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# Sulfur monoxide thermal release from an anthracene-based precursor, spectroscopic identification, and transfer reactivity

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**Sulfur monoxide (SO) is a highly reactive molecule and thus, eludes bulk isolation. We report here on synthesis and reactivity of a molecular precursor for SO generation, namely 7-sulfinylamino-7-azadibenzonorbornadiene (**1**). This compound has been shown to fragment readily driven by dinitrogen expulsion and anthracene formation on heating in the solid state and in solution, releasing SO at mild temperatures (<100 °C). The generated SO was detected in the gas phase by MS and rotational spectroscopy. In solution, **1** allows for SO transfer to organic molecules as well as transition metal complexes.**

microwave spectroscopy | reactive intermediate | molecular precursor | astrochemistry | sulfur monoxide

In contrast to the ubiquitous and well-studied chemistry of earth-abundant dioxygen (**1**), the chemistry of its heavier, valence-isoelectronic analogue sulfur monoxide (SO) is hardly explored and has been relegated to a niche existence, which is certainly due to its high reactivity: SO is unstable under ambient conditions toward disproportionation to SO<sub>2</sub> and elemental S (**2**) and eludes bulk isolation. However, in space, SO can accumulate and has been found in the interstellar medium (**3**, **4**) as well as in our solar system (**5–7**), which is important to note considering that both O and S are biogenic elements (**8**).

Fragmentation of suitable molecular precursors presents a potential entry point to explore the synthetic chemistry for such reactive species and opens new avenues for spectroscopic characterization (**9–16**). In the case of SO, a limited number of synthetic precursors have been reported that allow thermal transfer of SO (Fig. 1): well-investigated are the chemistries of episulfoxides (**A**) (**17–20**), a thiadiazepin *S*-oxide (**B**) (**21**), trisulfide oxides (**C**) (**22**, **23**), thianorbornadiene-*S*-oxides (**D**) (**24**), and *N*-sulfinylamine phosphinoborane adducts (**E**) (**25**). In explaining the SO transfer reactions of some of these substances, the intermediacy of free SO is assumed, while for others, the precursors fragment likely via associative mechanisms.

Our group has a longstanding interest in small reactive species, such as P<sub>2</sub> (**9–11**), AsP (**12**), HCP (**13**), phosphinidenes (**14**, **15**), and dimethylgermylene (**16**), generated by mild thermal activation of suitable precursors. The driving force of anthracene (C<sub>14</sub>H<sub>10</sub>, **A**) expulsion for the release of highly reactive molecules and subsequent characterization and synthetic transfer has been amply capitalized on (**13**, **14**, **26–31**). Against this backdrop and our reasoning that an additional N<sub>2</sub> unit should further increase the energy of the ground state of the precursor molecule, we envisioned our synthetic target, 7-sulfinylamino-7-azadibenzonorbornadiene, OSN<sub>2</sub>**A** (**1**), as promising for SO release simultaneously with **A** and dinitrogen formation. To probe this hypothesis, we compared the computed Gibbs free energies for singlet SO release from *A–E* and **1** (Fig. 1). Indeed, the formation of singlet SO was predicted to be thermodynamically strongly favorable only in case of **1**.

Free SO, amenable to spectroscopy, has been generated by electric discharge experiments of SO-containing gases (OCS, SO<sub>2</sub>) (**32**) or using ethylene episulfoxide at high temperature (180 °C to 580 °C) (**33**). To the best of our knowledge, spectroscopic observation of free SO provided by mild thermolysis of a well-defined, solid, and easy-to-handle precursor compound has not been achieved. With **1**, we present now the synthesis of such a compound that fragments at approximately 95 °C in the solid state and allows for direct detection of SO in the gas phase. In addition, examples of SO transfer with this reagent in solution to both organic molecules and transition metal complexes are outlined.

## Results and Discussion

The synthesis of **1** was achieved by reaction of Carpino's hydrazine (7-amino-7-azadibenzonorbornadiene, H<sub>2</sub>N<sub>2</sub>**A**) (**30**) with thionyl chloride in the presence of triethyl amine (Scheme 1) (**34**).

Compound **1** was isolated as a pale yellow solid in very good yield (83%). In solution (benzene-*d*<sub>6</sub>), the <sup>1</sup>H NMR chemical shift of the bridgehead protons at δ = 6.22 ppm, located 1.48 ppm downfield from that of H<sub>2</sub>N<sub>2</sub>**A**, is reflective of the strongly withdrawing effect of the sulfinyl group. Colorless crystals grew from a concentrated toluene solution of **1** layered with diethyl ether at –35 °C and were subjected to X-ray diffraction analysis.

## Significance

The generation of highly reactive molecules under controlled conditions is desirable, as it allows exploration of synthetic chemistry and enables spectroscopic studies of such elusive species. We report here on the synthesis and reactivity of a precursor molecule that readily fragments with concomitant expulsion of dinitrogen and anthracene to release the highly reactive sulfur monoxide, a compound of interest for both synthetic chemists and astrochemists.

Author contributions: M.J., M.N., W.J.T., and C.C.C. designed research; M.J., M.N., W.J.T., M.-A.M.-D., M.C.M., and D.P. performed research; M.J., M.N., W.J.T., M.-A.M.-D., M.C.M., and D.P. analyzed data; and M.J., M.N., W.J.T. and C.C.C. wrote the paper.

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The authors declare no conflict of interest.

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Data deposition: The crystallography, atomic coordinates, and structure factors have been deposited in the Cambridge Structural Database to [www.ccdc.cam.ac.uk/Solutions/CSDSystem/Pages/CSD.aspx](http://www.ccdc.cam.ac.uk/Solutions/CSDSystem/Pages/CSD.aspx) (accession nos. 1567576 and 1567577).

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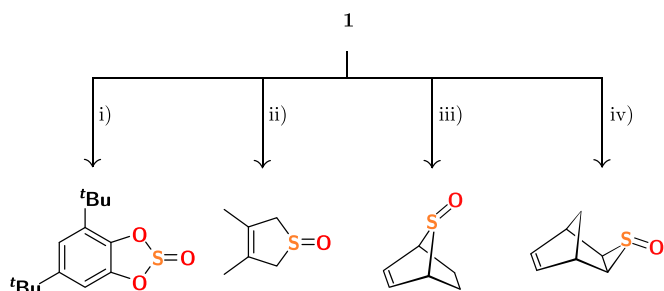
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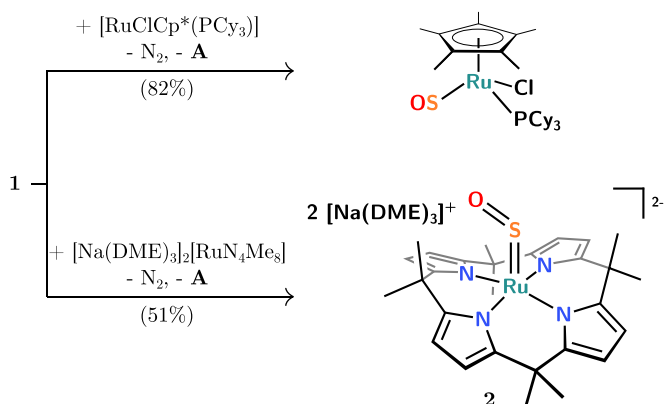


**Scheme 2.** Transfer of SO from **1**: (i) 3,5-di-*tert*-butyl-1,2-quinone (1 eq), benzene, 70 °C, 24 h (quantitative); (ii) DMB (neat), 80 °C, 16 h (60%); (iii) 1,3-cyclohexadiene (5 eq), benzene, 70 °C, 24 h (59%); (iv) norbornadiene (10 eq), benzene, 25 °C, 16 h (55%).

in principle: the radiative lifetimes of these species were determined to be approximately 7 ms (exp. value; calculated 13.6 ms) and 450 ms (calculated), respectively (47, 48). Singlet SO is thus sufficiently long-lived for detection with our apparatus with an approximate flight time of 2–3 ms assuming a reasonable molecular velocity in the gas phase (*SI Appendix*). However, an additional complication is the propensity of singlet SO for collisional relaxation to the  $^3\Sigma^-$  spin state with third bodies, such as the heated walls of the sample holder (49).

In addition, we were able to identify the rotational transitions of **1** itself, which partially transferred without fragmentation into the gas phase. The pure rotational spectrum of **1** was analyzed using a variant of the Automated Microwave Double Resonance technique (50). The small differences between the experimentally obtained rotational constants and those recovered from calculations based on the geometry of **1** in the crystal structure imply that the structure of **1** is nearly identical in the solid state and in the gas phase (*SI Appendix*). The differences in the observed products of thermolysis of **1** highlight the complementary nature of the techniques used but also illustrate how critical the thermolysis conditions are to what gases are evolved from **1**.

To comment on the thermolysis mechanism of **1**, quantum chemical calculations were carried out using the RI-B2PLYP-D3(BJ) density functional and the Def2-TZVP basis set. A single-step fragmentation with concerted loss of singlet SO and N<sub>2</sub> from the anthracene platform was found, featuring a single transition state corresponding to the breaking of the C–N bond opposed to the sulfinylamino moiety. The minimum energy path across this transition state revealed no further stationary points en route to the final products: the structure collapsed in a concerted, asynchronous way with breaking of the N–S and of the second C–N bond. This is likely due to the NNSO



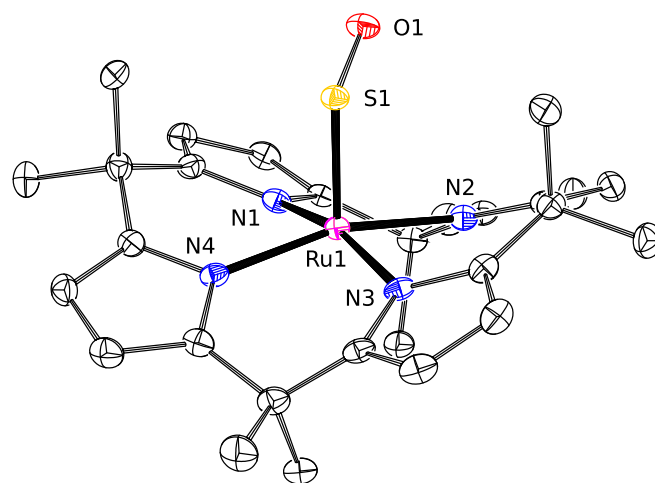
**Scheme 3.** Transfer of SO from **1** to ruthenium complexes.

intermediate not being a minimum on the potential energy surface (51). This quasimultaneous breaking and making of bonds in this fragmentation process may be classified as a coarctate reaction as defined by Herges (52). While for pericyclic reactions, the concerted breaking and forming of bonds occur in a cyclic manner, coarctate processes describe bond rearrangements with a transition-state topology of a constricted cycle. The constriction point is located at the coarctate atom(s) at which two bonds are broken and reformed in the transition state. In case of the fragmentation of **1**, both N atoms are coarctate atoms. In this process, SO may indeed be released in its singlet state: the formation of A, N<sub>2</sub>, and singlet SO is favored by  $-40.4 \text{ kcal}\cdot\text{mol}^{-1}$  with respect to the starting material (Fig. 5).

With evidence of SO release on thermolysis in the solid state, we explored the reactivity of **1** in solution. Heating **1** in benzene-*d*<sub>6</sub> in a sealed tube led to its decomposition, which was indicated visually by a color change from colorless to yellow likely due to production of sulfur. The sole product observable by <sup>1</sup>H NMR spectroscopy was A. This reaction obeyed a first-order rate law with  $k_{\text{obs}} = (2.743 \pm 0.436) \times 10^{-4} \text{ s}^{-1}$  at 80 °C in THF as determined by ultraviolet–visible spectroscopy. This barrier corresponded to an activation barrier of  $\Delta G^\ddagger(80^\circ\text{C}) = 26.55(11) \text{ kcal}\cdot\text{mol}^{-1}$  according to the Eyring equation, similar to the calculated value of  $\Delta G_{\text{calc}}^\ddagger(80^\circ\text{C}) = 23.3 \text{ kcal}\cdot\text{mol}^{-1}$ .

To assess the possibility of SO transfer from **1** to an acceptor, various representative reaction partners were used. We focused first on organic compounds suitable for SO trapping, such as quinones and olefins (Scheme 2). Compound **1** was heated with 3,5-di-*tert*-butyl-1,2-quinone for 24 h to 70 °C to convert quantitatively to the corresponding known sulfite (53).

Several SO-releasing precursors are capable of SO addition to 1,3-dienes (18, 21–24, 54, 55). Heating of **1** with an excess (5 eq) of 2,3-dimethyl-1,3-butadiene (DMB) led to decomposition of **1** without thiophene-*S*-oxide formation. However, when performing the reaction at 80 °C in neat DMB, this SO transfer product was observed (60%). Similarly, transfer of SO to 1,3-cyclohexadiene was successful. Likely due to the locked *cisoid* conformation of the double bond, just a fivefold excess of this diene was sufficient to lead to formation of 7-oxo-7-thianorbornene (59%) (23). Thermal SO transfer to other olefins or alkynes was unsuccessful: reactions using *cis*-stilbene, styrene,



**Fig. 6.** Molecular structure of **2** in the solid state with thermal ellipsoids at the 50% probability level. SO-coordinated sodium cation and separate sodium tris(1,2-dimethoxyethane) cation as well as one cocrystallized THF solvent molecule have been omitted for clarity. Selected distances (angstroms) and angles (degrees): Ru1–S1 2.0282(8), S1–O1 1.503(2), Ru1–N 2.044 (average); Ru1–S1–O1 118.17(11).

or phenyl acetylene in excess did not provide the respective addition products. Contrasting reactivity of **1** toward norbornadiene yielded the corresponding thiirane (55%): addition of SO occurred at 25 °C (16 h), well below the temperature required for fragmentation of **1**, and thus, it does not involve free SO but rather, proceeds via an associative mechanism. The SO transfer from **1** to transition metal complexes also occurs through an associative mechanism (Scheme 3). Stirring a solution of **1** and [RuCl(Cp\*)(PCy<sub>3</sub>)<sub>3</sub>] (Cp\* = η<sup>5</sup>-C<sub>5</sub>Me<sub>5</sub>) in THF at 25 °C for 30 min led to a gradual color change from blue to red–brown. After removal of **A**, the known SO ruthenium complex [RuCl(Cp\*)(SO)(PCy<sub>3</sub>)<sub>3</sub>] was isolated (82%) (56, 57). Compound **1** also reacted with the anionic ruthenium complex [Na(DME)<sub>3</sub>]<sub>2</sub>[Ru(N<sub>4</sub>Me<sub>8</sub>)] (H<sub>4</sub>N<sub>4</sub>Me<sub>8</sub> = octamethylporphyrinogen) (**58**) to give [Na(DME)<sub>3</sub>][Ru(N<sub>4</sub>Me<sub>8</sub>)(SO)] (**2**), obtained in 51% yield after removal of anthracene and selective crystallization as a brown–orange solid.

Cooling a solution of this compound in THF and DME to –35 °C yielded dark orange crystalline blocks. An X-ray diffraction analysis revealed a dimeric structure, with sodium ions bridging two units of the [Ru(N<sub>4</sub>Me<sub>8</sub>)(SO)]<sup>2–</sup> platform (Fig. 6). These units feature the ruthenium center in a square-pyramidal environment, in which the SO ligand occupies the apical position. The Ru–S bond [2.0282(8) Å] is slightly shorter and the S–O bond [1.503(2) Å] is slightly longer than in a related Ru(II)–SO complex [2.0563(11) and 1.447(3) Å, respectively] (59). Terminal SO transition metal complexes generally show a strong SO stretch around 1,046–1,126 cm<sup>–1</sup> (55, 56, 59, 60). Analysis of the IR spectrum of **2** revealed a band at 1,021 cm<sup>–1</sup> assigned to the SO stretching vibration. Both this notable red shift with regard to values for related compounds as well as the metrical data for **2** are in accord with strong backbonding from Ru to the

SO ligand due to the electron-rich porphyrinogen ligand. Coordination of the sodium cations to the oxygen atom of ligated SO may enhance this effect.

We did not observe selective reaction of **1** with *N*-heterocyclic carbenes to give the corresponding sulfines. However, heating a mixture of **1** and a phosphine (PPh<sub>3</sub> or P<sup>*t*</sup>Bu<sub>3</sub>) in benzene gave about equimolar mixtures of the respective phosphine oxides and phosphine sulfides via formal splitting of SO (25).

## Conclusion

We have shown here a well-controlled synthetic route to SO by thermal decomposition of **1**. Taking reactivity, computational studies, and spectroscopic detection of <sup>3</sup>Σ<sup>–</sup> SO into consideration, it is believed that **1** generates <sup>1</sup>Δ SO on thermolysis. <sup>3</sup>Σ<sup>–</sup> SO is detected by microwave spectroscopy, possibly originating from a small amount of <sup>1</sup>Δ SO that has had enough time to phosphoresce into the lower-energy triplet ground state. Regardless of the spin state of the SO evolved from **1**, this study firmly establishes that SO is in fact released from the molecular precursor, illustrating the power of synthesis in combination with spectroscopy to shed light on reactive intermediates of general importance.

## Methods

Experimental and computational details and crystallographic information are included in *SI Appendix*. Computed atomic coordinates are included in *Datasets S1–S4*.

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