Roemmele, Tracey L.

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Electrochemical and chemical reduction of disulfur dinitride: formation of \([\text{S}_4\text{N}_4]\), EPR spectroscopic characterization of the \([\text{S}_2\text{N}_2\text{H}]\cdot\) radical and X-ray structure of \([\text{Na}(15 \text{ crown-5})][\text{S}_3\text{N}_3]\)

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<td>Roemmele, Tracey; university of calgary, chemistry; University of Lethbridge Konu, Jari; University of Calgary, Department of Chemistry Boere, Rene; University of Lethbridge, Chemistry and Biochemistry Chivers, Tristram; university of calgary, chemistry</td>
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Abstract

Voltammetric studies of $S_2N_2$ employing both cyclic voltammetry (CV) and rotating disc electrode (RDE) methods on GC electrodes at RT revealed two irreversible reduction processes at ca. −1.4 V and −2.2 V in $CH_3CN$, $CH_2Cl_2$ and THF (vs. ferrocene) and no observable oxidation processes up to the solvent limit when the scan is initially anodic. However, after cycling the potential through −1.4 V, two new couples appear near −0.3 V and −1.0 V due to $[S_3N_3]^{-}/0$ and $[S_4N_4]^{-}/0$ respectively. The diffusion coefficient $D$ for $S_2N_2$ was determined to be $9.13 \times 10^{-6} \text{ cm}^2 \text{s}^{-1}$ in $CH_2Cl_2$ and $7.65 \times 10^{-6} \text{ cm}^2 \text{s}^{-1}$ in $CH_3CN$. Digital modeling of CVs fits well to a mechanism in which $[S_2N_2]^{-}$ couples rapidly with $S_2N_2$ to form $[S_4N_4]^{-}$, which then decomposes to $[S_3N_3]^-$. In situ EPR spectroelectrochemical studies of $S_2N_2$ in both $CH_2Cl_2$ and $CH_3CN$ resulted in the detection of strong EPR signals from $[S_4N_4]^{-}$ when electrolysis is conducted at −1.4 V; at more negative voltages, spectra from transient adsorbed radicals are observed. In moist solvent or with added HBF$_4$, a longer-lived spectrum is obtained due to the neutral radical $[S_2N_2H]^+$, identified by simulation of the EPR spectrum and DFT calculations.

The chemical reduction of $S_2N_2$ with Na[C$_{10}H$_{8}] or Na[Ph$_2$CO] produces [Na(15-crown-5)][S$_3N_3$], while reduction with cobaltocene gives [Cp$_2$Co][S$_3N_3$]. The X-ray structure of the former reveals a strong interaction (Na···N = 2.388(5) Å) between the crown ether-encapsulated Na$^+$ cation and one of the nitrogen atoms of the essentially planar six-membered cyclic anion $[S_3N_3]^-$. 

Introduction

The binary sulfur nitride $S_2N_2$ has been known for more than 50 years.$^1$ It is commonly prepared by the thermolysis of the cage molecule $S_4N_4$ over silver wool at ca. 220 °C,$^2$ although other sulfur-nitrogen rings, e.g. $[S_4N_3]Cl$,$^3$ and $Ph_3As=N-S_3N_3$,$^4$ have been used as a source of
S$_2$N$_2$. Disulfur dinitride has a square-planar structure with S–N bond distances intermediate between single- and double-bond values.\(^2\) The four-membered ring is an important precursor for the synthesis of the conducting polymer poly(sulfur nitride) (SN)$_x$\(^5\). In this solid-state polymerization which occurs at 0 °C over several days, colorless crystals of S$_2$N$_2$ are converted via a topochemical process into shiny bronze-colored fibers of (SN)$_x$. Recent work by Kelly et al. has shown that this polymerization process can be effected in a zeolitic matrix.\(^6\) In an intriguing contribution the same group has shown that the S$_2$N$_2$→(SN)$_x$ conversion is a potential device for latent fingerprint detection.\(^7\)

The electronic structure of this compositionally simple heterocycle continues to be a matter of debate.\(^8\) Simple electron-counting procedures lead to the conclusion that S$_2$N$_2$ is a six \(\pi\)-electron (aromatic) system.\(^9\) However, four of the six \(\pi\)-electrons occupy non-bonding \(\pi\) MOs (i.e. those with nodes either through both sulfur or both nitrogen atoms). S$_2$N$_2$ is more appropriately considered as a two \(\pi\)-hole \((4N+2)\)-electron system\(^{8g,h}\) with minor singlet diradical character (ca. 6 %), which is attributed solely to the nitrogen atoms.\(^{8d-f}\)

Apart from polymerization, investigations of the reactions of S$_2$N$_2$ have been limited. The four-membered ring forms \(N\)-bonded mono and di-adducts of the type S$_2$N$_2$:L and S$_2$N$_2$:2L with a variety of Lewis acids, e.g. AlCl$_3$,\(^{10}\) SbCl$_5$,\(^{11}\) and with transition-metal halides.\(^{12}\) In the presence of trace amounts of nucleophiles or reducing agents S$_2$N$_2$ dimerizes rapidly to S$_4$N$_4$,\(^{13}\) but this process is not well understood. It has been suggested that it may involve the initial formation of the radical anion [S$_2$N$_2$]$^-$.\(^{14}\) The interaction of the SOMO of the radical anion with the LUMO of a neutral S$_2$N$_2$ molecule would produce an adduct that could readily isomerize to the tetrascufur tetranitride radical anion [S$_4$N$_4$]$^-$.\(^{14}\) (Figure 1a). DFT calculations forecast that the $D_{2d}$ cage of neutral S$_4$N$_4$ unfolds to a $C_{2v}$ structure in which one of the transannular S⋯S bonds
opens up more than the other (Figure 1b) while the four nitrogen atoms remain equivalent, upon
addition of an electron to form the radical anion \([S_4N_4]^-\)\(^{15}\).

There have been no detailed studies of the electrochemistry of \(S_2N_2\),\(^{16}\) however DFT
calculations for the radical anion \([S_2N_2]^-\) predict a square planar (\(D_{2h}\)) structure with bond
lengths ca. 0.06 Å longer than those in the neutral molecule \(S_2N_2\), as expected for population of
the anti-bonding \(\pi\)-MO by a single electron.\(^{15}\) Numerous metal complexes of the chelating
acyclic dianion \([SNSN]^{2-}\) are known.\(^{17}\) Two methods for the in situ generation of this ligand are
(a) the reaction of \([S_4N_3]Cl\) with liquid ammonia\(^{18}\) or (b) deprotonation of the tetraimide \(S_4(NH)_4\)
in the presence of a base.\(^{19}\) Weak NMR resonances observed for reaction (a) have been
tentatively attributed to \([SNSN]^{2-}\).\(^{18}\) The protonated monoanion \([S_2N_2H^-]\) has been convincingly
shown to be formed by deprotonation of 99% \(^{15}\)N-enriched \(S_4(NH)_4\) with \(KNH_2\) in liquid
ammonia solutions using \(^{15}\)N NMR spectroscopy.\(^{20}\) However, attempts to detect the formation of
dianion \([S_2N_2]^{2-}\) via reaction (b) by this technique were unsuccessful.\(^{20a}\)

![Figure 1](attachment:image.png)

**Figure 1.** (a) Interaction between the LUMO of \(S_2N_2\) and the SOMO of \([S_2N_2]^-\) and (b) SOMO
of the \(C_{2v}\) symmetric \([S_4N_4]^-\) radical anion.

In recent publications we applied modern electrochemical and electron paramagnetic
resonance (EPR) methods to redox reactions of S,N heterocycles\(^{21}\) and binary sulfur nitrides.\(^{22}\) In
the latter contribution simultaneous electrochemical EPR (SEEPR) spectroscopy was used to
provide a comprehensive characterization of the \([S_4N_4]^+\) radical anion.\(^{22}\) Kinetic models for the
conversion of \(S_4N_4\) to \([S_3N_3]^-\) upon one-electron reduction and for the reverse process upon
oxidation were also developed. We have now carried out the first thorough voltammetric studies of $S_2N_2$ and have determined the identity of the redox products, including the neutral radical $[S_2N_2H]^-$ formed by proton transfer by using SEEPR spectroscopy in conjunction with DFT calculations. Digital modeling of CVs has been applied to determine the pathway for this transformation. Finally, we have identified the products of the chemical reduction of $S_2N_2$ with cobaltocene $Cp_2Co$, sodium naphthalenide $[Na][C_{10}H_8]$, and sodium benzophenone $[Na][Ph_2CO]$, and report details of the X-ray structure of $[Na(15\text{-crown-5})][S_3N_3]$.

**Experimental Section**

**Safety note!** In the pure form both $S_4N_4$ and $S_2N_2$ are friction-sensitive explosives. The latter has been reported to spontaneously detonate above 30° C, although we have not encountered such instability. All the appropriate safety precautions must be observed in their handling.

**General Procedures.** The reactions with $S_2N_2$ and the manipulations of products were performed under an argon atmosphere by using standard Schlenk techniques or an inert atmosphere glove box. The starting materials $S_4N_4$ and $S_2N_2$ were prepared by following the literature methods, and they were purified by recrystallization from CH$_3$CN and Et$_2$O, respectively. Isotopically labeled $S_2^{15}N_2$ was prepared from 99% $S_4^{15}N_4$. The solvents Et$_2$O and THF were dried by distillation over Na/benzophenone, and the solvents CH$_2$Cl$_2$ and CH$_3$CN were distilled from CaH$_2$ under an argon atmosphere prior to use. Electrochemical grade tetrabutylammonium hexafluorophosphate $[^nBu_4N][PF_6]$ (Fluka) was used as the supporting electrolyte and was kept in a desiccator prior to use. Ferrocene (Fc) was sublimed prior to use.

**Electrochemical Procedures.** Voltammetry, SEEPR, digital simulations and bulk electrolysis experiments were conducted as in our previous study using the same apparatus, electrodes, etc. except as noted below. Voltammograms were obtained at 20 ± 2 °C in CH$_3$CN,
CH$_2$Cl$_2$, and THF solutions containing 0.1 M, 0.4 M, and 0.4 M [$^n$Bu$_4$N][PF$_6$], respectively, as the supporting electrolyte. CV and RDE measurements were performed with a Bio-Analytical Systems (BASi) CV-50 computer-controlled potentiostat in conjunction with a PINE Model AFMSRXE Modulated Speed Rotator. The potentials for S$_2$N$_2$ are reported vs. the operative formal potential, $E^{0/0}_{Fe^{0+/+}}$, for the Fe$^{0+/+}$ redox couple, which was used as an internal standard, and literature data reported against SCE in CH$_3$CN/$[^n$Bu$_4$N][PF$_6$] have been converted to the Fc scale by subtraction of 0.38 V.\textsuperscript{25} In situ EPR experiments (Bruker EMX 113 spectrometer, 9.8 GHz) were conducted on CH$_2$Cl$_2$ solutions of S$_2$N$_2$ at temperatures between −90 and −20 °C using the miniature solution cell described previously and in CH$_3$CN solutions down to −35 °C.\textsuperscript{21,22} Experimental $g$ values were determined with reference to external, solid DPPH (2.0037 ± 0.0002).\textsuperscript{26} Simulations of EPR spectra were performed with Bruker Simfonia (version 1.25) and WinSim (version 0.98, 2002) software.\textsuperscript{27} Computed cyclic voltammograms were produced using DigiElch software (www.elchsoft.de).\textsuperscript{28}

**Quantum Calculations.** DFT calculations undertaken for this study employed geometry optimization at the (U)B3LYP/6-31G(d,p) level. Harmonic vibrational frequencies were calculated for all optimized geometries to confirm that they are stationary points. Calculation of energies and EPR hyperfine splitting (hfs) constants used (U)B3LYP/6-311G++(3df,3pd). All calculations were performed with the Gaussian 03 suite of programs.\textsuperscript{29}

**X-ray Crystallography.** A yellow, plate-shaped crystal of [Na(15-crown-5)][S$_3$N$_3$] (1a) was coated with Paratone 8277 oil and mounted on a glass fiber. Diffraction data were collected on a Nonius KappaCCD diffractometer using monochromated MoK$_{\alpha}$ radiation ($\lambda = 0.71073$ Å) at −100 °C. The data were corrected for Lorentz and polarization effects, and an empirical absorption correction was applied to the net intensities. The structure was solved by direct
methods using SHELXS-97 and refined using SHELXL-97. After the full-matrix least-squares refinement of the non-hydrogen atoms with anisotropic thermal parameters, the hydrogen atoms were placed in calculated positions (C–H = 0.99 Å). The isotropic thermal parameters of the hydrogen atoms were fixed at 1.2 times to that of the corresponding carbon. In the final refinement the hydrogen atoms were riding on their respective carbon atoms. One of the oxygen atoms in the 15-crown-5 unit showed positional disorder with site occupancy factors of ca. 0.5:0.5 in the final refinement.

**Reaction of [Na][C\textsubscript{10}H\textsubscript{8}] and S\textsubscript{2}N\textsubscript{2}**. A mixture of S\textsubscript{2}N\textsubscript{2} (0.037 g, 0.40 mmol) and 15-crown-5 (0.176 g, 0.80 mmol) in 30 mL of THF was added to a dark green solution of [Na][C\textsubscript{10}H\textsubscript{8}] (0.121 g, 0.80 mmol) in 30 mL of THF at 23 °C ([Na][C\textsubscript{10}H\textsubscript{8}] was prepared by stirring 0.018 g of Na metal and 0.103 g of Naphthalene in THF at 60 °C for 15h). The dark red reaction mixture was stirred for 2 h. The solvent was evaporated under vacuum and the resulting red, tarry product was washed with 2 x 10 mL of Et\textsubscript{2}O giving a yellow powder (yield = 0.099 g, 97%, calculated as [Na(15-crown-5)][S\textsubscript{3}N\textsubscript{3}] (1a), see discussion). X-ray quality crystals of 1a were obtained from THF after 1 h at −25 °C.

**Reaction of [Na][Ph\textsubscript{2}CO] and S\textsubscript{2}N\textsubscript{2}**. A mixture of S\textsubscript{2}N\textsubscript{2} (0.037 g, 0.40 mmol) and 15-crown-5 (0.176 g, 0.80 mmol) in 30 mL of Et\textsubscript{2}O was added to a dark blue solution of [Na][Ph\textsubscript{2}CO] (0.164 g, 0.80 mmol) in 30 mL of Et\textsubscript{2}O at 23 °C ([Na][Ph\textsubscript{2}CO] was prepared by stirring 0.018 g of Na metal and 0.146 g of benzophenone in Et\textsubscript{2}O at 23 °C for 6h). The reaction mixture was stirred for 2 h. The solvent was evaporated under vacuum and the resulting orange-red, tarry product was washed with 2 x 10 mL of Et\textsubscript{2}O giving a yellow powder (yield = 0.079 g, 77%, calculated as [Na(15-crown-5)][S\textsubscript{3}N\textsubscript{3}] (1a), see discussion). X-ray quality crystals of 1a were obtained from THF after 1 h at −25 °C and identified by a unit cell determination.
**Reaction of Cp₂Co and S₂N₂.** A solution of Cp₂Co (0.151 g, 0.80 mmol) in 30 mL of THF was added to a solution of S₂N₂ (0.037 g, 0.40 mmol) in 40 mL of THF at −80 °C. The reaction mixture was allowed to reach room temperature in 2 h and it was stirred at 23 °C for 2 h. The resulting red powder was allowed to settle and the solvent was decanted via a cannula. The product was washed with 20 mL of THF followed by drying under a vacuum (yield = 0.078 g, 90% calculated as [Cp₂Co][S₃N₃] (1b), see discussion). X-ray quality crystals of 1b were obtained from CH₃CN after 24 h at −25°C.³³,³⁴

**Results and Discussion**

**Voltammetry of S₂N₂.** S₂N₂ was studied by cyclic voltammetry (CV) in CH₃CN, CH₂Cl₂, and THF over scan rates of 0.05 – 20 V s⁻¹ and temperatures of 20 ± 2 °C. Data for the CV responses of S₂N₂ in all three solvents are reported in Table 1 with potentials quoted relative to Fc⁰⁺. Representative CVs of S₂N₂ in CH₂Cl₂ at a scan rate of 0.2 V s⁻¹ are shown in Figure 2. Two irreversible reduction processes appear upon sweeping the potential in the cathodic direction from the rest potential (−0.8 V vs. Fc⁰⁺), designated as $E_{p}^{c₃}$ and $E_{p}^{c₄}$, which are separated by approximately 0.8 V in all three solvents. $E_{p}^{c₃}$ appears at −1.37 V in CH₂Cl₂, −1.40 V in CH₃CN, and −1.29 V in THF. Further discussion will focus on voltammetry in CH₂Cl₂/0.4 M [⁷Bu₄N][PF₆] which is the medium that we found to be the most versatile for this investigation; similar behavior occurred in the other two solvent/electrolyte systems. CVs obtained over an electrochemical window that was extended in the anodic direction show the absence of any additional processes when the initial sweep direction is anodic out to the solvent limit (Figure 2a), but the presence of a new redox process, $E_{p}^{a₁}$ at −0.31 V which has a return wave of lower peak current height, $E_{p}^{c₁}$ at −0.38 V ($\Delta E_{p} = 72$ mV) when the potential is first swept through the initial reduction process (Figure 2b). This indicates that the new process is likely to be the
oxidation of a species generated from the decomposition of the product of electrochemical reduction at $E_p^{c3}$. Comparison of the peak potentials in CH$_3$CN with the earlier report of Banister and Hauptman $^{16}$ suggests that similar redox peaks were observed in their study. However, our extensive experience with the voltammetry$^{22}$ of S$_4$N$_4$ and the [PPN]$^+$ and [Cp$_2$Co]$^+$ salts of [S$_3$N$_3$]$^-$ allowed us to recognize the $E_p^{1}$ peaks immediately as those belonging to the [S$_3$N$_3$]$^{−}$/0 couple based on similarities in both the potentials and the $I_p^{a1}/I_p^{c1}$ ratio rather than the previous assignment of this process to oxidation of S$_2$N$_2$. $^{16}$ As further confirmation of this assignment, a small peak designated as $E_p^{c2}$ at $−1.02$ V appears subsequent to cycling through the [S$_3$N$_3$]$^{−}$/0 couple; such a process was anticipated based on our previous results and is unquestionably due to a small amount of S$_4$N$_4$ formed from the oxidation of [S$_3$N$_3$]$^−$ (see ref. 22; this peak was reported previously at $−1.06$ V in CH$_2$Cl$_2$).

**Figure 2.** CVs (black lines) obtained on 2.0 mM solutions of S$_2$N$_2$ in CH$_2$Cl$_2$ containing 0.4 M [′′Bu$_4$N][PF$_6$] at a GC electrode at 20 °C, $ν = 0.2$ V s$^{−1}$ with the potential starting at $−0.8$ V: (a) sweeping in the positive direction and (b) sweeping in the negative direction. (c) Calculated CV (red line) resulting from input of the kinetic parameters listed in the text for the specific case of $ν = 0.2$ V s$^{−1}$ and $k_{f3} = 10^{8}$ M$^{−1}$ s$^{−1}$, sweeping the potential first in the negative direction starting at $−0.8$ V.
In view of the absence of $E_p^{c2}$ in the initial cathodic scan (Fig. 2a) we tentatively assign $E_p^{c3}$ to the formation of $[S_2N_2]^*$. The behavior of this voltammetric peak when investigated over the full range of scan rates shows strong evidence of distortions that we attribute to adsorption on the electrode surface. For example, after several sequential scans, we observe a crossing over of the anodic and cathodic traces; cleaning the electrode surface by polishing then restores the signals to those observed originally (e.g. Fig. 2a). Even scans on clean electrode surfaces show evidence of unusual peak shapes. This voltammetric indication that absorption seems to accompany $E_p^{c3}$ is relevant to the SEEPR results (vide infra).

**Table 1.** CV Data for $S_2N_2$ in CH$_3$CN, CH$_2$Cl$_2$, and THF and Calculated Enthalpies of Electron Attachment.$^a$

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<th>solvent</th>
<th>conc (mM)</th>
<th>$E_{pa}^{a1}$ (V)</th>
<th>$E_{pc}^{a1}$ (V)</th>
<th>$E_{pa}^{b}$ (V)</th>
<th>$E_{pc}^{c2}$ (V)</th>
<th>$E_{pc}^{c3}$ (V)</th>
<th>$E_{pc}^{c4}$ (V)</th>
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<td>CH$_2$Cl$_2$</td>
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<td>−0.38</td>
<td>−0.35</td>
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</tr>
<tr>
<td>THF</td>
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<td>−0.39</td>
<td>−0.34</td>
<td>−0.99</td>
<td>−1.29</td>
<td>−2.25</td>
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</tbody>
</table>

$^a$ Obtained at a GC electrode at $\nu = 0.2$ V s$^{-1}$ and $T = 20 \pm 2$ °C.$^b$ $E_n = \frac{E_{pa}^{a1} + E_{pc}^{c1}}{2} \approx E_{o/0}(1)$.

$^c$ As reported in Ref. 16. Converted from the SCE scale to Fe by subtraction of 0.38 V (see Ref. 25). $^d$ Reported as $E_{p/2}$, i.e. the potentials at half-peak-height. $^e$ Calculated at the (U)B3LYP/6-311G++(3df,2dp)// (U)B3LYP/6-31G(d,p) level in Gaussian 03 (gas phase).

CVs extended further in the cathodic direction (Figure 3) show evidence of continuous current flow beyond $E_p^{c3}$ with another identifiable peak ($E_p^{c4}$) at −2.2 V that has a peak current ~2/3 that of $E_p^{c3}$. While $E_p^{c4}$ might be due to generation of the dianion $[S_2N_2]^{2-}$, its identification by voltammetric methods is unreliable. We chose instead to study this second process using a chemical approach (vide infra). Significantly, $E_p^{c4}$ is not the same process as that observed from CV studies of solutions containing bulk $S_4N_4$, for which the (irreversible) second reduction peak is observed at the more anodic potential of −1.6 V vs. Fe$^{0/+}$.22
Figure 3. CV of a 2.0 mM soln of S$_2$N$_2$ in CH$_2$Cl$_2$ (0.4 M [$_n$Bu$_4$N][PF$_6$]), v = 0.2 V s$^{-1}$, T = 20 °C, with the potential sweep starting at −0.8 V and extending further negative to include the second reduction process at −2.3 V (labeled $E_p^{c4}$).

**Determination of the Diffusion Coefficient of S$_2$N$_2$.** Both CV and RDE measurements on solutions of S$_2$N$_2$ were undertaken in order to determine the diffusion coefficient $D$. The same approach was used as for the determination of $D$ for S$_4$N$_4$ and [S$_3$N$_3$]$^-$.$^{22}$ CV measurements provided estimates of $D$ of 8.16 x 10$^{-6}$ cm$^2$ s$^{-1}$ in CH$_2$Cl$_2$ and 7.65 x 10$^{-6}$ cm$^2$ s$^{-1}$ in CH$_3$CN. RDE measurements employed rotation rates between 1000 – 2250 rpm over a potential range of −0.6 to −2.2 V to a 2.1 mM solution of S$_2$N$_2$ in CH$_2$Cl$_2$. The Levich current ($I_L$) values were negatively affected by the electrochemical generation of [S$_4$N$_4$]$^-$ at rotation rates greater than 1000 rpm (Fig. 4a). Therefore, all calculations had to take account of the number of moles of S$_2$N$_2$ converted to [S$_4$N$_4$]$^-$ during the course of RDE measurements in order to obtain accurate $I_L$ values. A plot of the reciprocal Levich current ($I_L^{-1}$) vs. the reciprocal square root of the rotational velocity ($\omega^{-1/2}$) yielded a straight line (Fig. 4b) with a slope = (0.62$nFAC_v^{1/6}D^{2/3}$)$^{-1}$. Solving for $D$ yielded a value of 8.73 x 10$^{-6}$ cm$^2$ s$^{-1}$, in good agreement with the values obtained from CV measurements using the Randles-Sevcik equation.$^{35}$ Values for $D$ based on both RDE and CV measurements are listed in Table 2. A best estimate for $D$ used in modeling the voltammetric response of S$_2$N$_2$ (vide infra) was taken to be 9.13 x 10$^{-6}$ cm$^2$ s$^{-1}$, the average of the values obtained by both CV and RDE methods in CH$_2$Cl$_2$. (Note that the diffusion coefficient of [S$_3$N$_3$]$^-$ is smaller by ~50%, which accounts for the discrepancy in the peak currents ($I_p^{al}$ vs.
1. \( I_p^c \) for the two species shown in Figure 2; for further background on such phenomena see Ref 22.

![Figure 4](image)

Figure 4. (a) RDE measurements on a 2.1 mM solution of \( \text{S}_2\text{N}_2 \) in \( \text{CH}_2\text{Cl}_2 \) with 0.4 M \([n\text{Bu}_4\text{N}][\text{PF}_6]\) at a 0.5 mm GC electrode, rotation rates = 1000 – 2250 RPM, \( \nu = 0.010 \text{ V s}^{-1} \); and (b) Koutecky-Levich \( ^{35} \) plot for the reduction of \( \text{S}_2\text{N}_2 \) (\( R^2 = 0.998 \)) after correction for \( \text{S}_4\text{N}_4 \) formation (see text).

Table 2. Diffusion Coefficient Values for \( \text{S}_2\text{N}_2 \) in \( \text{CH}_2\text{Cl}_2 \) at \( T = 21 \pm 2 \) °C.

<table>
<thead>
<tr>
<th>Method</th>
<th>Electrode</th>
<th>RPM</th>
<th>( D ) (( \text{S}_2\text{N}_2 )) / 10\text{(^{-6})} cm(^2) s(^{-1})</th>
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<td>RDE(^a)</td>
<td>( \text{CH}_2\text{Cl}_2 )</td>
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<tr>
<td></td>
<td>( \text{CH}_2\text{Cl}_2 )</td>
<td>—</td>
<td></td>
</tr>
<tr>
<td></td>
<td>CV</td>
<td>GC</td>
<td></td>
</tr>
<tr>
<td></td>
<td>( \text{CH}_3\text{CN} )(^e)</td>
<td>GC</td>
<td></td>
</tr>
</tbody>
</table>

\(^a\) 0.4 M \([n\text{Bu}_4\text{N}][\text{PF}_6]\), \( \nu = 0.01 \text{ V s}^{-1} \). \(^b\) 2.13 mM. \(^c\) 0.4 M \([n\text{Bu}_4\text{N}][\text{PF}_6]\), \( \nu = 0.1 – 1.0 \text{ V s}^{-1} \). \(^d\) 2.00 mM. \(^e\) 0.1 M \([n\text{Bu}_4\text{N}][\text{PF}_6]\), \( \nu = 0.1 – 0.5 \text{ V s}^{-1} \). \(^f\) 1.50 mM.
Digital modeling of CVs of S$_2$N$_2$. In order to gain a better understanding of both the decay of [S$_2$N$_2$]$^-$ and the interconversion between S$_2$N$_2$, S$_4$N$_4$, and [S$_3$N$_3$]$^-$, we have built a model of the voltammetric behavior of S$_2$N$_2$ using DigiElch software. Starting from the robust square scheme previously established from the excellent curve fitting simulations of CVs for the S$_4$N$_4$ ↔ [S$_3$N$_3$]$^-$ interconversion under the same conditions, we developed a ladder scheme in which the common parameters have the same meaning as in Ref. 22 (Scheme 1). For the digital models, the scan rate $\nu$ was varied between 0.1 – 10 V s$^{-1}$ and the shapes of the simulated CVs were compared to those measured experimentally. In this study it proved impossible to perform computerized line-fitting of the model to the experimental curves, most likely due to the abnormal peak shapes caused by adsorption (vide supra). However, reasonable visual fits were obtained over the full range of scan speeds (Figure 2c). Our model assumes a coupling reaction between the initially formed [S$_2$N$_2$]$^-$ with excess S$_2$N$_2$ in solution to form the known [S$_4$N$_4$]$^-$ (an EC mechanism with a second-order C step). In view of the highly irreversible nature of the [S$_2$N$_2$]$^{\circ}$/0 redox couple, the value for $K_{eq3}$ was set to be very large ($10^6$), rendering the C step irreversible regardless of scan rate. Values for $k_{f3}$ which resulted in the greatest similarity between experimental and modeled CVs were $\geq 10^8$ M$^{-1}$ s$^{-1}$, which converts to a half-life of $\leq 4 \times 10^{-6}$ seconds. Another feature of our model is that at faster scan rates ($> 1$ V s$^{-1}$), a return current peak occurs for the [S$_4$N$_4$]$^{\circ}$/0 redox couple which is not evident in the experimental CVs at similar scan rates. In order for the models to follow the experimental behavior over the full range of scan rates, $k_{f1}$ values needed to be at least 25 $\times$ greater than was observed previously, i.e. the half-life of [S$_4$N$_4$]$^-$ when in the presence of [S$_2$N$_2$]$^{\circ}$/0 appears to be significantly shorter than in bulk S$_4$N$_4$ solutions. Final values for the adjustable parameters are: $k_{s1} = 0.03$ cm s$^{-1}$; $k_{s2} = 0.02$ cm s$^{-1}$; $k_{s3} = 0.02$ cm s$^{-1}$; $k_{f1} = 50$ s$^{-1}$; $k_{f2} = 0.4$ s$^{-1}$; $k_{f3} \geq 10^8$ M$^{-1}$ s$^{-1}$. This model thus
predicts that the major product of the reduction of S$_2$N$_2$ in an aprotic environment will be [S$_3$N$_3$]$^-$ via the intermediate [S$_4$N$_4$]$^-$•. This is fully supported by both EPR spectroscopic evidence and the results of chemical reduction, in which *quantitative* production of [S$_3$N$_3$]$^-$ salts is obtained (vide infra).

**Scheme 1.** The proposed "ladder" scheme interrelating S$_2$N$_2$, [S$_4$N$_4$]$^-$ and [S$_3$N$_3$]$^-$.

\[
\begin{align*}
S_2N_2 + 1e^-, k_{s3} & \rightarrow [S_2N_2]^+ + S_2N_2 \\
S_4N_4 + 1e^-, k_{s4} & \rightarrow [S_4N_4]^+ \\
S_3N_3^+ - 1e^-, k_{s2} & \rightarrow [S_3N_3]^-
\end{align*}
\]

**EPR Spectroscopy.** In order to characterize the nature of the redox process occurring at $E_{p}^{c3}$, tentatively ascribed to a one-electron reduction of S$_2$N$_2$, we conducted SEEPR experiments on solutions of S$_2$N$_2$ in CH$_2$Cl$_2$, CH$_3$CN, and THF containing [°Bu$_4$N][PF$_6$] as the supporting electrolyte. For these experiments, our newly-developed low temperature solution electrochemical cell was used exclusively.\textsuperscript{21,22} Temperatures were varied between $-100$ and $-20^\circ$C in CH$_2$Cl$_2$, $-35$ and $0^\circ$C in CH$_3$CN, and $-100$ and $0^\circ$C in THF. In practice, we found that in these systems accurate control of the electrolysis potentials was difficult. There is a strong tendency for over-reduction accompanied by adsorption of paramagnetic species exhibiting anisotropic EPR spectra onto the electrode surface (Figure 5). Significantly, both the isotropic (solution phase) and anisotropic (adsorbed) EPR signals decay rapidly even at low temperatures.
Figure 5. (a) Experimental first derivative EPR spectrum obtained during in situ reductive electrolysis at −1.4 V of S₂N₂ (82 s single scan) at a gold foil electrode at −70 °C in CH₂Cl₂ (0.4 M [⁸⁹Bu₄N][PF₆]), 20.6 mM solution, modulation amplitude = 0.15 mT. (b) Spectrum obtained under identical conditions at −1.8 V. The g value of the apparent nine-line signal between 334.5 and 336 mT (green asterisks) is 2.0039(1). (c) Spectrum obtained at an intermediate potential for 99% S₂¹⁵N₂. The red line is a simulation of the broadened three-line signal (red asterisks) for two equivalent ¹⁵N nuclei with a(¹⁵N) = 0.20 mT. Note that a little of the solution-phase [S₄¹⁵N₄]⁻ persist at the r.h.s. of this spectrum (blue asterisks).

The most reliable results were obtained by using thin platinum or gold foil working electrodes and a Teflon-insulated silver wire reference electrode in CH₂Cl₂ solution at potentials that corresponded closely to $E_p^{c3}$ as determined by room temperature voltammetry studies. When electrolysis is conducted at −1.4 V vs. Fe⁰/⁺ at temperatures between −90°C and −40°C, strong signals from [S₄N₄]⁻ are obtained (Figure 5a; nine-line, $g = 2.0008(1)$, $a(N_{1-4}) = 0.115$ mT). This spectroscopic result lends support to the mechanism whereby [S₄N₄]⁻ is a direct and rapid product of the reduction of S₂N₂ (Scheme 1). Starting at −1.5 V and continuing to more negative potentials, the solution-phase spectrum of [S₄N₄]⁻ begins to be replaced by an anisotropic spectrum, likely from species adsorbed onto the electrode surface. At −1.8 V, no solution-phase signal remains (Figure 5b), but there appears to be a nine-line pattern at the right-hand side (green asterisks) that resembles solution-phase [S₄N₄]⁻; however, the g value is different and the lines are broader than observed in solution at the same temperature. Similarly, the pattern
between 333 and 334 mT (red asterisks) can be fit to a 1:2:3:2:1 quintet with hfs of 0.14 mT which could belong to $[S_2N_2]^-$. When identical experiments were performed with 99% isotopically labeled $S_2^{15}N_2$ (Figure 5c), the replacement of $^{14}N$ (I = 1) by $^{15}N$ (I = ½) results in the expected change in the fine structure of the EPR spectra from nine-line (green) and five-line (red) signals (Figure 5b) to five-line (green) and three-line (red) patterns. The ratio $a^{15}N:a^{14}N$ is ca. 1.4, as expected from the relative values of the nuclear gyromagnetic ratios. However, all the signals are shifted, albeit by a small amount, which is indicative of a species or species with anisotropic $g$ tensors. In view of this complexity, assignment of these signals to one or more specific sulfur-nitrogen radicals is not warranted. Note that similar signals with rather small $g$ anisotropy have been reported for intermediates in the preparation of the (SN)$_x$ polymer.

![Figure 6](image_url)

**Figure 6.** (a) Experimental EPR spectrum (black line) obtained during in situ reductive electrolysis of 1.5 mM $S_2N_2$ at −1.5 V (164 s single scan) at a platinum foil electrode in CH$_3$CN (0.1 M $[^{n}Bu_4N][PF_6]$) at −20 °C after noise reduction, and (b) simulation (red line), with $a^{^{14}N_1}=1.12$ mT, $a^{^{14}N_2}=0.65$ mT, with $a^{(1)H}=0.60$ mT, $g$ value $= 2.0136(1)$, mod. amp. $= 0.2$ mT.

In CH$_3$CN reductive electrolysis at −1.5 V at temperatures between −20 °C and −35 °C produced weak eight-line signals (Figure 6a). This spectrum is readily simulated by assuming coupling to two inequivalent nitrogen and single hydrogen atoms (Figure 6b), attributable to the neutral radical $[S_2N_2H]^+$ through a proton transfer. Significantly, the hfs values for N2 and H show significant variation with changes in temperature and environment (Table 3) as expected.
for a non-planar radical. Since adventitious moisture is the most likely source of protons for forming $[S_2N_2H]^\cdot$, this radical was generated deliberately by electrolysis of "wet" 2:1 mixtures of CH$_2$Cl$_2$/CH$_3$CN (where the latter was degassed but not dried). The use of lower temperatures resulted in much stronger signals for $[S_2N_2H]^\cdot$ than obtained in CH$_3$CN (Table 3).

In an experiment designed to produce $[S_2N_2H]^\cdot$ via in situ electrochemical reduction of the protonated cation $[S_2N_2H]^+$ electrolysis of a solution of $S_2N_2$ in CH$_2$Cl$_2$/0.4 M [${^8}$Bu$_4$N][PF$_6$] containing one equivalent of HBF$_4$·2OEt$_2$ was performed. At low temperatures, EPR spectra obtained from this solution resemble the signal for the absorbed species depicted in Fig. 5b, but at higher temperatures the signals attributed to $[S_2N_2H]^\cdot$ grow in steadily until this is the dominant signal at $-40^\circ$C (Fig. 7a,b). Further warming to $-20 ^\circ$C resulted, after several minutes, in the generation of a persistent five-line signal (Fig. 7c,d) that is readily attributed to $[S_3N_2]^\cdot$ ($g = 2.0111, a(N) = 0.31$ mT). This stable free radical is known to result from the treatment of either $S_2N_2$ or $S_4N_4$ solutions with strong acid. By contrast, the spectra obtained in wet CH$_3$CN at $-20 ^\circ$C show that the $[S_2N_2H]^\cdot$ signal in absence of strong acid decays with no indication of the formation of other EPR-active species. The EPR characterization of $[S_2N_2H]^\cdot$ provides strong support for the production of $[S_2N_2]^\cdot$ in the one-electron electrochemical reduction of $S_2N_2$ associated with the voltammetric process at $E_p$. The failure to observe $[S_2N_2]^\cdot$ directly is attributed to the short lifetime of this radical in the presence of excess $S_2N_2$ as predicted by the voltammetric model (Scheme 1) and evidenced by spectroscopic observation of rapid and copious production of $[S_4N_4]^\cdot$ from the main reaction.
Figure 7. Experimental EPR spectra from single scans obtained during in situ reductive electrolysis of 1.5 mM $S_2N_2$ at $-1.5$ V (164 s single scan) at a platinum foil electrode in CH$_2$Cl$_2$ (0.4 M $[^\text{a}]{\text{Bu}_4N}\text{[PF}_6\text{]}$) containing HBF$_4$·2OEt$_2$: (a) at $-40$ °C (black line) and (b) the simulation of this spectrum (red line); note that in the simulation the small contribution by $[S_4N_4]^-\cdot$ (14% by area) has been omitted to more clearly show the high-field line component. (c) At $-20$ °C after about 10 minutes (black line). (d) Simulation (red line) of the spectrum in (c).

Table 3. Experimental and calculated EPR Data for $[S_2N_2]^-\cdot$ and $[S_2N_3H]^\cdot$

<table>
<thead>
<tr>
<th>Species/condition</th>
<th>Temp. (°C)</th>
<th>$a(N_1)$ (mT)</th>
<th>$a(N_2)$ (mT)</th>
<th>$a(H)$ (mT)</th>
<th>LW (mT)</th>
<th>g value</th>
</tr>
</thead>
<tbody>
<tr>
<td>$[S_2N_2]^-\cdot$/Calculated/gas phase $^a$</td>
<td>—</td>
<td>0.52</td>
<td>0.52</td>
<td>—</td>
<td>—</td>
<td>—</td>
</tr>
<tr>
<td>$[S_2N_3H]^\cdot$/CH$_3$CN $^b$</td>
<td>$-20$</td>
<td>1.12</td>
<td>0.65</td>
<td>0.60</td>
<td>0.090</td>
<td>2.0136(1) $^e$</td>
</tr>
<tr>
<td>$[S_2N_3H]^\cdot$/CH$_3$CN $^b$</td>
<td>$-30$</td>
<td>1.12</td>
<td>0.64</td>
<td>0.64</td>
<td>0.093</td>
<td>2.0136(1) $^e$</td>
</tr>
<tr>
<td>$[S_2N_3H]^\cdot$/CH$_3$CN $^b$</td>
<td>$-35$</td>
<td>1.12</td>
<td>0.63</td>
<td>0.69</td>
<td>0.090</td>
<td>2.0136(1) $^e$</td>
</tr>
<tr>
<td>$[S_2N_3H]^\cdot$/mixture $^c$</td>
<td>$-20$</td>
<td>1.11</td>
<td>0.69</td>
<td>0.71</td>
<td>0.078</td>
<td>2.0137(1) $^e$</td>
</tr>
<tr>
<td>$[S_2N_3H]^\cdot$/mixture $^c$</td>
<td>$-50$</td>
<td>1.10</td>
<td>0.70</td>
<td>0.75</td>
<td>0.081</td>
<td>2.0137(1) $^e$</td>
</tr>
<tr>
<td>$[S_2N_3H]^\cdot$/acid $^d$</td>
<td>$-40$</td>
<td>1.11</td>
<td>0.68</td>
<td>0.73</td>
<td>0.056</td>
<td>2.0142(1) $^e$</td>
</tr>
<tr>
<td>$[S_2N_3H]^\cdot$/Calculated/gas phase $^a$</td>
<td>—</td>
<td>0.88</td>
<td>$-0.10$</td>
<td>1.17</td>
<td>—</td>
<td>—</td>
</tr>
</tbody>
</table>

$^a$ Computed at the UB3LYP/6-311G++(3df,3pd)/UB3LYP/6-31G(d,p) level with Gaussian03 (see text). $^b$ Containing 0.1 M $[^\text{a}]{\text{Bu}_4N}\text{[PF}_6\text{]}$. $^c$ 2/3 CH$_2$Cl$_2$, 1/3 wet CH$_3$CN containing 0.4 M $[^\text{a}]{\text{Bu}_4N}\text{[PF}_6\text{]}$. $^d$ CH$_2$Cl$_2$ containing 1 equiv. HBF$_4$·2OEt$_2$ and 0.4 M $[^\text{a}]{\text{Bu}_4N}\text{[PF}_6\text{]}$. $^e$ Referenced internally to $[S_4N_4]^-\cdot$ at 2.0008(1) (Ref. 22).
**DFT Calculations.** The \([S_2N_2H]\) neutral radical is calculated at the UB3LYP/6-31G(d,p) level of theory to adopt a slightly butterfly ring geometry in the gas phase (dihedral angle between SNS planes of 6.9°) with the hydrogen atom lying out of the ring plane (Figure 8; see figure and legend for metric parameters). The calculated hfs constants of \(a(N_1) = 0.88\) mT, \(a(N_2) = -0.10\) mT, and \(a(H) = 1.17\) mT are in reasonable agreement with the experimental values (Table 3). The geometry of \([S_2N_2]^-\) at this same level of theory is close to that previously calculated using a specialized basis set.\(^{15}\)

![Figure 8. Optimized gas-phase UB3LYP/6-31G(d,p) structure of \([S_2N_2H]\); \(\angle S1-N1-S2 94.7°\), \(\angle N1-S1-N2 87.8°\), \(\angle S2-N2-S1 89.4°\), \(\angle S1-N2-H 108.9°\), \(\angle N1-N2-H 120.4°\), S1N1S2-S1N2S2 dihedral 6.9°.](image)

DFT calculations were also undertaken to determine the enthalpies of electron attachment of \(S_2N_2\), \(S_3N_3\) and \(S_4N_4\) in the gas phase in order to rationalize the measured redox potentials in solution (Table 1). The calculations show that in the gas phase, \(\Delta H_{EA} S_3N_3 > S_4N_4 > S_2N_2\), i.e. the same trend as the redox potentials in solution. An important consequence is that the solution electron transfer reaction (Eq. 1) is predicted to be non-spontaneous.\(^{36}\)

\[
[S_4N_4]^- + S_2N_2 \rightarrow S_4N_4 + [S_2N_2]^- \quad E^\circ = -0.35 \text{ V} \quad (1)
\]

This reaction would seem to be a necessary component of the proposed catalytic cycle for the rapid conversion of solutions of \(S_2N_2\) to \(S_4N_4\) with just a small amount of reducing agent.\(^{14,44}\)
Chemical Reduction of $\text{S}_2\text{N}_2$: Synthesis and Crystal Structure of $[\text{Na(15-crown-5)}][\text{S}_3\text{N}_3]$. The electrochemical experiments described above were complemented by investigations of the chemical reduction of $\text{S}_2\text{N}_2$ with the primary objective of determining the possible production of the dianion $[\text{S}_2\text{N}_2]^{2-}$, a potential alternative source of the short-lived radical anion $[\text{S}_2\text{N}_2]^{-}$ via one-electron oxidation. These reactions were conducted with three reagents that display increasing strength for reduction (formal potential vs. $\text{Fe}^{0+/+}$): $\text{Cp}_2\text{Co}$ (−1.33 V in $\text{CH}_2\text{Cl}_2$), $[\text{Ph}_2\text{CO}]^{-}$ (−2.30 V in THF) and $[\text{C}_{10}\text{H}_8]^{-}$ (−3.10 V in THF). The formal potentials of the first two reducing agents are close to the measured values for $E_p^{c3}$ and $E_p^{c4}$, respectively, while the naphthalenide radical anion is a significantly stronger reducing agent than $E_p^{c4}$ (vide supra).

The reactions were performed in a 2:1 molar ratio by using the sodium salts of radical anions ($[\text{Ph}_2\text{CO}]^{-}$ and $[\text{C}_{10}\text{H}_8]^{-}$) in the presence of 15-crown-5 and cobaltocene $\text{Cp}_2\text{Co}$. The products were identified by X-ray crystallography, which revealed that in all three reactions, despite the various strengths of the reducing agents, formation of the known $[\text{S}_3\text{N}_3]^{-}$ monoanion as $[\text{Cp}_2\text{Co}]^{+}$ and $[\text{Na(15-crown-5)}]^{+}$ salts in excellent yields occurred (Scheme 2). The voltammetry and EPR results revealed the rapid conversion of $[\text{S}_2\text{N}_2]^{-}$ to $[\text{S}_4\text{N}_4]^{-}$ and subsequently, $[\text{S}_3\text{N}_3]^{-}$. Thus, the outcome of the chemical reductions is consistent with these observations made for the electrochemical reduction of $\text{S}_2\text{N}_2$, even in the presence of an excess of reducing agent. Consistent with previous attempts to generate the $[\text{S}_2\text{N}_2]^{2-}$ dianion by deprotonation of $[\text{S}_2\text{N}_2\text{H}]^{-}$, no evidence for the formation of the $[\text{S}_2\text{N}_2]^{2-}$ dianion was garnered from either the chemical or electrochemical reductions.
Scheme 2. Formation of the $[S_3N_3]^{-}$ anion from chemical reduction of $S_2N_2$

The crystal structure of $[\text{Cp}_2\text{Co}][S_3N_3]$, obtained from the reduction of $S_4N_4$, has been reported previously. However, $[\text{Na}(15\text{-crown-5})][S_3N_3]$ (1a) is the first structurally characterized alkali-metal salt of $[S_3N_3]^{-}$. A large, non-coordinating cation is usually required to stabilize the six-membered ring; simple alkali-metal salts of $[S_3N_3]^{-}$ are explosive in the solid state. As illustrated in Figure 9, the crystal structure of 1a shows a $C_s$-symmetry with the O1, Na1, N1 and S2 atoms in the mirror plane. The Na$^+$-cation, coordinated by the crown ether, displays a fairly strong Na···N interaction of 2.388(5) Å (cf. for example Na···N of 2.375(3) and 2.393(5) Å in $[\text{Na}(18\text{-crown-6})(\text{THF})_2][\text{NCS}]$ and $[\text{Na}(15\text{-crown-5})][\text{NCS}]$, respectively). The most germane comparison with 1a, perhaps, is the salt $[\text{Na}(15\text{-crown-5})][\text{SN}_2\text{C}_2\text{N}_2\text{S}]$ containing a heterocyclic C,N,S radical anion, in which the Na—N distances are 2.511 and 2.837 Å. The larger values in the latter salt are likely attributable to its infinite chain arrangement of N···Na$^+$···N···Na$^+$ contacts, whereas the $[S_3N_3]^{-}$ salt has a single N···Na$^+$ contact on one side of the crown. The six-membered $[S_3N_3]^{-}$ ring in 1a shows a slight, albeit almost negligible, distortion towards a chair conformation with the maximum deviation from planarity of 0.023(3) Å. All the bond lengths and angles at S and N atoms, respectively, are equal within experimental

...
errors (Table 4). The bond parameters are also comparable to the average values of 1.61 Å (S–N), 124° (∠SNS) and 116° (∠NSN) observed in the previously reported salts of the [S$_3$N$_3$]$^-$ anion (Table 4). The only notable intermolecular close contacts in 1a are the two N···H connections of 2.700(4) and 2.703(4) Å involving the N2 (and by symmetry N2$^9$) atom and the adjacent crown ether (cf. sum of the van der Waals radii for nitrogen and hydrogen 2.74 Å).$^{48}$

![Figure 9](image_url)

**Figure 9.** Crystal structure of 1a with the atomic numbering scheme. Hydrogen atoms have been omitted for clarity. $^a$ Symmetry operation: x, 0.5-y, z.

Comparison of the solid-state structures of [S$_3$N$_3$]$^-$ salts demonstrates that the subtle distortion in the six-membered ring is surprisingly independent of the cation-anion interactions. The diversity in the anion bond parameters in 1c and 1d ([(PhCN$_2$S$_2$)$_2$Cl]$^+$ and [PhCN$_2$S$_2$]$^+$ salts, respectively) was attributed mainly to interionic S···S interactions,$^{49}$ and in 1e ([PPN]$^+$) to the strong hydrogen-bonding with the MeOH solvate.$^{22}$ On the other hand, there are no close contacts, even to the hydrogen atoms of the cations, in the [$^4$Bu$_4$N]$^+$ (1f)$^{50}$ and [Ph$_4$P]$^+$ (1g)$^{19b}$ salts, but significant distortions in the S-N bond lengths are also observed in these salts. Curiously, structures 1a and 1b display no significant variation in the bond parameters, despite the presence of strong interionic Na···N(1a) and N···H interactions in 1b. We note that, unlike 1c-1g, the [S$_3$N$_3$]$^-$ anion in 1a and 1b exhibits crystallographic mirror symmetry. In summary,
the 10 \( \pi \)-electron \([\text{S}_3\text{N}_3]^-\) anion\(^8c\) exhibits minor distortions from regular \(D_{3h}\) symmetry in the solid state that are not consistently determined by anion-cation interactions.

**Table 4.** Selected bond lengths (Å) and bond angles (°) in \(1\alpha\) and other \([\text{S}_3\text{N}_3]^-\) salts.\(^a,b\)

<table>
<thead>
<tr>
<th>Bond Length or Angle</th>
<th>(1\alpha)</th>
<th>(1\beta^d)</th>
<th>(1\gamma)</th>
<th>(1\delta)</th>
<th>(1\epsilon)</th>
<th>(1\zeta)</th>
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<tr>
<td>N1-S1</td>
<td>1.617(2)</td>
<td>1.623</td>
<td>1.587(4)</td>
<td>1.623(4)</td>
<td>1.615(5)</td>
<td>1.604(12)</td>
<td>1.641(9)</td>
</tr>
<tr>
<td>N1-S1(^c)</td>
<td>1.617(2)</td>
<td>1.623</td>
<td>1.603(3)</td>
<td>1.616(4)</td>
<td>1.630(5)</td>
<td>1.609(12)</td>
<td>1.601(9)</td>
</tr>
<tr>
<td>N2-S1</td>
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<td>1.625</td>
<td>1.597(4)</td>
<td>1.564(4)</td>
<td>1.616(6)</td>
<td>1.626(12)</td>
<td>1.566(8)</td>
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<tr>
<td>N2-S2</td>
<td>1.605(4)</td>
<td>1.618</td>
<td>1.615(3)</td>
<td>1.634(4)</td>
<td>1.646(5)</td>
<td>1.584(12)</td>
<td>1.628(8)</td>
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<tr>
<td>N2(^c)-S2</td>
<td>1.605(4)</td>
<td>1.618</td>
<td>1.590(4)</td>
<td>1.630(4)</td>
<td>1.646(5)</td>
<td>1.599(12)</td>
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<tr>
<td>N2(^c)-S1(^c)</td>
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<td>1.625</td>
<td>1.593(4)</td>
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<td>1.609(7)</td>
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<td>Na1...Nl</td>
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<tr>
<td>S1-N1-S1(^c)</td>
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<td>123.7</td>
<td>124.9(3)</td>
<td>122.6(2)</td>
<td>123.8(3)</td>
<td>122.9(8)</td>
<td>122.6(5)</td>
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<td>S1-N2-S2</td>
<td>124.1(2)</td>
<td>123.4</td>
<td>122.9(2)</td>
<td>125.2(3)</td>
<td>124.2(3)</td>
<td>122.6(8)</td>
<td>124.3(5)</td>
</tr>
<tr>
<td>S2-N2(^c)-S1(^c)</td>
<td>124.1(2)</td>
<td>123.4</td>
<td>124.9(2)</td>
<td>125.5(3)</td>
<td>123.2(4)</td>
<td>124.2(8)</td>
<td>124.4(5)</td>
</tr>
<tr>
<td>N1-S1-N2</td>
<td>115.3(2)</td>
<td>116.7</td>
<td>116.2(2)</td>
<td>116.7(2)</td>
<td>117.5(3)</td>
<td>116.5(7)</td>
<td>116.6(4)</td>
</tr>
<tr>
<td>N2-S2-N2(^c)</td>
<td>116.4(3)</td>
<td>116.1</td>
<td>116.1(2)</td>
<td>113.4(2)</td>
<td>115.4(3)</td>
<td>117.0(7)</td>
<td>115.6(4)</td>
</tr>
<tr>
<td>N2(^c)-S1(^c)-N1</td>
<td>115.3(2)</td>
<td>116.7</td>
<td>114.8(2)</td>
<td>116.1(2)</td>
<td>115.5(3)</td>
<td>116.5(7)</td>
<td>116.4(4)</td>
</tr>
</tbody>
</table>

\(^a\) Atomic numbering scheme of \(1\beta\)\(^d\)-\(1\gamma\) has been changed to match that in \(1\alpha\). \(^b\) \((1\alpha) [\text{Na}(15\text{-}

Conclusions

Both electrochemical and chemical reduction of \(\text{S}_2\text{N}_2\) produce the \([\text{S}_3\text{N}_3]^-\) anion. The former process can be modeled by a mechanism in which the initially formed \([\text{S}_2\text{N}_2]^-\) radical anion reacts with excess \(\text{S}_2\text{N}_2\) to afford \([\text{S}_4\text{N}_4]^-\), verifying a literature prediction.\(^14\) The latter species is known to decay rapidly to give \([\text{S}_3\text{N}_3]^-\).\(^22\) In the presence of a proton donor EPR spectra indicate that trapping of \([\text{S}_2\text{N}_2]^-\) occurs to form the neutral radical \([\text{S}_2\text{N}_2\text{H}^-]\), which was
identified by simulation of the EPR spectrum as supported by DFT calculations. Consistent with the results of previous chemical approaches, investigations of the electrochemical reduction of S$_2$N$_2$ provided no evidence for the formation of the dianion [S$_2$N$_2$]$^{2-}$.

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Supporting information available: X-ray crystallographic file in CIF format. This material is available free of charge via the Internet at http://pubs.acs.org.

References and Notes


(16) To our knowledge, there is only one prior literature report of a voltammetric study of S$_2$N$_2$. In a paper that deals with the electrochemical reduction of the [S$_5$N$_3$]$^+$ cation it is reported that $E_{p/2}^{red} = -0.85$ V, $E_{p/2}^{ox} = 0.1$ V (vs. aq. SCE); S$_2$N$_2$ (ca. 10$^{-3}$ mol dm$^{-3}$),
[NBu₄][BF₄] (0.1 mol dm⁻³) in MeCN at 16.5°C". No further details are provided.


(32) Crystal data for 1a: C₁₀H₂₀N₃Na₅O₅S₃, \( M_r = 381.46 \), orthorhombic, space group \( Pnma \), \( a = 16.158(3) \), \( b = 14.383(3) \), \( c = 7.371(2) \AA, \( V = 1713.1(6) \AA^3, Z = 4, \rho_{\text{calc}} = 1.479 \) g cm\(^{-3}, \mu = 0.481 \) mm\(^{-1}, \) \( T = 173(2) \) K, 2809 reflections collected (\( \theta \) range = 2.83-25.03°), 1557 unique (\( R_{\text{int}} = 0.0482 \)), \( R_I = 0.0546 \) [for 1018 reflections with \( I > 2\sigma(I) \)] and \( wR_2 = 0.1479 \) (for all data).

(33) The crystal structure of 1b has been reported previously (ref. 34) but the identification of the product of the \( \text{S}_2\text{N}_2/\text{Cp}_2\text{Co} \) reaction was confirmed by a full data collection and structure solution.


(37) Values for the kinetic parameters \( k_{s1}, k_{s2}, k_{f1}, \) and \( k_{f2} \) as well as the \( E^{o/} \) values for each redox couple along with values for the working electrode area \( A \) (6.6 x 10\(^{-2} \) cm\(^{2} \)) and \( D \) (\( \text{S}_4\text{N}_4[/\text{S}_4\text{N}_4]^- = 1.17 \times 10^{-5} \) cm\(^{2} \) s\(^{-1} \), \( [\text{S}_3\text{N}_3]^-/[\text{S}_3\text{N}_3]^+ = 4.00 \times 10^{-6} \) cm\(^{2} \) s\(^{-1} \)) were taken
over from the previous study and initially held invariant (see Ref 22). Other parameters employed were: T = 22 °C; concentration = $2 \times 10^{-3}$ mol L$^{-1}$; $v = 0.1 - 10$ V s$^{-1}$; $R_u = 250 \Omega$; working electrode area = 0.066 cm$^2$, $K_{eq1} = K_{eq3} = 10^6$, $K_{eq2} = 1.804 \times 10^5$, $D (S_2N_2, [S_2N_2]^{-}) = 9.13 \times 10^{-6}$ cm$^2$ s$^{-1}$.

(38) For the EC and ECC classifications of heterogeneous electron transfer reactions (E) with coupled homogeneous chemical reactions (C), see chapter 12 of ref. 35, pp. 471.

(39) We tested this hypothesis during the EPR investigations by measuring the decay of the $[S_4N_4]^{-}$ signal when electrolysis was conducted at −1.0 V at −70°C in a solution containing bulk $S_2N_2$. These measurements yielded a half-life for $[S_4N_4]^{-}$ in this environment of only 38±1 s compared to an expected value of 280±5 s based on the previous study (ref. 22) on solutions containing pure $S_4N_4$.


(42) EPR signal decay measurements from spectra obtained at −70°C for peaks attributed to $[S_2N_3H]^-$ fit best to a first-order decay process, with an estimated half-life of 7±1 s. Thus,
the lifetime of \([S_2N_2H]^+\) is about 5 × shorter than that of \([S_4N_4]^-\) under the same conditions (see ref. 39).


(44) For a catalytic cycle to operate (albeit slowly) it is only necessary that a small amount of \([S_2N_2]^+\) forms in eqn. 1 as turnover may be possible. Since voltammetry under quiescent conditions is not a good test for what happens in the bulk, we attempted to address this issue by performing punctuated electrolysis in stirred solution, halting after 1, 10 and 50% of the required current had passed, waiting for a catalytic reaction to transpire and interrogating the mixture via RDE voltammetry. However, our results were inconclusive probably because of equipment limitations. Testing of the catalytic hypothesis by chemical or electrochemical approaches is left for future work.


How does reduction (electrochemical or chemical) of $S_2N_2$ produce the $[S_3N_3]^{-}$ anion in high yield? Coupling of initially formed $[S_2N_2]^{-}$ with unreacted $S_2N_2$ produces $[S_4N_4]^{-}$; the latter is known to decay rapidly to give $[S_3N_3]^{-}$. EPR spectroscopy also provides evidence for protonation of $[S_2N_2]^{-}$ to produce the longer lived radical $[S_2N_2H]^{+}$. 

Tracey L. Roemmele,† Jari Konu,‡ René T. Boeré†* and Tristram Chivers‡*
(a) Interaction between the LUMO of S2N2 and the SOMO of [S2N2]−• and (b) SOMO of the C2v symmetric [S4N4]−• radical anion.
CVs (black lines) obtained on 2.0 mM solutions of S2N2 in CH2Cl2 containing 0.4 M [nBu4N][PF6] at a GC electrode at 20 °C, v = 0.2 V s−1 with the potential starting at −0.8 V: (a) sweeping in the positive direction and (b) sweeping in the negative direction. (c) Calculated CV (red line) resulting from input of the kinetic parameters listed in the text for the specific case of v = 0.2 V s−1 and kf3 = 108 M−1 s−1, sweeping the potential first in the negative direction starting at −0.8 V.
CV of a 2.0 mM soln of S2N2 in CH2Cl2 (0.4 M [nBu4N][PF6]), v = 0.2 V s−1, T = 20 °C, with the potential sweep starting at −0.8 V and extending further negative to include the second reduction process at −2.3 V (labeled Epc4).

82x54mm (600 x 600 DPI)
(a) RDE measurements on a 2.1 mM solution of S2N2 in CH2Cl2 with 0.4 M [nBu4N][PF6] at a 0.5 mm GC electrode, rotation rates = 1000 – 2250 RPM, ν = 0.010 V s−1; and (b) Koutecky-Levich 35 plot for the reduction of S2N2 (R2 = 0.998) after correction for S4N4 formation (see text).
(a) Experimental EPR spectrum (black line) obtained during in situ reductive electrolysis of 1.5 mM S2N2 at −1.5 V (164 s single scan) at a platinum foil electrode in CH3CN (0.1 M [nBu4N][PF6]) at −20 °C after noise reduction, and (b) simulation (red line), with $a(14N1) = 1.12$ mT, $a(14N2) = 0.65$ mT, with $a(1H) = 0.60$ mT, g value = 2.0136(1), mod. amp. = 0.2 mT.
Experimental EPR spectra from single scans obtained during in situ reductive electrolysis of 1.5 mM S2N2 at −1.5 V (164 s single scan) at a platinum foil electrode in CH2Cl2 (0.4 M [nBu4N][PF6]) containing HBF₄•2OEt₂: (a) at −40 °C (black line) and (b) the simulation of this spectrum (red line); note that in the simulation the small contribution by [S4N4]−• (14% by area) has been omitted to more clearly show the high-field line component. (c) At −20 °C after about 10 minutes (black line). (d) Simulation (red line) of the spectrum in (c).
Optimized gas-phase UB3LYP/6-31G(d,p) structure of [S2N2H•]; S1-N1-S2 94.7°, N1-S1-N2 87.8°, S2-N2-S1 89.4°, S1-N2-H 108.9°, N1-N2-H 120.4°, S1N1S2-S1N2S2 dihedral 6.9°.

80x47mm (300 x 300 DPI)
Crystal structure of 1a with the atomic numbering scheme. Hydrogen atoms have been omitted for clarity. a Symmetry operation: x, 0.5-y, z.
The proposed "ladder" scheme interrelating $\text{S}_2\text{N}_2$, $[\text{S}_4\text{N}_4]^\cdot$ and $[\text{S}_3\text{N}_3]^-$.

$\text{S}_2\text{N}_2 \xrightleftharpoons{+1e^-, k_{s3}} [\text{S}_2\text{N}_2]^\cdot + \text{S}_2\text{N}_2$

$\text{S}_4\text{N}_4 \xrightleftharpoons{+1e^-, k_{s1}} [\text{S}_4\text{N}_4]^\cdot$

$\text{[S}_3\text{N}_3]^\cdot \xrightleftharpoons{+1e^-, k_{s2}} [\text{S}_3\text{N}_3]^-$
Formation of the [S3N3]− anion from chemical reduction of S2N2

54x38mm (600 x 600 DPI)
How does reduction (electrochemical or chemical) of S2N2 produce the [S3N3]− anion in high yield? Coupling of initially formed [S2N2]−• with unreacted S2N2 produces [S4N4]−•; the latter is known to decay rapidly to give [S3N3]−. EPR spectroscopy also provides evidence for protonation of [S2N2]−• to produce the longer lived radical [S2N2H]•.