**Proline-based phosphoramidite reagents for the reductive ligation of S-nitrosothiols**

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S-Nitrosothiols (RSNOs) have many biological implications but are rarely used in organic synthesis. In this work we report the development of proline-based phosphoramidite substrates that can effectively convert RSNOs to proline-based sulfenamides through a reductive ligation process. A unique property of this method is that the phosphine oxide moiety on the ligation products can be readily removed under acidic conditions. In conjugation with the facile preparation of RSNOs from the corresponding thiols (RSHs), this method provides a new way to prepare proline-based sulfenamides from simple thiol starting materials.

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**INTRODUCTION**

Sulfur-nitrogen (-S-N-) linkages are unique chemical moieties. These structures often show interesting bioactivities (Figure 1). For example, sulfonamides are a class of important herbicides.¹,² Sulfanilamides have been used as common drugs for infections.³,⁴ N-Thiolated β-lactams are found to have antibacterial, antifungal and anticancer effects.⁵–¹² Sulfenamides exhibit antimicrobial activities against various infectious pathogens.¹³ Proline-based sulfenamides have been recognized as potential antiproliferative and anti-infective agents.¹⁴–¹⁶ Because of these activities, the preparation of these molecules, especially the formation of the -S-N- linkages, has become an active area in organic synthesis. So far, many methods have been developed for the construction of the -S-N- bonds, including: (1) sulfenylation between sulfenyl chlorides and amines,¹⁷,¹⁸ (2) amination of thiols with N-halo compounds or amines in the presence of oxidizing reagents to give unsubstituted and N-substituted sulfenamides,¹⁹ (3) the treatment of disulfides with ammonia or amines in the presence of silver or mercuric salts in alkaline medium,²⁰,²¹ (4) sulfenylation of primary and secondary amines with the esters of sulfenic acids,²² (5) derivations of N-chlorothio-compounds,²³,²⁴ (6) the [2,3]-sigmatropic rearrangement of S-allylsulfynimines,²⁵ (7) reductive cleavage of sulfaminic acids in the presence of thiophenols,²⁶ (8) electrolysis of 2-mercaptothenazolium-amine or bisbenzothiazol-2-ylsulfide-amine²⁷,²⁸ and (9) the preparation of sulfin- or sulfon-amides by the oxidation of sulfenamides in the presence of the oxidants.²⁹,³⁰

**RESULTS AND DISCUSSION**

Small organic thiols (RSHs) can also be easily converted to the corresponding RSNOs under mild nitrosation conditions. Normally, RSNOs are unstable species that makes their detection in biological systems very challenging. Moreover, the instability of RSNOs makes these compounds unattractive for synthetic chemists. As a result, the application of RSNOs in synthesis has been rarely reported. Interestingly, in our recent work on RSNO bioorthogonal reactions (aiming at the development of novel detection methods for protein RSNO formation),³¹–³⁴ we have discovered some unusual reactivity and properties of RSNOs. In our opinion, RSNOs are powerful synthons that can be used to introduce S, N and/or O atoms into molecular structures. Herein, we wish to report a method to prepare proline-based sulfenamides from readily available RSH substrates via RSNO intermediates. The novel reactions and synthetic strategies introduced in this article could be applied for an effective synthesis of bioactive molecules including a proline moiety.

**Scheme 1**

In our previous work, we discovered that RSNOs and triarylphosphines (2 equivalents) can rapidly react to generate reactive thioazaylide intermediates in high yields under mild conditions.³⁴ Thioazaylides are potent nucleophilic species. Upon manipulating the electrophilic groups attached to the phosphine reagents, thioazaylides can be trapped as stable products. For example, as shown in Scheme 1, when an *ortho*-ester group was attached to the phosphine reagent (compound 3), the reactive thioazaylde 4 could be trapped via an intramolecular acyl transfer to afford sulfenamide. Small organic thiols (RSHs) can also be easily converted to the corresponding RSNOs under mild nitrosation conditions. Normally, RSNOs are unstable species that makes their detection in biological systems very challenging. Moreover, the instability of RSNOs makes these compounds unattractive for synthetic chemists. As a result, the application of RSNOs in synthesis has been rarely reported. Interestingly, in our recent work on RSNO bioorthogonal reactions (aiming at the development of novel detection methods for protein RSNO formation),³¹–³⁴ we have discovered some unusual reactivity and properties of RSNOs. In our opinion, RSNOs are powerful synthons that can be used to introduce S, N and/or O atoms into molecular structures. Herein, we wish to report a method to prepare proline-based sulfenamides from readily available RSH substrates via RSNO intermediates. The novel reactions and synthetic strategies introduced in this article could be applied for an effective synthesis of bioactive molecules including a proline moiety.

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simple RSH starting materials. However, the use of phosphine substrates like 3 would lead to products like 5 that contain an unnecessary triphenylphosphine oxide moiety. The removal of this bulky group from the final products would require harsh conditions. Therefore, seeking phosphine substrates that can undergo the reductive ligation while leading to a readily removable phosphine oxide moiety from the products is desirable. With this idea in mind, we proposed that proline-based phosphoramidites like 6 would be suitable substrates. As shown in Scheme 1, the reaction between RSNO and 6 should still follow the reductive ligation process. The resultant phosphoramidate moiety can be considered as a NH-protecting group and removable under acidic condition. As such, proline-linked sulfenamides could be prepared from simple RSH and phosphoramidite 6.

To test this idea, three proline-based phosphoramidites (Scheme 2, 9–11) were prepared. These substrates were treated with TrSNO (12a) to explore their reactivity. The selection of TrSNO as the RSNO model compound was because of its remarkable stability and ease of synthesis. We found the characteristic green color of TrSNO disappeared immediately when treated with all three phosphoramidites, suggesting the formation of thioazaylides was fast. For 9 and 10, the reactions stopped at this stage as no desired ligation product was obtained even when the reaction time was extended to 24 h. With 11, however, we obtained the desired ligation product 16a. Apparently, this can be explained by the factor that phenyl ester is a better leaving group than the methyl and t-butyl esters. We tested a series of different solvents for this reaction. The best was found to be a mixture of THF/phosphate buffer (pH 7.4, 20 mM, 3:1) that gave 16a in 84% yield.

As phosphoramidite 11 was found to be the most reactive substrate for this ligation, it was applied to other RSNO substrates to test the generality of the reaction. A series of primary, secondary and tertiary RSNOs were freshly prepared from RSH and then used in this study without any purification. The results were summarized in Table 1. In all cases, the desired ligation products were obtained. For relatively stable tertiary RSNO substrates, the corresponding ligation products were isolated in modest to high yields (entries 1–4). Secondary and primary RSNO substrates gave slightly lower yields (entries 5–9).

Interestingly, the purification of secondary RSNO products (entries 5 and 6) was found to be very difficult because they overlapped with the

Figure 1  Representative molecules having sulfur-nitrogen (-S-N-) linkages.
phosphine oxide by-product. Nevertheless, the formation of the products was clearly confirmed by NMR and mass spectroscopy analysis. These results demonstrated that RSH can be readily converted to proline-based sulfenamides by this two-step SNO-ligation process.

Although phosphoramidite 9 did not show good reactivity to convert RSNO to the desired sulfenamides, we did observe some unique reactivity of 9. As shown in Scheme 3, the treatment of a cysteine SNO derivative 12g with 9 gave dehydroalanine 18 in a modest yield (37%) together with the disulfide product (46%).

These results indicated that the reaction indeed proceeded to form the thioazaylide intermediate 17. Presumably, the methyl ester was not reactive enough. Hence, the acyl transfer was slow and nonproductive. Instead, the azaylide underwent an intramolecular β-elimination on the cysteine substrate to form dehydroalanine 18.

In the proline-based sulfenamide products, the diphenylphosphoryl moiety could be considered as the protecting group of proline. We expected it could be removed under acidic conditions.

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**Table 1 Summary of the reactions between 11 and S-nitrosothiols (RSNOs)**

| entry | RSNO          | product            | yield (%) |
|-------|---------------|--------------------|-----------|
| 1     | Ph—SNO        | 16a                | 84        |
| 2     | BnHN—SNO      | 16b                | 92        |
| 3     |                 | 16c                | 54        |
| 4     | SNO            | 16d                | 57        |
| 5     |                 | 16e                | 82a       |
| 6     |                 | 16f                | 40a       |
| 7     | MeO—SNO       | 16g                | 62        |
| 8     |                 | 16h                | 42        |
| 9     |                 | 16i                | 40        |

*aYields determined by NMR analysis.*
conditions. We then tested the deprotection of 16g (Scheme 4).

Indeed, a cleavage cocktail of 10% water in TFA provided sulfenamide almost quantitatively. This result revealed that the -S-N- bond on the sulfenamide compounds was quite stable under acidic conditions. Given the efficiency of this protocol, it can find applications in making water-soluble proline-based sulfenamides.

HNO is the one-electron reduced/protonated form of nitric oxide. It shows distinct physiology and pharmacology from nitric oxide. The reductive ligation was also found to be effective for HNO. Several specific fluorescent probes for HNO detection have been developed based on the triarylphosphine template. We next wondered whether the proline-based phosphoramidite would work for HNO and tested the reaction between 11 and HNO (generated from Angeli’s salt). As shown in Scheme 5, the reaction proved to be fast and efficient. The desired ligation product 21 was obtained in a comparable yield (50%) as the one obtained from triarylphosphines, suggesting the proline-based phosphoramidite may be useful for the design of novel HNO sensors.

CONCLUSIONS
In summary, we have developed a reductive ligation of S-nitrosothiols using N-diphenylphosphine proline ester substrates. This reaction was found to be effective for all small-molecule S-nitrosothiols (primary, secondary and tertiary), as well as for HNO. The ligation products bear a removable phosphine oxide moiety on the proline residue. In conjunction with the facile preparation of RSNOs from the corresponding RSHs, this novel method provides a unique way to prepare proline-based bioactive sulfenamides from simple thiol starting materials.

METHODS

Instrumentation
1H NMR spectra, 13C NMR and 31P NMR were recorded at 300 MHz (VX 300, Varian, Palo Alto, CA, USA) and are reported in p.p.m. on the δ scale relative to residual CHCl3 (δ 7.25 for 1H and δ 77.0 for 13C). These experiments were performed at room temperature. Mass spectra were recorded using an electrospray ionization mass spectrometry (ESI, Thermo Finnigan LCQ Advantage, San Jose, CA, USA) or MALDI-TOF (matrix-assisted laser desorption ionization time-of-flight) mass spectrometry. Mass data were reported in units of m/z for [M+H]⁺ or [M+Na]⁺.

Preparation of proline-based phosphoramidites
To a solution of the proline ester HCl salt in CH2Cl2 (total c = 0.30 M) was added freshly distilled 2.5 equivalent of triethylamine followed by PPh2Cl (1 equivalent) at 0 °C. The mixture was slowly warmed to room temperature and stirred for 3–4 h. Upon completion (monitored by TLC), the reaction mixture was filtered to remove the triethylammonium salt. The filtrate was concentrated and diluted with EtOAc, and washed with satd NaHCO3, water and brine. The organic layer was dried with anhydrous MgSO4, filtered and concentrated. The crude product was purified by flash column chromatography (with preneutralized silica gel by 3% triethylamine in hexanes).

Compound 9: 1H NMR (300 MHz, CDCl3) δ 7.59–7.27 (m, 10H), 4.22 (ddd, J = 8.8, 6.0, 3.5 Hz, 1H), 3.63 (s, 3H), 3.17–3.06 (m, 1H), 2.88 (m, 1H), 2.22–1.70 (m, 4H); 31P NMR (122 MHz, CDCl3) δ 49.31; 13C NMR (75 MHz, CDCl3) δ 176.00, 138.93, 132.88, 132.61, 132.23, 131.98, 128.90, 128.40, 128.33, 128.27, 128.19, 65.28, 64.87, 51.96, 47.64, 47.57, 31.80, 31.71, 25.90; MS (ESI) m/z calcd for C18H21NO2P [M+H]⁺ 314.1, found 314.1.

Compound 10: 1H NMR (300 MHz, CDCl3) δ 7.49 (ddd, J = 8.1, 5.4, 1.5 Hz, 2H), 7.39–7.28 (m, 8H), 4.07 (ddd, J = 8.6, 6.6, 3.4 Hz, 1H), 2.83 (ddd, J = 9.2, 6.8, 2.7 Hz, 1H), 2.07 (dq, J = 12.0, 8.3 Hz, 1H), 1.99–1.82 (m, 2H), 1.76–1.64 (m, 1H), 1.40 (s, 9H). 13C NMR (75 MHz, CDCl3) δ 174.76, 139.27, 132.83, 132.26, 132.01, 128.81, 128.38, 128.29, 128.19.
Preparation of S-nitrosothiols

The thiol starting material (RSH, 0.2 mmol) was dissolved in 1 mL of MeOH followed by the addition of 1 N HCl (1 mL) at room temperature. To this solution was then added freshly prepared 1 N NaNO2 (1 mL) in water in dark (total c = 0.07 M). The color of the reaction was immediately turned to red (for primary and secondary RSNO) or green (for tertiary RSNO). The mixture was stirred for 10–15 min at room temperature. Upon completion (monitored by TLC), the RSNO product was directly extracted with cold diethyl ether (1 mL x 3) in dark. The organic layers were collected and dried. The solvent was removed to provide the RSNO product that was then used for the ligase reaction without further purification.

General reductive ligation procedure

To the freshly prepared RSNO product was added a solution of 2 equiv of 11 in 3:1 THF-aqueous buffer (pH 7.4, degassed by bubbling with argon). The final concentration was ~0.1 M. The reaction was monitored by TLC and it was usually completed within 15–30 min at room temperature. The reaction mixture was extracted with ethyl acetate. The combined organic layers were washed with water and brine, dried by anhydrous Na2SO4 and concentrated. The crude product was purified by flash column chromatography.

Thioacetamide 15: 1H NMR (300 MHz, CDCl3) δ 7.81–7.55 (m, 4H), 7.47–7.17 (m, 2CH3), 6.94–6.80 (m, 3H), 4.35 (d, J = 8.8, 3.5 Hz, 1H), 3.14 (qd, J = 6.5, 2.8 Hz, 2H), 2.30 (m, 1H), 2.13–1.99 (m, 1H), 1.99–1.78 (m, 2H); 31P (122 MHz, CDCl3) δ 18.3; FT-IR (thin film) 3025.4, 2985.3, 1728.8 (strong, C=O, carbonyl group), 1610.3, 1450.0, 1372.3, 1286.0, 1247.1, 1071.0 cm⁻¹; MS (ESI) m/z calc for C2H6N2NaO2PS [M+Na]+ 672, found 671.

Thioacetamide 16: 1H NMR (300 MHz, CDCl3) δ 7.81 (s, 1H), 7.76–7.24 (m, 25H), 3.95 (dt, J = 9.2, 5.3 Hz, 1H), 3.01 (p, J = 6.1, 1H), 2.66–2.51 (m, 1H), 1.92 (m, 2H), 1.68–1.61 (m, 1H), 1.34–1.26 (m, 1H); 31P (122 MHz, CDCl3) δ 30.2; 13C NMR (75 MHz, CDCl3) δ 178.99, 150.71, 145.25, 133.24, 132.12, 132.87, 132.74, 132.01, 131.59, 130.54, 130.04, 129.81, 129.51, 128.59, 128.41, 128.22, 128.15, 127.56, 126.42, 126.00, 126.11, 126.01, 115.57, 63.10, 60.11, 49.03, 32.18, 25.76; MS (ESI) m/z calc for C2H6N2NaO2PS [M+Na]+ 672, found 671.

Dehydroxylamine (Dha) 18: 1H NMR (300 MHz, CDCl3) δ 8.54 (br, 1H), 7.85–7.81 (m, 2H), 7.57–7.43 (m, 3H), 6.79 (s, 1H), 5.98 (d, J = 1.4 Hz), 3.88 (s, 3H); 13C NMR (75 MHz, CDCl3) δ 171.22, 170.56, 170.56, 170.08, 170.0, 170.5, 170.54, 170.57, 132.54, 132.54, 132.57, 132.57, 132.51, 132.42, 132.37, 132.30, 132.24, 131.76, 131.52, 130.05, 129.84, 129.19, 129.19, 128.93, 126.02, 126.02, 53.70, 52.66, 48.47, 48.46, 48.01, 42.04, 31.27, 31.19, 25.26, 25.17, 23.24, 19.77, 17.57; MS (ESI) m/z calc for C2H6N2NaO2PS [M+Na]+ 659, found 659.

Deprotection of the diphenylphosphoryl group

A sulfenamidic product (0.05 mmol) was treated with 1 mL of cold 10% water in TFA at 0 °C (total c = 0.05 M). The resulting solution was stirred for 30 min at 0 °C. Upon completion, the excess TFA was removed with hexanes as the co-solvent under vacuum. To the remaining mixture was added cold diethyl ether to solidify the proline-sulfenamide TFA salt. The solid product was further washed with cold diethyl ether (2 mL x 5) and dried to provide the final product.

The reaction between 11 and HNO

To an argon sparged mixture of acetonitrile and water was added 11 (0.21 mmol), and to this stirring mixture was added freshly prepared Angelis’s salt (0.1 mmol, Na2N3O5). The resulting solution was allowed to stir until the reaction was completed by (TLC) or 20 min. The product 21 (50% yield) was isolated by extraction and flash column chromatography.

To the 11 (300 MHz, CDCl3) δ 0.04 (br, 1H, NH); 7.96–7.68 (m, 5H), 7.64–7.39 (m, 5H), 5.85 (br, 1H, NH), 4.03 (d, J = 8.1, 5.7, 2.3 Hz, 1H), 3.18 (m, 2H), 2.46–2.18 (m, 1H), 2.18–1.99 (m, 1H), 1.89 (dd, J = 13.5, 8.9, 4.2 Hz, 2H); 31P (122 MHz, CDCl3) δ 28.6; 13C NMR
The authors declare no conflict of interest.

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