Studies on the Electroactive Heterocycles $\text{C}_6\text{S}_8^{0/2-}$ and $\text{C}_6\text{S}_6\text{O}_2^{0/2-}$ and Related Metal Complexes

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Abstract: The properties of the carbon sulfide $\text{C}_6\text{S}_8$ (1) as well as its dioxo analogue $\text{C}_6\text{S}_6\text{O}_2$ (2) have been studied in the fully oxidized and fully reduced forms, i.e., $1^0$, $1^{2-}$, $2^0$, $2^{2-}$. A crystallographic study shows that compound 2 is a tricyclic species with a twisted 1,2-dithiol (unsaturated $\text{C}_6\text{S}_2$ ring) core. Reduction of 1 with 2 equiv of LiBHET gave $\text{C}_6\text{S}_8^{2-}$ ($1^{2-}$), which was isolated as its PPh$_3$ salt. A crystallographic study of this salt shows that reduction has resulted in cleavage of the $\text{S}--\text{S}$ bond and that, in the solid state, the dihedral angle between the two rings is 180°. Both $1^{2-}$ and $2^{2-}$ undergo two sequential and reversible 1e$^-$ oxidations near 0 V (vs Ag/AgCl), thus establishing that $1^{0}$ and $2^{0}$ are good oxidants relative to other disulfides. Treatment of ($\text{PPH}_3$)$_2[\text{C}_6\text{S}_8]$ with [$\text{Ni(H}_2\text{O}_4)\text{C}_2\text{Cl}_2$] afforded the dark red ($\text{PPH}_3$)$_2[\text{Ni(C}_6\text{S}_8)\text{C}_2]$ (3). The analogous Pt complex (4) was also prepared. Crystallographic analysis of 3 shows that the nickel center is square planar, comprising two seven-membered NiS$_4$C$_4$ rings. The paramagnetism observed for solutions of 3 are attributed to the formation of a tetrahedral isomer.

Introduction

The redox behavior of the $\text{S}--\text{S}$ bond is important in many areas of chemistry.¹ For example, the scission of the $\text{S}--\text{S}$ bond in lipoic acid derivatives is an important step in the Krebs Citric Acid Cycle.² The cysteine-cystine equilibrium³ plays a determining role in many proteins: Synthetic polymers also have been prepared with redox active $\text{S}--\text{S}$ linkages in their backbones.⁴,⁵ Energy storage technologies have been based on the reduction of $\text{S}--\text{S}$ bonds, e.g., sulfur–sodium batteries.⁶ The charging and discharging cycle in such devices involves the shortening or elongation of polysulfido chains (eq 1).

$$\text{Na}_2\text{S}_3 + 2\text{S} \rightleftharpoons \text{Na}_2\text{S}_5$$ (1)

In recent years, research on the use of sulfur-based batteries has been invigorated through the development of both new cathodes and new anodes.⁷ Of relevance to the present work, organic disulfides have become of interest as replacement for inorganic sulfur in the cathodes.⁸ The redox cycle of the organic disulfides is similar to that involving alkali metal polysulfides, i.e., the reductive cleavage of $\text{S}--\text{S}$ bonds.⁹ Although the cycling rates of batteries based on organic disulfides is less than that for their inorganic counterparts,¹⁰ the organic disulfides offer a number of advantages including low operating temperatures. In contrast, alkali metal inorganic polysulfides have high melting points which necessitates that the battery operates at temperatures in the 300–400 °C range.¹¹ The organic disulfides are available with a wide range of substituents, thus allowing one to fine-tune the melting point, redox potential, and equivalent weight. Particular attention has focused on the use of polymeric organic disulfides—the so-called solid redox polymerization electrodes (SRPEs). Illustrative is the redox cycle for dimercaptothidiazolate (eq 2).

Since the battery’s cell potential is directly related to its energy density, there is interest in the synthesis of new disulfides which are highly oxidizing.

(9) NEXAFS studies establishing that the charging and discharging cycles of the Li-organic disulfide battery involves the formation and breaking of $\text{S}--\text{S}$ linkages: Yang, X. Q.; Xue, K. H.; Lee, H. S.; Guo, Y. H.; McBreen, J.; Skotheim, T. A.; Okamoto, Y.; Lu, F. J. Electrochem. Soc. 1993, 140, 943.

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The present results are derived from our research on new compositions composed only of sulfur and carbon. Our Structure of $C_6S_8O_2$ (eq 3). The corresponding $C_6S_8O_2$ is easily prepared from $C_6S_8$ (eq 3). The present studies on the redox chemistry of $C_6S_8$ and $C_6S_8O_2$ show the following: (i) that they are exceptionally oxidizing persulfido compounds, (ii) that their electrochemistry is reversible on the time scale of cyclic voltammetry, and (iii) that the reduction products, $C_6S_8^{2-}$ and $C_6S_8O_2^{2-}$, which are 1,4-dimercaptobutadiene derivatives, form stable coordination complexes.

**Results**

**Structural Studies on $C_6S_8$ and $C_6S_8O_2$.** We were initially interested in confirming the structure of at least one of these tricyclic species. We were unable to grow crystals of $C_6S_8$ (eq 1) because of its poor solubility. The related bis(dithiocarbonate) $C_6S_8O_2$ (eq 2), following trends noted earlier, is significantly more soluble and crystallized well from organic solvents. Single-crystal X-ray analysis showed that 1 has a tricyclic core and a single $S-S$ bond (Figure 1). The $S-S$ bond length is 2.054(2) Å (Table 1). The linkage $C2-C3-C4-C5$ exhibits single-double bond alternation as follows: 1.341(6), 1.458(6), 1.353(6) Å, respectively. Perhaps of greatest interest is the $C2-S5-S2-C5$ dihedral angle of 25.3°, the nonplanarity of which gives the molecule idealized $C_2$ symmetry. Schroth and coworkers have conducted the only other crystallographic characterization of tricyclic 1,2-dithiins (i.e., 1,2-dithiabenzenes), where the fused rings are thiophenes. In Schroth’s species the dihedral angles for the $C-S-S-C$ linkages are within a $2^\circ$ of $52^\circ$, i.e., they are much less planar than 2. In acyclic disulfides, the $C-S-S-C$ dihedral angles are typically $90^\circ$.

**Synthesis and Properties of $C_6S_8^{2-}$.** Suspensions of $C_6S_8$ in THF dissolve upon the addition of 2 equiv of LiBHET$_3$ to give violet solutions which contain the dianion $2^-$. Addition of methanolic PPh$_4$Cl to these solutions afforded an almost quantitative yield of violet-black microcrystals of (PPh$_4$)$_2[C_6S_8]$ (eq 4). This salt can be recrystallized from DMF:Et$_2$O. The same reduction can also be achieved using sodium-ammonia solutions. Chemical reduction of 2 with LiBHET$_3$ produces brown colored solutions containing $C_6S_8O_2^{2-}$; this anion was also isolated as its PPh$_4^+$ salt. The compound, (PPh$_4$)$_2[C_6S_8O_2]$ or (PPh$_4$)$_2[C_6S_8]$, is also soluble in DMF and DMSO. The 2:1 salts were characterized by several spectroscopic techniques. The $^{13}$C NMR spectrum of (PPh$_4$)$_2[C_6S_8]$ features three resonances for the anion at $\delta$ 207.3, 159.2, and 123.7, assigned to the thiocarbonyl and two olefinic carbon centers, respectively. The corresponding $^{13}$C NMR spectrum of (PPh$_4$)$_2[C_6S_8O_2]$ displays three resonances at $\delta$ 196.6, 148.4, and 111.6 in DMSO-d$_6$, all three of which are slightly shifted with respect to $C_6S_8^{2-}$. The IR spectrum of (PPh$_4$)$_2[C_6S_8]$ exhibits strong bands at 1439 and 1024 cm$^{-1}$, assigned to $v(C-S)$ and $v(S-S)$, respectively. These salts exhibit rich optical spectra.

This structure of (PPh$_4$)$_2[C_6S_8]$ was confirmed by single-crystal X-ray diffraction (Figure 2). The anion is centrosymmetric, consisting of a butadiene-like $C_4$ chain terminated in cyclic trithiocarbonates and bearing thiolato groups at C2 and C2A. The greatest difference between 2$^{-}$ and its neutral parent, aside

| Table 1. Selected Distances (Å) and Angles (Deg) in $C_6S_8O_2$ (2$^-$) with Standard Deviations in Parentheses |
|-----------------|-----------------|-----------------|
|                | S1–C2 1.745(4)  | S1–C1 1.786(5)  |
|                | S2–S5 2.054(2)  | S3–C3 1.758(4)  |
|                | S4–C4 1.755(5)  | S3–C1 1.780(5)  |
|                | S6–C5 1.743(5)  | S4–C6 1.777(5)  |
|                | S6–C6 1.778(5)  | S5–C2 1.773(4)  |
|                | O1–C1 1.198(6)  | O1–C1–S1 118.3(3) |
|                | C2–C6 1.202(6)  | C2–C3–S4 123.6(3) |
|                | C4–C5 1.356(3)  | C4–C3–S4 123.0(4) |

with bromine gave a precipitate of $C_6S_8$, as confirmed by the
resulted in rotation by 180° from the cleaved S–S bond, is the fact that reduction has
exhibited reversible couples at 0.09 and −0.07 V vs SSCE, respectively. These couples also show reversible
couples at 0.09 and −0.07 V vs SSCE, which were examined by cyclic voltammetry on DMF
solutions. Using a glassy carbon electrode, solutions of
2 gave two oxidative currents (Figure 3). There is a linear dependence of the peak
anodic and cathodic currents, indicative of the redox behavior of the thiolato centers in 2, examined the properties of two simple metal complexes. The complex
(Ph$_4$P)$_2$[Ni(C$_6$S$_8$)$_2$] (3) was prepared by combining DMF solutions of
(Ph$_4$P)$_2$C$_6$S$_6$O$_2$ and methanolic solution of [Ni(H$_2$O)$_6$]$^{2+}$Cl$_2$ (eq 5).
($\text{Ph}_4\text{P})_2\text{[Ni(C}_6\text{S}_8\text{)]}_2$) is a dark reddish brown
compound that dissolves in a variety of polar and weakly polar organic solvents. The compound is relatively air stable. In its
IR spectrum, the band at 1434 cm$^{-1}$ is assigned as vC–S. We also isolated the analogous complex ($\text{Ph}_4\text{P})_2\text{[Pt(C}_6\text{S}_8\text{)]}_2$) (4) via the reaction of 1$^2$− with K$_2$PtCl$_4$.
The crystallographic analysis of 3 confirms the presence of Ni$^{2+}$ in a square planar coordination sphere defined by four thiolatosulfur atoms (Figure 4, Table 3). The seven-membered C$_4S_2Ni$ ring is puckered, and the two C$_3S_2$ rings are twisted out of coplanarity with a dihedral angle of 37.8°. The easy formation of the nickel complex indicates that the rotation about the C3–C4 bond has a low barrier.

Whereas square planar nickel(II) complexes are diamagnetic, solutions of 3 are clearly paramagnetic. Attempts to record the
$^{13}$C NMR spectrum of the complex were unsuccessful, although strong signals were seen for the phenyl carbon atoms of the
Ph$_4$P cations. Solution magnetic measurements on CH$_2$Cl$_2$ solutions gave a moment of 1.05 $\mu_B$ under ambient conditions. Since CH$_2$Cl$_2$ is poorly coordinating, the paramagnetism is attributed to the formation of a complex of tetrahedral geometry, as might be expected in light of the tetrahedral geometry observed for Ni(SPh)$_2$$^{2−}$.

In contrast, the corresponding platinum complex remains diamagnetic in solution indicating

![Figure 2](image1.png)

**Figure 2.** Structure of the anion $2^{2−}$ in $(\text{Ph}_4\text{P})_2\text{[C}_6\text{S}_8\text{]}$ with thermal ellipsoids drawn at the 50% probability level.

**Figure 4.** Structure of the anion in $(\text{Ph}_4\text{P})_2\text{[Ni(C}_6\text{S}_8\text{)]}_2$ (3) with thermal ellipsoids drawn at the 50% probability level.

**Table 2.** Selected Distances (Å) and Angles (deg) for the Anion $1^{2−}$ in $(\text{Ph}_4\text{P})_2\text{[C}_6\text{S}_8\text{]}$ with Standard Deviations in Parentheses

<table>
<thead>
<tr>
<th>Bond/Angle</th>
<th>Distance/Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>S1–C1</td>
<td>1.726(2) Å</td>
</tr>
<tr>
<td>S2–C1</td>
<td>1.779(2) Å</td>
</tr>
<tr>
<td>S3–C2</td>
<td>1.718(2) Å</td>
</tr>
<tr>
<td>S4–C1</td>
<td>1.704(2) Å</td>
</tr>
<tr>
<td>S5–C3</td>
<td>1.751(2) Å</td>
</tr>
<tr>
<td>S6–C3</td>
<td>1.744(2) Å</td>
</tr>
<tr>
<td>C1–C2</td>
<td>1.704(2) Å</td>
</tr>
<tr>
<td>C2–S3</td>
<td>1.129(1) Å</td>
</tr>
<tr>
<td>C3–S4</td>
<td>1.129(1) Å</td>
</tr>
<tr>
<td>C4–S5</td>
<td>1.129(1) Å</td>
</tr>
</tbody>
</table>

![Table 2](image2.png)

Although we defer to a future more detailed report, some general aspects of the half oxidized species, e.g., 1$^+$ and 2$^+$,
are evident from the CV data. Using the calibration curve provided by Richardson and Taube for differential pulse voltammograms of two-electron couples,$^{18}$ $\Delta E_{1/2}$ for the two redox couples in 1 was determined to be 142 mV. The stability of the intermediate species, 1$^{−}$, can be expressed by the comproportionation constant, $K_{\text{com}}$, which was calculated using the equation $\log K_{\text{com}} = \frac{\Delta E_{1/2} - 0.0591}{2}$. These data indicates that solutions of 1$^{−}$ (or (1)$^{2−}$) should be relatively free of the 1$^0$ and 1$^{2−}$. A future report will deal with the characterization of the monoanion.$^{19}$

**Studies on $[\text{M(C}_6\text{S}_8\text{)]}_2$$^{2−}$.** As a further means of manipulating the redox behavior of the thiolato centers in 2, examined the properties of two simple metal complexes. The complex
(Ph$_4$P)$_2$[Ni(C$_6$S$_8$)$_2$] (3) was prepared by combining DMF solutions of
(Ph$_4$P)$_2$C$_6$S$_6$O$_2$ and methanolic solution of [Ni(H$_2$O)$_6$]$^{2+}$Cl$_2$ (eq 5).

$$2C_6S_8^{2−} + [\text{Ni(H}_2\text{O)}_6]^{2+} + 2\text{PPH}_4^{+} \rightarrow (\text{Ph}_4\text{P})_2\text{[Ni(C}_6\text{S}_8\text{)]}_2}$$

It is evident from the relative values of the peak anodic and cathodic currents (Figure 3). There is a linear dependence of the peak
anodic and cathodic currents, which are reversible and diffusion controlled. Complementary electrochemical studies were conducted on the diamons 1$^2$− and 2$^2$−, each of which displayed two oxidative couples at 0.09 and −0.06 V and at −0.07 and −0.21 V vs SSCE, respectively. These couples also show reversible behavior. Preparative scale oxidation of a MeOH slurry of 1$^{2−}$ with bromine gave a precipitate of C$_6$S$_8$, as confirmed by the
IR spectrum of the isolated solid.

![Figure 3](image3.png)

**Figure 3.** Cyclic voltammogram of (Ph$_4$P)$_2$C$_6$S$_6$O$_2$ (2$^{2−}$) and differential pulse voltammogram of C$_6$S$_6$O$_2$ (2), both for DMF solutions.

Although we defer to a future more detailed report, some general aspects of the half oxidized species, e.g., 1$^+$ and 2$^+$,
that it is square planar both in the solid state and in solution. The $^{13}$C NMR spectrum of Pt(C$_6$S$_8$)$_2^{2-}$ shows signals for both the organic cation and the inorganic anion, the latter consisting of three signals.

**Discussion**

The bicyclic species C$_6$S$_8$ is an unusual example of an electroactive inorganic heterocycle whose redox properties are noteworthy in the context of S–S containing species. While the structure of C$_6$S$_8$ was not determined directly, its analogue C$_6$S$_6$O$_2$ was fully characterized. A close structural similarity of the bis(trithiocarbonate) C$_6$S$_8$ and the bis(dithiocarbonate) C$_6$S$_6$O$_2$ is supported by the fact that the dioxo species is generated in good yield from C$_6$S$_8$. Furthermore, the structure of the dianion C$_6$S$_8^{2-}$ is that expected to result by cleavage of the S–S bond in C$_6$S$_8$. Finally the close structural relationship between 1 and 2 is indicated by the similar electrochemical properties of C$_6$S$_8$O$_2^{2-}$ and C$_6$S$_8^{2-}$. Crystallographic and $^{13}$C NMR analyses clearly show that the reduction of 1 and 2 is accompanied by scission of the S–S bond. The reduction of the S–S bond in 1,2-dithiins had not been previously observed.

Compounds 1 and 2 are most distinctive because their S–S bonds can be reversibly reduced, not only chemically but also analogously. A close structural relationship between 1 and 2 is indicated by the similar electrochemical properties of C$_6$S$_8$O$_2^{2-}$ and C$_6$S$_8^{2-}$. Crystallographic and $^{13}$C NMR analyses clearly show that the reduction of 1 and 2 is accompanied by scission of the S–S bond. The reduction of the S–S bond in 1,2-dithiins had not been previously observed.

The reversibility of the reduction process is attributable to the relative short time scale of cyclic voltammetry. The bonds can be reversibly reduced, not only chemically but also analogously. Examples of the reversible intramolecular scission of disulfides are most distinctive because their S–S forming process. Other examples of the reversible intramolecular scission of disulfides are known, e.g., Fe$_2$S$_2$(CO)$_2^{2-}$ and its inorganic analogue FeS$_2$, but these require potent redoxants, e.g., Na. Furthermore the oxidative coupling of thiocarbonylates on the time scale of cyclic voltammetry is quite uncommon.

Another noteworthy aspect of the redox chemistry of 1 and 2 is the mildness of the redox potentials, which are within 100 mV of SCE. Most organic disulfides require ~0.5 V more driving force to effect reduction. One factor favoring reduction is the electronic withdrawing character of the 1,3-dithiacyclopentene-2-thione substituents. The redox potential might also be influenced by the strain associated with the small dihedral angle for the central C–S–S–C linkage in 2 (and 1). This angle is normally near 90° for acyclic disulfides. The preference for 90° dihedral angles has been much discussed. More recent discussions have pointed out that 90° dihedral angles maximize overlap of the C–S $\sigma^*$ orbital and an otherwise nonbonding 2p electron ("lone pair") on sulfur: This explanation is consistent with the finding that stabilization increases with the electronegativity of R, since the energy of the $\sigma^*$ level will be lower for electronegative R. The stabilization arising from 90° dihedral angles is supported by studies on the rotational barriers for S–S bonds in R$_2$S$_2$. For organic disulfides, these barriers approach 6 kcal/mol.

To appreciate the high oxidizing power of 1 and 2, it is instructive to review the redox properties of other disulfides (Table 4). At pH = 7, organic disulfides typically reduce within 100 mV of ~0.25 V, e.g., ~0.19 V for Ph$_2$S$_2$ and ~0.32 V for lipoic acid (vs NHE). In contrast, the inorganic disulfides discussed in this work are reduced at 0.331 (C$_6$S$_6$) and 0.171 V (C$_6$S$_8$O$_2$). Thus, these new species are more than 0.5 V more oxidizing that organic disulfides. Again reflecting their electrophilic character of these species, the dithioloates are not readily protonated. Thus the redox for 1 and 2 is uncomplicated by protonation, in contrast to the situation for conventional organic disulfides. Polargraphic studies show that the oxidation (NC)$_2$C$_2$S$_2^{2-}$ (where (NC)$_2$C$_2$S$_2^{2-}$ is the maleonitriledithiolate or mnt$^-$) occurs at 0.321 V to give the acyclic disulfide [(CN)$_2$C$_2$S$_2$]$^{2+}$, thus the resulting disulfide is more oxidizing than 1 by ca. 150 mV. In contrast to 1, however, the oxidized form of this cyano-substituted 1,2-dithiolyte, i.e., [mnt$^-$]$_2$S, is too unstable for isolation.

The small separation of the two redox processes, $\Delta E/2$ for 1 and 2 is also unusual, but there are few precedents in sulfur.

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(19) Studies of Brietzer, J. G.; Chou, J.-H.; Smirnov, A.; Szczepura, L. F.; Rauchfuss, T. B.
(20) Qualitative measurements suggest that [Ni(C$_6$S$_6$O$_2$)$_2$]$^{2-}$ forms analogously.
chemically, partly because of the interfering effects of protonation, which in turn is due to the high basicity of most thiolates. Small values of Δ$E_{1/2}$ typically indicate either a (fast) chemical rearrangement that encourages the second redox process or low on-site Coulomb energy, which essentially means that the charges are effectively dispersed such that the effect of the first reduction (or oxidation) does not inhibit the second step. The latter explanation would be consistent with the π-delocalized character of the carbon sulfide framework. An additional factor that facilitates the second reduction is the possibility of strain inherent in 1 and 2 by virtue of the small C–S–S–C dihedral angle.

The lack of reversible redox activity in the complexes [M(C_6S_8)_2]^{2-} reaffirms the central importance of the S–S bond in the behavior of 1 and 2. These ligands are potentially interesting as homologs of the dithiolene (1,2-alkenedithiolate) ligands. The 1,2-dithioleines are renown as “noninnocent” due to their ability to stabilize metals in multiple oxidation states by virtue of the strong M–S π interactions. In contrast, the 1,4-dithioleines reported in this work appear to serve as innocent, chelating thioliates, perhaps due to the incompatibility of the nonplanar MS_C_4 ring with π-delocalization. We have previously reported a second family of 1,4-dithioleine complexes in the form of their titanocene derivatives which, like 3 and 4, feature seven-membered MS_C_4 rings:

Because C_6S_8^{2-} is easily prepared, this class of ligands may be of further interest as precursors to metal dithiolates with large S–M–S angles.

Experimental Section

Materials and Methods. All reagents and inorganic salts were used as received, while solvents were dried and deoxygenated by standard methods. We synthesized C_6S_8^{2-} by the addition of DMF to a suspension of Na in C_2S_2.

IR spectra were recorded on KBr disks and are reported in cm$^{-1}$. IR and $^{13}$C NMR absorptions for PPh_4^+ cation are seen in all samples at 527, 688, 722, 942, 1107, and 1439 cm$^{-1}$ and at δ117.6 (d), 130.5 (d), 134.5 (d), and 135.4 (d), respectively.

Electrochemical measurements were conducted in a three-electrode one-compartment cell containing a glassy carbon working electrode, a saturated sodium calomel electrode (0.2360 V vs NHE) reference electrode, and a Pt auxiliary electrode in 0.2 M Bu_4NPF_6 in DMF. Cyclic voltammetric measurements were measured at a scan rate of 100 mV/s, and differential pulse voltammograms were measured at a scan rate of 20 mV/s with a pulse amplitude of 10 mV, a pulse width of 50 ms, and a pulse period of 100 ms.

(PPh_4)[C_6S_8]. A dark purple suspension of 0.328 g (1 mmol) of C_6S_8 in 30 mL of THF was cooled with an acetone-dry ice bath and treated dropwise with 2.2 mL of 1 M THF solution of LiBHEt. The color of the solution changed to violet red almost immediately. The THF was removed under vacuum and 50 mL of O_2-free water was added to redissolve the dark brown residue. A solution of 0.75 g (2 mmol) of PPh_4Cl in 30 mL of O_2-free water was added to the dark brown solution to give a violet microcrystals. The crude solid was washed with water and dried under vacuum. Yield: 0.985 g (98%). X-ray quality crystals were grown by fractional crystallization from DMF/Me_2O as violet platelet crystals. Alternatively, the reduction of the C_6S_8 can be achieved by using Na metal in liquid NH_3. IR: 1365, 1602, 1721 (cm$^{-1}$) from the published values. The product was purified by reprecipitation from DMF/Et_2O as violet platelike crystals. Alternatively, the reduction of the C_6S_8 can be achieved by using Na metal in liquid NH_3. IR: 1365, 1602, 1721 (cm$^{-1}$) from the published values.

(PPh_4)[C_5S_6O_2]. A suspension of 0.317 g (1.1 mmol) of C_5S_6O_2 in 30 mL of THF was cooled with an acetone-dry ice bath and treated with 3.5 mL of 1.0 M THF solution of LiBHEt. The color of the solution changed to violet red almost immediately. The THF was removed under vacuum and 50 mL of O_2-free water was added to redissolve the dark brown residue. A solution of 0.75 g (2 mmol) of PPh_4Cl in 30 mL of O_2-free water was added to the dark brown solution to give a violet microcrystals. The crude solid was washed with water and dried under vacuum. Yield: 0.985 g (98%). X-ray quality crystals were grown by fractional crystallization from DMF/Me_2O as violet platelet crystals. Alternatively, the reduction of the C_6S_8 can be achieved by using Na metal in liquid NH_3. IR: 1365, 1602, 1721 (cm$^{-1}$) from the published values. The product was purified by reprecipitation from DMF/Et_2O as violet platelike crystals. Alternatively, the reduction of the C_6S_8 can be achieved by using Na metal in liquid NH_3. IR: 1365, 1602, 1721 (cm$^{-1}$) from the published values.


Table 5. Crystal and Data Collection Details for C₆S₆O₂, (PPh₄)₃C₆S₈, and (PPh₄)₂[Ni(C₆S₈)]

| Crystallographic Analysis of (P₄PhP)₃[C₆S₈]. Purple-red platelike crystals of were grown by fractional diffusion of Et₂O into a DMF solution of (P₄PhP)₃[C₆S₈]. The data crystal was attached to a thin glass fiber using Paratone-N oil (Exxon). The data crystal was bound by the (1 0 0), (−1 0 0), (0 0 1), (0 −1 0), (0 −1 0), (0 −1 1), and (−1 1 0) faces. Distances from the crystal center to these facial boundaries were 0.130, 0.130, 0.120, 0.120, 0.250, 0.250, 0.200, and 0.200 mm, respectively. Diffraction data were collected at 198 K on a Siemens 3-circle platform diffractometer with CCD area detector. Crystal and refinement details are given in Table 5. Systematic conditions suggested the ambiguous space group P1 (no. 2). Three standard intensities monitored every 90 min showed no decay. Step-scanned intensity data were reduced by profile analysis²⁸ and corrected for Lorentz and polarization effects and for absorption (SHELXLT version 5.03). Scattering factors and anomalous dispersion terms were taken from standard tables.²⁹

The structure was solved by direct methods⁴⁰ and refined by full-matrix least squares on F² (SHELXL 93).⁴¹ Correct positions for all nonhydrogen atoms were deduced from a vector map. H atoms U's were assigned as 1.2 times Ueq of the adjacent non-H atoms. All non-hydrogen atoms were refined with anisotropic thermal coefficients. The highest peak in the final map had no other significant features. A final analysis of variance between observed and calculated structure factors showed no dependence on amplitude or resolution.

Crystallographic Analysis of (P₄PhP)₂[Ni(C₆S₈)]. Dark-brown crystals were grown by fractional diffusion of Et₂O into a DMF solution of (P₄PhP)₂[Ni(C₆S₈)]. The data crystal was attached to a thin glass fiber using Paratone-N oil (Exxon). The data crystal was bound by the (0 0 1), (0 −1 0), (0 1 0), (−1 0 0), (−1 0 1), (−1 0 −1), and (1 0 1) faces. Distances from the crystal center to these facial boundaries were 0.18, 0.18, 0.28, 0.28, 0.32, and 0.24 mm, respectively. Diffraction data were collected at 198 K on a Siemens 3-circle platform diffractometer with CCD area detector. Crystal and refinement details are given in Table 5. Systematic conditions suggested the space group P2₁/n (no. 14). Three standard intensities monitored every 90 min showed no decay of the crystal. Step-scanned intensity data were reduced by profile analysis⁴⁸ and corrected for Lorentz–polarization effects and for absorption. Scattering factors and anomalous dispersion terms were taken from standard tables.²⁹

The structure was solved by direct methods⁴⁰ and refined by full-matrix least squares on F² (SHELXL 93).⁴¹ Correct positions for all nonhydrogen atoms were deduced from a vector map. H atoms U's were assigned as 1.2 times Ueq of the adjacent non-H atoms. All non-hydrogen atoms were refined with anisotropic thermal coefficients. The highest peak in the final map had no other significant features. A final analysis of variance between observed and calculated structure factors showed no dependence on amplitude or resolution.

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Supporting Information Available: Crystallographic data for compounds 2, (PPh$_4$)$_2$[1], and 3 including positional and thermal parameters (13 pages). See any current masthead page for ordering and Internet access instructions.

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