, and (c) the unique structure and properties of the conducting polymer (SN)x. In the last few years, in addition to unexpected developments in the chemistry of well-known sulfur nitrides, the emphasis of these studies has changed to include nitrogen-rich species formed under high pressures, as well as the selenium analogues of well-known S,N compounds. Novel applications have been established or predicted for many binary S/Se,N molecules, including their use for fingerprint detection, in optoelectronic devices, as high energy-density compounds or as hydrogen-storage materials. The purpose of this perspective is to evaluate critically these new aspects of the chemistry of neutral, binary chalcogen–nitrogen molecules and to suggest experimental approaches to the synthesis of target compounds. Recently identified ternary S,N,P compounds will also be considered in light of their isoelectronic relationship with binary S,N cations.

1. Introduction

In contrast to the oxides of nitrogen NO, NO₂, and N₂Oₓ (x = 1, 3, 5), which all have acyclic structures, sulfur and nitrogen form a variety of neutral binary compounds with cyclic, polycyclic or polymeric structures.¹ These include the intriguing,
The structures and properties of the sulfur–nitrogen nitrides 1,3-S3N2 and 1,3-S4N2 (4) provide an interesting comparison. The putative five-membered ring cyclo-1,3-S3N2 is an antiaromatic (π-electron) system, whereas the known six-membered ring cyclo-1,3-S4N2 is formally a 10π-electron molecule, which adopts a half-chair conformation in the solid state. The optimised geometries for 4 gives values of S–S and S–N bond lengths that are in close agreement with the experimental data. For example, the calculated S–S bond length is 2.090 Å, cf. 2.061 Å from the X-ray structure. This congruence establishes confidence in the calculated S–S bond length of 2.302 Å for cyclo-1,3-S3N2. The elongated S–S bond is presumably a result of inherent strain in the five-membered ring, as suggested by the predicted value of the internal bond angle < NSS = 94.5°, cf. 105.2° in the six-membered ring 4.

As an alternative to a cyclic structure, in 2018 Chen and co-workers have proposed a stable two-dimensional trisulfur dinitride with three polymorphs comprised solely of σ bonds. The crystal structure of the most stable form, α-S3N2 (space group Pmn21), consists of condensed 12-membered (S6N6) rings (Fig. 1). The S3N2 crystal is predicted to be dynamically,
thermally and chemically stable on the basis of the computed phonon spectrum and *ab initio* molecular dynamics simulations. The predicted “chemical stability” is surprising in view of the known susceptibility of sulfur–nitrogen compounds to degradation by both nucleophiles and electrophiles.\(^{10a,b}\) Calculations also indicate that α-S\(_3\)N\(_2\) will be a wide, direct band gap (3.92 eV) semiconductor with possible optoelectronic applications such as blue or UV light-emitting diodes and photodetectors.

A subsequent study of 2D monolayer crystals of E\(_3\)N\(_2\) (E = S, Se) employing DFT calculations predicted α-heart and ß-heart structures, which exhibit different mechanical and electrical properties.\(^{10b}\) The former structure resembles that depicted for α-S\(_3\)N\(_2\) in Fig. 1. The E\(_3\)N\(_2\) materials are predicted to have unusual mechanical properties as indicated by the values of Negative Poisson’s Ratios indicating auxetic behaviour.\(^{10c}\)

The recent computational predictions of the unique structure and mechanical properties of S\(_3\)N\(_2\) provide a strong incentive for pursuing experimental approaches to this unknown binary sulfur nitride. A carefully controlled cyclocondensation reaction of CISSCI with Me\(_2\)SiNSNSiMe\(_3\) under high-dilution conditions is a plausible first attempt.\(^{14}\) An alternative route is the one-electron chemical or electrochemical reduction of the known radical cation [S\(_3\)N\(_2\)]\(^+\), which is prepared by oxidation of S\(_3\)N\(_4\) with AsF\(_3\) in liquid SO\(_2\).\(^{16}\) The reaction of 1,3-S\(_3\)N\(_2\) (4) with one equivalent of PPH\(_3\) is likely to result in attack of the nucleophile at the S(π) centre rather than the required removal of a single sulfur atom from the –S–S–S– unit.

3. Molecules with equal numbers of chalcogen and nitrogen atoms

3.1. Tetrasulfur and tetr selenium tetranitride E\(_4\)N\(_4\) (E = S, Se)

The S\(_4\)N\(_4\) (E = S) molecule is regarded as the quintessential binary sulfur–nitrogen compound in view of its multifarious applications as a reagent in inorganic and organic chemistry, despite a tendency to detonate under the influence of heat or friction.\(^{10b}\) Nonetheless, the availability of simple synthetic procedures, the selenium analogue 1 (E = Se) has undergone limited use for the preparation of selenium–nitrogen compounds owing to the explosive nature of this reagent.\(^{10a,b}\) Some new aspects of the chemical reactivity of 1 (E = S), including the electrochemical reduction and photochemical behaviour, are discussed in a recent book chapter.\(^{18}\) This section will consider more recent developments in the chemistry of S\(_4\)N\(_4\), especially those involving an understanding of the electronic structure and unusual reaction chemistry of this cage compound.

### 3.1.1. Electronic structure of 1 (E = S, Se); transannular E··E interactions

The strength of the intramolecular S··S interaction in 1 (E = S) has been estimated to be of the order of a hydrogen bond on the basis of the noncovalent interaction index.\(^{15}\) Tuononen et al. have investigated the nature of this interaction in 1 (E = S, Se) with high-level theoretical methods.\(^{6}\) Their results showed that the elongation of the transannular E··E bonds compared to a typical single-bond value can be attributed to surprisingly large correlation effects. Furthermore, there is some singlet diradical character associated with each E··E interaction. Although the consequences of this feature of the electronic structure of 1 (E = S) have not been investigated intentionally by experimentalists, an early report by Mews described the preparation of the di-substituted derivative S\(_2\)S’-1,5-S\(_4\)N\(_4\)ON(CF\(_3\))\(_2\) upon reaction of S\(_3\)N\(_2\) with two equivalents of the nitroxide radical (CF\(_3\))\(_2\)NO\(^{16}\). The reaction of S\(_3\)N\(_4\) with molecular H\(_2\) is worthy of investigation computationally and experimentally in view of this diradical character and the potential use of sulfur–nitrogen compounds as hydrogen-storage materials as discussed in section 3.2.5.

A different aspect of the bonding interactions in 1 (E = S) was reported in 2016 by Jenkins and co-workers.\(^{17}\) By applying QTAIM (quantum theory of atoms in molecules) methodology to all 18 modes of vibration of the cage molecule, these authors found a considerable degree of metallicity in the S–S and S–N bonding interactions, on the basis of the delocalisation of the electron density away from the atoms. In particular, considerable metallic behaviour was apparent in the S–S bond critical points for all modes.\(^{17}\) The experimental implications of this finding are yet to be investigated.

### 3.1.2. Unusual reaction chemistry involving S\(_4\)N\(_4\)

Numerous complexes of S\(_4\)N\(_4\) with Lewis acids have been structurally characterised and the bonding trends in these N-bonded adducts have been investigated.\(^{16,18}\) In 2018 Si and Ganguly described first principles calculations of the interaction of S\(_4\)N\(_4\) with benzene and naphthalene.\(^{19}\) These authors found that the σ hole of the sulfur atoms of S\(_4\)N\(_4\) engages in non-covalent interactions with these aromatic π-systems. For example, the interaction energy of S\(_4\)N\(_4\) with three naphthalene rings was estimated to be ca. 20 kcal mol\(^{-1}\).\(^{15}\) The S··π interaction is the main driving force in these complexes of S\(_4\)N\(_4\) with only a marginal contribution from the N atoms. The reactions of S\(_4\)N\(_4\) with alkynes have been investigated in con-
Considerable details and shown to cause cleavage of the cage molecule to form the heterocycle 1,2,5-thiadiazole as the major product along with the 7-membered rings (RC)2SN3 and RCSN3. The only example of adduct formation between S4N4 and an organic compound was reported more than 20 years ago by Konarev et al. These authors determined the single crystal X-ray structural analysis of the non-stoichiometric complex C40(S4N4)1.33(C6H6)0.67 and found that only the sulfur atoms participate in bonding to the fullerene. A more detailed experimental investigation of complexes of S4N4 with aromatic systems is warranted in view of the computational predictions.

A convenient synthesis of S4N4 involves the cyclocondensation reaction of [(Me3Si)2N]2S with an equimolar mixture of SCl2 and SO2Cl2. The complexity of the reaction pathway for this transformation is highlighted on the cover page of a recent textbook with a focus on “arrow-pushing” in inorganic chemistry. Ghosh and Berg propose a twelve-step process with the sequential participation of the reagents SCl2 followed by SO2Cl2. However, the combination of SCl2 and SO2Cl2 generates SCl4 (+SO2) spontaneously, so a cyclocondensation via a series of thermodynamically favoured eliminations of Me3SiCl is suggested (Scheme 2).

Kumar and Muralidharan have reported the hexamethyl-disilazane (HMDS)-assisted synthesis of β-In2S3, CuS and Ag2S nanoparticles via reactions of sulfur with an appropriate metal salt in the presence of HMDS. In order to explain the role of HMDS in the synthesis of these metal sulfides, the formation of a novel polymeric intermediate [SN(SiMe3)]n, was invoked. The characterization of this purported polymer was based upon GPC (gel permeation chromatography) analysis of the distilled material, as well as 29Si NMR spectra and HRMS (high resolution mass spectrometry) data. The fact that this intermediate could not be isolated as a pure compound was attributed by the authors to the similarity of its boiling point with that of HMDS. Since the boiling point of HMDS is only 125 °C, this explanation is highly unlikely. Orange crystals obtained as a minor by-product in the synthesis of Ag2S were identified by XRD as S3N4 as a minor by-product; however, the formation of this binary sulfur nitride from the purported sulfur imide polymer is not readily explained.

3.2. Disulfur and diselenium dinitride E2N2 (2, E = S, Se)

In contrast to the numerous applications of S4N4 as a source of novel inorganic and organic sulfur–nitrogen compounds, the use of S2N2 in synthesis is rare owing to its highly labile nature. However, both electrochemical and chemical reduction produce the [S2N2]− anion, which can be isolated as ion-separated salts. This section will consider recent developments in our understanding of the chemistry of S2N2 and its heavier chalcogen homologues, including electronic structures, spectroscopic characterisation, the isomers formed upon photoysis, topochemical polymerization to give (SN)3, and possible use as a hydrogen-storage material.

3.2.1. Electronic structures. The square-planar structure of S2N2 (2, E = S), a volatile white crystalline solid, was established 45 years ago. Despite this longevity, the electronic structure of this four-membered ring and its heavier chalcogen analogues continues to generate controversy centred around their aromaticity and diradical character. The most recent developments are discussed below.

In 2004 Head-Gordon et al. concluded that the diradical character of S2N2 is not significant on the basis of a large HOMO–LUMO gap (5.1 eV), a large singlet–triplet vertical transition energy (3.6 eV), and the lack of a spin-unrestricted solution at the level of DFT using the B3LYP functional. They asserted, however, that S2N2 is aromatic based on its negative NICS value (−26.2 ppm), albeit with a small aromatic stabilisation energy, since only two of the six π electrons contribute to the bonding.

**Scheme 2** Formation of S4N4 via cyclocondensation of [(Me3Si)2N]2S with SCl4.
Concurrently, Tuononen and co-workers investigated the electronic structures and molecular properties of $S_2N_2$ \(^{27b}\) as well as the unknown heavier chalcogen analogues $Se_2N_2$ and $Te_2N_2$ \(^{27b}\) by employing various computational methods. These authors agreed that all of these four-membered rings can be described as 2π-electron aromatics, but with minor singlet diradical character ranging from 6% in $S_2N_2$ to 10% $Te_2N_2$ located solely on the two nitrogen atoms. \(^{27b}\)

In 2012 Braida et al. carried out an \textit{ab initio} valence bond study of $E_2N_2$ which confirmed that the diradical sites are the two nitrogen atoms ($E = Se, Te$), however they estimated that the contribution from diradical structures is close to 50%. \(^{28}\) Their calculations of vertical resonance energies gave values of about 80% of the predicted value of benzene for $S_2N_2$ with somewhat lower values for $Se_2N_2$ and $Te_2N_2$, consistent with the notion of aromatic character for $S_2N_2$ based on magnetic criteria. An important conclusion from these results is that diradical character and aromaticity are not mutually exclusive. \(^{28}\)

In a sequel to his earlier work on valence bond structures for $S_2N_2$, Harcourt \(^{29}\) postulated that the six Lewis structures assumed by Braida \textit{et al.} \(^{28}\) are equivalent to resonance between two \textit{increased-valence} structures with one-electron and fractional electron-pair π bonds.

In 2018 Karadakov \textit{et al.} employed magnetic criteria to study the bonding and aromaticity in the ground state, first singlet excited and lowest triplet electronic states of $S_2N_2$. \(^{25}\) Their results indicate that (a) the $S_0$ electronic state is aromatic, but less so than the electronic ground state of benzene, (b) $S_1$ is strongly antiaromatic, and (c) $T_1$ is moderately antiaromatic to a similar extent to that observed in the electronic ground state of square cyclobutadiene. Thus, they assert that $S_2N_2$ is the first example of an inorganic ring for which theory predicts substantial changes in aromaticity upon vertical transition from the ground state to the first singlet or lowest triplet electronic states. \(^{25}\)

### 3.2.2. Spectroscopic characterisation

Tuononen \textit{et al.} \(^{27b}\) found that coupled cluster and multiconfigurational approaches, as well as density functional methods, were the most reliable for the prediction of spectroscopic properties such as vibrational frequencies, $^{14}N$, $^{15}N$ and $^{77}Se$ NMR chemical shifts, and excitation energies of sulfur and selenium nitrides. The congruence between the computational results and experimental data for $S_2N_2$ provides some confidence in the predictions for selenium-containing analogues $SSeN_2$ and $Se_2N_2$.

The calculated IR-active vibrational frequencies \(^{27b}\) are in reasonable agreement with experimental data from the IR and Raman spectra of $S_2N_2$ obtained as a solid condensate and in $N_2$ or $CH_4$ matrices at 15–35 K, which were reported by Downs \textit{et al.} \(^{30}\) By using 30% $^{15}N$-enrichment, these authors confirmed that the isolated $S_2N_2$ molecule has essentially the same square-planar geometry as the crystalline solid; calculations of the $S$–$N$ stretching force constant indicated a bond order only slightly greater than 1. \(^{30}\) In 2014 the high-resolution gas-phase FTIR spectrum of $S_2N_2$ together with a re-investigation of the IR spectrum in solid argon at 16 K were reported by Perrin, Beckers and co-workers. \(^{31}\) The main difference between the gas-phase and solid-state structures of $S_2N_2$ involves the bond angles; in the gas phase the angle at nitrogen $\langle SNS \rangle$ is smaller than $\langle NSN \rangle$, whereas the opposite is true in the solid-state structures. \(^{31}\) These small disparities may be the consequence of the intermolecular interactions that occur between $S_2N_2$ rings in the plane of the chain of rings in the solid state. Cremer \textit{et al.} have computed a pseudorotation barrier of $<2$ kJ mol$^{-1}$ for $S_2N_2$ indicative of a floppy molecule, however these authors maintain that this characteristic does not alter the electronic structure of the four-membered ring. \(^{32}\)

The predicted IR and Raman spectra of the selenium-containing systems $Se_2N_2$ and $SSeN_2$ are depicted in Fig. 2. \(^{27b}\) The band at $\sim590$ cm$^{-1}$ in the IR spectrum of $Se_2N_2$ should be considerably less intense than the other two bands. In a similar vein, the intensity of the band at 822 cm$^{-1}$ is estimated to be more than twice those of the other two bands.

**Fig. 2** (a) IR/Raman spectra of $S_2N_2$. \(^{30}\) (Adapted with permission from R. Evans, A. J. Downs, R. Köppe and S. C. Peake, \textit{J. Phys. Chem. A}, 2011, \textit{115}, 5127–5137. @ 2011 American Chemical Society), and the theoretically predicted IR/Raman spectra of (b) $Se_2N_2$ and (c) $SeSN_2$ \(^{27b}\) (adapted with permission from H. M. Tuononen, R. Suontamo, J. Valkonen and R. S. Laitinen, \textit{J. Phys. Chem. A}, 2005, \textit{109}, 6309–6317. @ 2005 American Chemical Society).
Tuonen et al., have also calculated the $^{14}$N, $^{34}$N and $^{77}$Se NMR chemical shifts for the chalcogen nitrides $S_2N_2$, $SeSN_2$, $Se_2N_2$, $S_4N_4$ and $Se_2S_2N_4$. Surprisingly, no experimental NMR data are available for $S_2N_2$. However, the calculated $^{14}$N and $^{34}$N NMR chemical shifts for $S_2N_2$ and the known mixed selenium–sulfur nitride $Se_2S_2N_4$ are in excellent agreement with the experimental data providing some confidence in the predicted values of 95 ± 20, 125 ± 20 and 185 ± 20 ppm for $S_2N_2$, $SeSN_2$ and $Se_2N_2$, respectively. The calculated $^{77}$Se chemical shifts for 1,5-$Se_2S_2N_4$ at different levels of theory are $ca.$ 200 ppm lower than experimental values. By contrast, the predicted PBEPE and CAS $^{77}$Se chemical shifts of 196 and 1886 ppm for $Se_2N_2$ are only slightly lower than the experimental value of 1936 ppm. Assuming a similar error for $SeSN_2$ and $Se_2N_2$, $^{77}$Se chemical shifts of 1820 ± 20 and 1780 ± 20 ppm, respectively, are estimated for these structurally analogous four-membered rings. The ternary chalcogen nitride $SeSN_2$ has been proposed as an intermediate in the formation 1,5-$Se_2S_2N_4$ from reactions of $[Me_3SiNSN]_2E$ ($E = S, Se$) with $ECl_2$ ($E = S, Se$), but this potential precursor of a mixed sulfur–selenium nitride polymer (SNSeN)$_6$ has eluded isolation.

The experimental UV spectrum of $S_2N_2$ consists of a broad band in the range 4.5–5.8 eV comprising two overlapping absorptions centred at approximately 5.0 eV (~250 nm) which is attributed to the $\pi \rightarrow \pi^*$ electronic transition. The calculated band-gap energies decrease as the selenium content in the ring increases as illustrated by colourless $S_2N_2$ (3.32 eV, ~375 nm) the predicted orange-yellow colour of $SSeN_2$ (2.50 eV, ~495 nm) and red colour of $Se_2N_2$ (2.20 eV, ~560 nm). This information should be valuable for the identification of the selenium-containing four-membered rings.

### 3.2.3. Photolysis: isomers of $S_2N_2$

In 2015 Zeng, Beckers and co-workers investigated the UV photolysis of $S_2N_2$ in solid argon matrices using two different light sources ($\lambda = 248$ or 255 nm). Although calculations predict both linear SNNS and cyclic 1,2,3,4-dithiadiazete ($C_{2v}$) to be in energy the same as the cyclic $S_2N_2$ ($D_{2h}$), neither of these two isomers has been observed experimentally. Instead, by employing IR spectra of both natural abundance and $^{34}$N-enriched $S_2N_2$, these authors were able to identify two open shell isomers, trans-SNNS ($8$) and cis-SNNS ($9$), as well as a closed shell $C_{2v}$ dimer ($SN)_2$ ($10$), as the primary photolysis products (Scheme 3). The identification of the open shell isomers is relevant to the mechanism of polymerisation of $S_2N_2$ into $(SN)_n$, which is discussed in the next section.

### 3.2.4. Topochemical polymerisation to $(SN)_n$

**Applications in forensic science.** The spontaneous topochemical polymerisation of $S_2N_2$ into $(SN)_n$, a polymer with fascinating conducting properties, has been known for more than 40 years. Kelly et al., have demonstrated that this process can be achieved in the zeolite Na-ZSM-5, which has a pore size that is able to accommodate $S_2N_2$ as well as infinite channels along the $b$-axis to allow for polymer growth. The same group also reported that the exposure of fingerprints on various surfaces, e.g. glass, metal, paper or ceramic surfaces, to $S_2N_2$ vapour generated an image of the fingerprint due to the formation of blue-black $(SN)_n$ polymer (Fig. 3a). It was suggested that this serendipitous observation could have applications in forensic science, although the design of a practical and portable apparatus for the production of $(SN)_n$ was recognised as a major impediment.

In 2013 Takaluoma et al. carried out molecular dynamics simulations of the topochemical polymerisation of $S_2N_2$ in the solid state using DFT methods and periodic functions. The calculations involved both high pressures and slightly elevated temperatures in order to increase the reaction rate. The formation of $(SN)_n$ is initiated by cleavage of one S–N bond in $S_2N_2$ followed by a very rapid attack of the resulting open-chain isomer, e.g. 8 in Scheme 3, on a neighbouring ring (Fig. 4). Propagation occurs along the $a$ axis throughout the lattice. The packing changes from the herringbone structure of the $S_2N_2$ lattice to the layered structure of $(SN)_n$. Although the metrical data for the polymer chain are in good agreement with the experimental crystal structure, there is less long-range order between neighbouring chains, possibly due to the influence of pressure or the poor description of dispersion by density functional theory.

### 3.2.5. A hydrogen-storage material?

Datta has described quantum chemical investigations of the interaction of H$_2$ molecules with binary S,N rings including $S_2N_2$. An adduct of $S_2N_2$ with two molecules of H$_2$ located in an eclipsed fashion above and below the ring was found to have only dispersive interactions with a low binding energy ($\sim 1.5$ kcal mol$^{-1}$). However, when the ring is doped with a transition metal such as Ni(0), Pd(0) or Pt(0), complexes that incorporate

![Scheme 3](image-url)
three \( \text{H}_2 \) molecules for each metal centre were found to have binding energies of \(-67.75\), \(-18.73\) and \(-211.75\) kcal \( \text{mol}^{-1} \) for \( M = \text{Ni}, \text{Pd} \) and \( \text{Pt} \), respectively. As illustrated in Fig. 5, the binding modes of \( \text{H}_2 \) to the metal centre are markedly different. In the Ni and Pd complexes four H atoms appear as hydrides while one \( \text{H}_2 \) molecule is \( \eta^2 \)-coordinated to the metal. By contrast, in the most stable Pt complex all six hydrogen atoms exist as hydrides with five of them forming an almost regular pentagon.\(^{35}\) Although the practicality of using binary S, N compounds as hydrogen-storage materials may be questioned by experimentalists, investigations of the interaction of extended systems similar to \((\text{SN})_x\) may be worthy of experimental interrogation.

4. **Nitrogen-rich chalcogen nitrides**

Early work on nitrogen-rich sulfur nitrides such as \( \text{SN}_2 \) and \( \text{SN}_4 \) was mainly limited to the purported involvement of these species as fleeting intermediates. However, recent experi-
mental and theoretical investigations of the formation and structures of this class of sulfur nitrides and their selenium analogues under high pressure have revealed fascinating properties with potential applications in optoelectronics and as high-energy-density materials. This section is organised according to increasing nitrogen content of these binary molecular species. Nitrogen-rich selenium nitrides are discussed in section 4.4.

4.1. The isomers NNS and NSN

The lowest energy isomer NNS, a heavier analogue of the stable molecule N₂O, decomposes above 160 K. It is generated by flash photolysis of 5-phenyl-1,2,3,4-thiatriazole and has been characterised by high-resolution mass spectrometry and IR spectroscopy. The symmetrical isomer NSN has been invoked as an elimination product during the transformations of polycyclic S–N compounds, but the synthesis and structure of this simple sulfur nitride remained elusive until very recently (vide infra).

In 2016 Zhang and co-workers investigated the stability, electronic structure and optical properties of a P3m1 phase of 1T-EN₂ (E = S, Se, Te) by means of DFT calculations (the prefix 1T refers to one layer per trigonal unit cell). The calculated crystal structure of the two-dimensional material 1T-SN₂ is comprised of six-coordinated S atoms and three-coordinate N centres (Fig. 6a). These chalcogen nitrides 1T-EN₂ (E = S, Se, Te) are endothermic materials. 1T-SN₂ has dynamical stability based on phonon spectra and calculated cohesive energies, but the thermal and mechanical stabilities have yet to be determined. Band-structure calculations revealed indirect band gaps of 2.825, 2.351, and 2.336 eV for 1T-SN₂, 1T-SeN₂, and 1T-TeN₂, respectively. Furthermore, the application of biaxial...
strain induced a transition from a wide-band-gap semiconductor to a metal. These findings suggest that the binary chalcogen nitrides 17-EN$_2$ are promising materials for applications in optoelectronic devices.

Earlier this year Chen and co-workers proposed a new phase of the SN$_2$ monolayer, namely S–SN$_3$N, by using first-principles calculations combined with the particle-swarm optimisation method. The structure of S–SN$_3$N belongs to the space group P4M2 and incorporates four-coordinate S atoms and two-coordinate nitrogen centres (Fig. 6b); the prefix S is used to indicate its square lattice and, hence, distinguish this new structure from 17-SN$_2$. The S–SN$_3$N monolayer is a remarkably stable semiconductor with an indirect band gap of 2.79 eV, cf. 2.825 eV for 17-SN$_2$. The co-existence of both out-of-plane and in-plane negative Poisson’s ratios indicates 3D auxetic properties for this new material, which may have applications in mechanical or optoelectronics devices.

In 2017 Cui and co-workers performed particle-swarm optimisation calculations on the sulfur–nitrogen system up to 200 GPa. Three binary sulfur–nitrogen solids were determined to be thermodynamically stable: Pnma (SN)$_3$, Pnnm SN$_2$, and C2/c SN$_4$ at pressures of 37, 43 and 48 GPa, respectively. Their crystal structures are depicted in Fig. 7. A polymeric 3D arrangement was predicted for SN$_2$ (Fig. 8c). Partial density of states calculations indicated that SN$_2$ is a semiconductor with a direct band gap of 0.66 eV at 60 GPa.

Last year a seminal experimental investigation of the formation of sulfur nitrides in the S–N$_2$ system at high pressures by Laniel and co-workers confirmed the predictions for SN$_2$. These authors found that the reaction of elemental sulfur and N$_2$ gas in a diamond anvil cell at pressures above 60 GPa with laser heating produces an orthorhombic ($P_{nnm}$) SN$_3$ compound. The lattice parameters of this material obtained from powder diffraction data match closely the theoretically predicted crystalline structure of SN$_2$. Single crystal X-ray diffraction analysis revealed a CaCl$_2$-type structure composed of edge-sharing SN$_6$ octahedra with N atoms that are triply coordinated by S atoms (Fig. 8). The SN$_2$ solid was found to be metastable down to ca. 20 GPa, at which point it dissociates into its elements.

### 4.2. Thiatetrazole SN$_4$

The nitrogen-rich thiatetrazole SN$_4$ has attracted the attention of both experimental and theoretical chemists for more than 20 years. Klapötke and Schulz attempted to prepare SN$_4$ from the reaction of the cation [NS]$^+$ in [NS][AsF$_6$] with the azide ion [N$_3$]$^-$ in cesium azide. The final product was a black precipitate identified as the polymer (SN)$_4$; no experimental evidence ($^{14}$N NMR) could be found for the expected SN$_4$ intermediate.

Furthermore, no spectroscopic evidence for the formation of SN$_4$ was garnered from the recent study of the S–N$_2$ system at high pressures, which led to the characterisation of metastable SN$_2$ (section 4.1).

The main focus of early theoretical investigations of SN$_4$ was an estimate of the aromaticity of the cyclic isomer, which is a 6e-electron five-membered ring formally related to thiophene by the replacement of four CH units by four nitrogen atoms. Schleyer and co-workers calculated NICS (nucleus independent chemical shift) and NICS(1) values of −18.40 and −17.48 for cyclo-SN$_4$; the large negative values are indicative of substantial aromatic character in the ring.

This conclusion is supported by the calculated value of 18.56 for the ASE (aromatic stabilisation energy). Nitrogen-rich sulfur nitrides are of considerable interest as high energy-density materials. The energy content of cyclo-SN$_4$ was first computed by Wang and co-workers using DFT methods at the B3LYP/6-311+G** level. It was found that cyclo-SN$_4$ is ca. 100 kcal mol$^{-1}$ higher in energy than the...
decomposition products $1/8\text{S}_8 + \text{N}_2$. However, the low dissociation barrier of 7.0 kcal mol$^{-1}$ for the decomposition of SN$_4$ into N$_2 + \text{N}_2\text{S}$ (Scheme 4) precludes the isolation of this sulfur nitride under ambient conditions.$^{44}$

4.3. Dithiatetrazine S$_2$N$_4$

A planar six-membered ring cyclo-S$_2$N$_4$ would be an 8π-electron (antiaromatic) system isoelectronic with dithiatrazines (RC)S$_2$N$_3$. The parent dithiatrazine (R = H) is not known, but aryl derivatives (R = Ph) exist as cofacial dimers in the solid state.$^{1}$ Numerous acyclic and cyclic isomers are possible for the putative nitrogen-rich sulfur nitride S$_2$N$_4$. Schleyer and co-workers have carried out computational investigations of the thermochemical stability and kinetic persistence of various S$_2$N$_4$ isomers at the B3LYP/6-311+G(3df) DFT level.$^{13}$ The results were compared with the corresponding data for the known sulphur nitrides S$_2$N$_2$, S$_4$N$_2$ and S$_4$N$_4$, as well as the nitrogen-rich species SN$_4$ and S$_2$N$_2$. A boat conformation of 1,4-S$_2$N$_4$ (11), cf. 1,4-S$_2$(CH)$_4$, was predicted to be the most stable of the three possible six-membered rings. The other most thermodynamically stable isomers include two five-membered rings SN$_4(=\text{S})$ (12 and 13) and two acyclic species SNSNNN (14 and 15) (Scheme 5). From a consideration of the kinetic stability (dissociation barriers) of these five isomers it was concluded that the acyclic isomer 14 is the most viable candidate for synthesis. However, its dissociation barrier to give N$_2$ and cyclo-S$_2$N$_2$ is only 21.6 kcal mol$^{-1}$, cf. 51 kcal mol$^{-1}$ for cyclo-S$_2$N$_2$ (2, E = S)$_2$. Although this barrier is considerably higher than the value of 7 kcal mol$^{-1}$ determined for the decomposition of cyclo-SN$_4$ (section 4.2), the facile formation of S$_2$N$_2$ from decomposition of SNSNNN (14) may account for the rapid production of [SN]$_x$ observed in the reaction of [SNS][AsF$_6$] with CsN$_3$ in SO$_2$(l) at $-20$ °C.$^{45}$

As a general conclusion, it is important to note that kinetic stability is more important than thermodynamic stability in determining which binary sulfur–nitrogen species can be isolated under ambient conditions. The isomers of the nitrogen-rich cation [S$_2$N$_3$]$^+$ a 6π-electron system, provide a compelling example of this principle. Despite being less thermodynamically stable than the 1,3-isomer, the 1,2-isomer has been isolated and structurally characterised in the salt [S$_2$N$_3$]$^+\text{[Hg}_2\text{Cl}_6]$.$^{46}$ Although it is the only known monocyclic, nitrogen-rich S,N cation, several salts of the tricyclic cation [S$_4$N$_5$]$^+$ have been structurally characterised.$^{47}$ The relative stability of 1,2-[S$_2$N$_3$]$^+$ is attributed to the higher dissociation barriers for its fragmentation compared to the 1,3-isomer for which the dissociation pathway to form [SNS]$^+$ + N$_2$ has a barrier of only 2.6 kcal mol$^{-1}$.$^{44a}$

4.4. Trithiatetrazepine S$_3$N$_4$

Planar, cyclic S$_3$N$_4$ (16) would be a 10π-electron (aromatic) system (see Scheme 6) isoelectronic with the known trithiatetrazepine (HC)S$_3$N$_3$.$^{49}$ However, the isolation of 16 at room temperature is unlikely in view the low energy barrier

![Scheme 4](image-url)
5. Nitrogen-rich selenium nitrides

As mentioned in section 1, the only binary selenium nitride that has been isolated under ambient conditions and structurally characterised is Se$_4$N$_4$ (1, E = Se),$^{3a}$ although metal complexes of Se$_2$N$_2$ (2, E = Se) are also known.$^{3c,d}$ In that context investigations of the high-pressure phase diagrams of the binary selenium–nitrogen system by Cui and co-workers reported in 2019 are of considerable interest.$^{52}$ These authors identified four stable compounds at high pressures: Cmc$_2$-SeN$_2$, P$_2$/m-SeN$_3$, PI-SeN$_4$ and PI-SeN$_5$. As illustrated in Fig. 9, these novel nitrogen-rich materials are predicted to incorporate a variety of polytellurium arrangements in the solid state.$^{52}$

The binary selenium nitride Cmc$_2$-SeN$_2$ has a layer structure (Fig. 9a), while P$_2$/m-SeN$_3$, PI-SeN$_4$ and PI-SeN$_5$ incorporate $\text{N}_6$ chains (Fig. 9b), oligomeric $\text{N}_6$ chains (Fig. 9c) or distorted [N$_6$]$^{1-}$ anionic rings alternating with layers of $\text{N}_6$ chains (Fig. 9d). The high energy-content of the latter three phases is reflected in the values of their energy densities, which are 3.27, 3.26 and 4.08 kJ g$^{-1}$, respectively.$^{52}$ Information on the dissociation barriers for these binary selenium nitrides is not available.

6. Ternary S,N,P molecules

A phosphorus atom P has the same number of electrons as an S$^+$ cation, consequently there can be an isoelectronic relationship between certain ternary S,N,P neutral molecules and well-known binary S,N cations. The latter species can be isolated as thermally stable salts, which have been structurally characterised, and the cations have been shown to exhibit a range of chemical reactions.$^{1a,b}$ The recent identification of the ternary systems SNP$^{12a}$ and SP$_2$N$_2^{12b}$ provide cogent examples of this isoelectronic connection.

Francisco, Zeng et al., conducted an initial investigation of the pyrolysis and photolysis of the triazide SP(N$_3$)$_3$.$^{12a}$ The experimental work was presaged by DFT calculations of the three triatomic SNP isomers, viz. linear SNP (18), linear SPN (19) and the three-membered ring cyclo-PSN (thiazaphosphirine) (20) at the B3LYP/6-311+G(3df). As indicated in Fig. 10, the cyclic isomer 20 is separated from the linear isomers 18 and 19 by barriers of 138 and 95 kJ mol$^{-1}$, respectively; the SNP arrangement 18 was predicted to be the global minimum. In one experimental approach the pyrolysis of SP(N$_3$)$_3$ at
1000 °C followed by trapping of the products in an argon matrix at 16 K generated SPN (eqn (1)). This species was identified with the help of $^{15}$N-labelling by comparison of calculated and experimental IR frequencies, which also revealed the formation of other nitrogen-containing species, PN, SN and SN$_2$.

$$\text{SP(N$_3$)$_3$} \rightarrow \text{SPN} + 4\text{N}_2 \quad (1)$$

In an alternative procedure the photolysis of SP(N$_3$)$_3$ in solid argon resulted in a stepwise dissociation to give the most stable SNP isomer 18 as the final product. In both the pyrolysis and photolysis experiments selective UV irradiation produced the linear SPN (19) and cyclo-PSN (20), which were identified by IR spectroscopy.$^{12a}$

The structure of SNP (18) can be represented by the two Lewis structures shown in Fig. 11a. The calculated P–N bond length is 1.502 Å, slightly greater than the sum of the triple bond radii for P and N (1.48 Å), and the calculated P–N stretching frequency of 1314 cm$^{-1}$ is close to that of diatomic PN (1327 cm$^{-1}$).$^{12a}$ The SNP isomer (18) is isoelectronic with the...
linear sulfur–nitrogen cation [SNS]⁺, which has an extensive
cycloaddition chemistry.⁵¹ The similarity of the frontier orbi-
tals of SNP (18) (Fig. 11b) with those of [SNS]⁺ suggest that the
ephemeral SNP species could be trapped via cycloaddition
reactions, e.g. with alkynes or nitriles.

A subsequent IR analysis of the products of the flash pyro-
lysis of SP(N₃)₃ under conditions used to generate SNP (18)
(vide supra) revealed a new species with IR frequencies that
increased in intensity with prolonged deposition time.¹² The
mid- and far-IR frequencies of this product appeared as quar-
tets when ¹⁴N labelled SP(N₃)₃ was used as a precursor, indicat-
ing a molecule with two structurally non-equivalent N
atoms.¹² In order to elucidate the identity of this species, theo-
technical calculations of the dimerisation of SNP were
conducted. As illustrated in Fig. 12, the activation energy for
head-to-tail association of SNP to give cyclo-SNPSNP is low.
However, the structural equivalence of the two nitrogen atoms
in this six-membered ring rules it out as the source of the new
IR bands. Instead, the energetically favourable loss of a sulfur
atom from cyclo-SNPNP, which is formally an 8
π-electron system, to give the five-membered ring
atom from cycloaddition chemistry.⁵³ The similarity of the frontier orbi-
tals of SNP (vide supra) to give [S₄N₄]²⁺ gives [S₄N₄]²⁺, which is formally a planar, cyclic cation [S₄N₄]²⁺, a fully delocalised
π-electron system.¹ The replacement of the two antipodal S
atoms in the supposed planar molecule S₄N₄ by P atoms gener-
ates 1,5-P₂N₄S₂, isoelectronic with [S₄N₄]²⁺. According to
Gleiter, this ternary S,N,P molecule would incorporate an
unpaired 3p electron at each P centre (21).² The diradical
species 21 would be stabilised by addition of two radicals R’ at
the phosphorus centres to give RP(NSN)₂PR (1,5-diphospha-
dithiatetrazocine (HC)₂N₂S₂ using density functional
methods.⁵⁵,⁵⁶ It was found that replacement of the two HC
groups by the more electronegative P atoms decreases the aromatic character.⁵⁵ However, the ternary ring system was
suggested to be a reasonable synthetic target.

In contrast to the planar P(III)-containing heterocycle 22,
the corresponding P(V) systems R₂P(NSN)₂PR (1,5-diphospha-
dithiatetrazocines) form bicyclic structures with a weak trans-
nular S⋯S interaction similar to that discussed for S₄N₄ in
section 3.1.1. A recent fragment-based energy analysis by
Jacobsen estimates that the cross-ring connection in 1,5-diphos-
phadithiatetrazocines is about half as strong as a typical S–S
bond.⁵⁷ This class of S,N,P heterocycle is considered to be an
unusual inorganic example of bishomoaromaticity.⁷⁶,⁵⁸

7. Summary and outlook

Although the complexities of the electronic structures of the
well-known binary sulfur nitrides S₃N₂ and S₄N₄ and their selen-
iurn analogues are still under consideration, the focus of
recent investigations of this class of inorganic compounds has
been on the potential applications of both known and
unknown systems. The foremost example is the use of the facile transformation of $\text{S}_2\text{N}_2$ into the blue-black polymer $[\text{SN}]_x$ for the detection of fingerprints on metal surfaces, which is being considered as an alternative to existing processes for this application in forensic science (section 3.2.4). A more speculative function involves the possible use of $\text{S}_2\text{N}_2$ or other sulfur–nitrogen rings as hydrogen-storage materials (section 3.2.5). Although the feasibility of this application is questionable in the case of $\text{S}_2\text{N}_2$, sulfur–nitrogen systems with higher chemical and thermal stability, e.g. C,S,N heterocycles, may be worthy of computational and experimental interrogation for this purpose.

In the area of sulfur-rich nitrides, the unknown molecule $\text{S}_2\text{N}_2$ is predicted to have quite different properties compared to those of the well-characterised six-membered ring $1,3\text{-S}_2\text{N}_2$, which is a labile, low melting compound. By contrast, computational studies indicate that crystalline $\text{S}_2\text{N}_2$ will form a two-dimensional network involving only $\sigma$ bonds rather than an $\pi$-electron five-membered ring. Furthermore, it has been proposed that $\text{S}_2\text{N}_2$ will exhibit chemical and thermal stability and behave as a wide direct band gap semiconductor with possible optoelectronic applications. As suggested in section 2, this neutral sulfur nitride should be experimentally accessible under ambient conditions so that these predictions can be tested.

Nitrogen-rich sulfur nitrides, e.g. thiattetrazole $\text{cyclo-SN}_1$, have long been considered as potential high energy-density materials (section 4.2). Until recently, this purported 5-membered ring and the acyclic triatomic molecule $\text{NSN}$ have only been invoked as fleeting intermediates in the decomposition of sulfur–nitrogen ring systems. In the last few years, however, seminal experimental and computational studies of the combination of a chalcogen (sulfur or selenium) and nitrogen gas ($\text{N}_2$) at very high pressures have yielded unanticipated information about the structures and properties of this new class of chalcogen nitride. In general, the computational work indicates that at high pressures a variety of nitrogen-rich compounds with structures that are quite different from the metastable sulfur nitrides known to exist at ambient temperatures and pressures (Scheme 1) will be formed. Experimental verification of this prediction has been provided by the identification of $\text{SN}_2$ solid at pressures above 64 GPa. This novel material has a $\text{CaCl}_2$ structure comprised of $\text{SN}_6$ octahedra with triple coordinated $\text{N}$ atoms, as forecasted by earlier computational work (section 4.1). In the light of these experimental results, the predicted formation of stable binary selenium nitrides $\text{SeN}_x$ ($x = 2, 3, 4$ or 5) at high pressures should provide an incentive for an experimental study of the selenium–N$_2$ system with the characterisation of the elusive $[\text{SeN}]_x$ polymer as one of the target molecules.

Finally, recently characterised acyclic and cyclic neutral $\text{S,N,P}$ molecules are considered in the context of the known structures and reactions of binary sulfur–nitrogen cations, which are isolectic with the ternary $\text{S,N,P}$ systems. This comparison is intended to provide some guidance to the future development of the chemistry of these interesting ternary systems.

Conflicts of interest

There are no conflicts of interest to declare.

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