2,2-Bis(phenylsulfonyl)ethyl sulfides as efficient precursors of sulfenic acids

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This paper is dedicated to Professor Nicolò Vivona on the occasion of his 70th birthday

Abstract
The use of 2,2-bis(phenylsulfonyl)ethyl sulfides as efficient precursors of sulfenic acids is reported. These starting compounds, carrying both sulfone and sulfide residues, are obtained in one step from commercially available thiols and 1,1-bis(phenylsulfonyl)ethylene. Mild decomposition of the derived 2,2-bis(phenylsulfonyl)ethyl sulfoxides to sulfenic acids, in the presence of suitable acceptors, opens the way to the generation of a wide library of new sulfoxides not easily accessible through different synthetic pathways.

Keywords: 1,1-Bis(phenylsulfonyl)ethylene, β-syn-addition, β-syn-elimination, sulfenic acids

Introduction

Sulfenic acids 1 are generally transient species (Scheme 1), not isolable when no stabilizing structural feature is present in the R moiety directly linked to the sulfur atom.1

Scheme 1
Many methods are reported in the literature\(^2\) for the \textit{in situ} generation of 1. The most exploited procedure is the thermolysis of suitable sulfoxides: this reaction is an intramolecular β-\textit{syn}-elimination, that often happens at high temperatures, and follows a concerted mechanism (Scheme 1).

The temperature of the sulfoxide decomposition into sulfenic acid and alkene depends on the mobility of the hydrogens in β-position to the sulfinyl moiety. The presence of one or two electron-withdrawing groups directly linked to the β-carbon enhances the mobility of this hydrogen atom,\(^3\) consequently lowering the activation energy of the decomposition. The transient sulfenic acid can add, once formed, to unsaturations, carbon-carbon double or triple bonds, \textit{via} a \textit{syn}-addition concerted mechanism, to produce various kinds of functionalized sulfoxides in regio- and stereo-selective manner, if the case. When the sulfenic acid bears at least one stereogenic structural feature, two sulfur epimers are formed, in some cases with a certain stereoselection.\(^4\) These sulfoxides can easily be separated to constitute enantiopure starting products of many stereoselective reactions where the sulfoxide moiety acts as chiral auxiliary.\(^5\)

Since the ‘90s part of our chemistry has been devoted to the employment of thio-functionalized compounds in the synthesis of molecules of applicative interest.\(^6\) In many cases the key step of the procedure has been the concerted \textit{syn}-addition of sulfenic acids 1 to unsaturated systems. If severe conditions are needed for the thermolysis step - sulfenic acid formation / addition to unsaturation as a unique domino process - the yields of the desired products are notably affected. In fact, elevated temperatures favour the self-condensation of the sulfenic acids as well as their redox transformations.\(^7\)

Finally, most of the reported syntheses involving sulfenic acid intermediation lead to the formation of vinyl sulfoxides, resulting from sulfenic acid addition on an alkyne, while only few cases involving the addition of 1 to double bonds are reported.\(^2\)

This paper intends to demonstrate the central importance of 1,1-bis(phenylsulfonyl)ethylene (2) as a key reagent in the generation of sulfenic acids. The conversion of the sulfinyl precursors takes place even at room temperature (RT), thus limiting the formation of side products to very low yields. Moreover the mildness of the reaction conditions allows the easy addition of the intermediate sulfenic acid even to double bonds, in quantitative yields.

**Results and Discussion**

The first step of our project has been the preparation of sulfides 5 by reacting thiols 4 with an almost equimolar amount of strongly electrophilic 1,1-bis(phenylsulfonyl)ethylene (2)\(^8\) in the presence of Triton B as a base (Scheme 2).\(^9\) In order to evaluate the applicability of the method, four thiols have been chosen as models: an aliphatic one [cyclohexanethiol (4a)], an aromatic one [benzenethiol (4b)], phenylmethanethiol (4c), and \(N\)-(\textit{tert}-butoxycarbonyl)-L-cysteine methyl ester (4d).
The nucleophilic addition of the obtained thiolate to the electron-poor unsaturation of 2 led to the formation of bis-sulfone sulfides 5 in quantitative yields. Subsequent controlled oxidation of 5 with meta-chloroperoxybenzoic acid (m-CPBA) at –78 °C gave sulfoxides 3 within a few minutes (Scheme 2).

In order to evaluate the capability of the newly formed sulfinyl systems in being good sulfenic acid precursors, sulfoxides 3b (racemic mixture) was thermolyzed in the presence of a typical alkyne acceptor, dimethyl acetylenedicarboxylate (DMAD, Scheme 3).

Scheme 3

The reaction was performed in dichloromethane (DCM) at 40°C, in the presence of an excess (1:6) of acceptor, using TLC to follow the consumption of the starting products 3b. The results obtained from the thermolysis of sulfoxides 3b in the presence of DMAD, in terms of reaction times and yields of the final addition products 7b, were compared with those obtained from sulfoxides 6 (racemic mixture), coming from the synthetic path shown in Scheme 3. In both cases the vinyl sulfoxides 7b (racemic mixture) were the main products. The great difference in the reaction times between 3b and 6 (20 min versus 12 h), both in DCM, at 40 °C and with the same sulfoxide/DMAD molar ratio (1:6), gave us the confirmation that the
thermolysis of the sulfoxide precursor is the rate limiting step of the total process and that the new precursor 3b requires shorter time with respect to 6 for decomposition to sulfenic acid 1b. This means that the two geminal sulfanyl groups render the β-hydrogen more mobile than in many other known starting products. All the sulfoxides 3 reacted in the same conditions, giving the addition products 7a-d, with comparable reaction times. Results are reported in Table 1 (entries 1, 4, 9, 12). The first outcome of these results is that sulfoxides 3 are too reactive to be stored, and therefore they have to be utilized as a crude in the subsequent reaction, at once after their formation by sulfide oxidation.

<table>
<thead>
<tr>
<th>Entry</th>
<th>Sulfenic acid</th>
<th>Acceptor</th>
<th>Reaction conditions</th>
<th>Product</th>
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<td>1</td>
<td>1a</td>
<td>DMAD</td>
<td>DCM, 40 °C, 1:6, 30 min</td>
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<td>1a</td>
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<td>1b</td>
<td>DMAD</td>
<td>DCM, 40 °C, 1:6, 20 min</td>
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<td>1b</td>
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<td>9</td>
<td>1c</td>
<td>DMAD</td>
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<td>12</td>
<td>1d</td>
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<td>15</td>
<td>1d</td>
<td>Methyl Acrylate</td>
<td>DCM, RT, 1:6, 100 h</td>
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This result could appear, at first sight, a limitation to our synthetic process. In other respects, the very high yields observed for the final vinyl sulfoxides prompted us to test the same process, but in milder conditions. Kept in the presence of DMAD in the same sulfoxide/acceptor molar ratio 1:6 at RT, crude led after 6 h to vinyl sulfoxides in quantitative yield (Table 1, entry 5). No products originated by self-condensation of were isolated, thus demonstrating that the mild conditions applied for sulfinic acid generation are also optimal for the addition step.

The same method was applied for the addition to double bonds. In fact, while the addition reaction of sulfinic acids to triple bonds has largely been exploited in the synthesis of vinyl sulfoxides, a comment must be made about the addition to the carbon-carbon double bonds. In this last case the adduction has to obey to more strict steric demands and, before the sulfinic intermediate can succeed in reaching the unsaturation in a proper geometry, it can easily undergo self-condensation to produce thiosulfinates. For this reason very few are the examples of efficient syntheses of sulfoxides via sulfinic acid / alkene concerted adduction, most of them being intramolecular, because of the easier accessibility of a close double bond to be attached. Otherwise, most of the reported examples of intermolecular addition of sulfinic acids to double bonds involve electron-poor unsaturations: it is suggested that in such cases the reaction follows a less-concerted, dipolar-like mechanism.

Sulfoxides were then reacted at RT in neat methyl acrylate (Table 1, entries 2, 6, 10, 14), and the unique products obtained in quantitative yields were the methyl sulfinylpropanoates resulting from the completely regioselective adduction of the sulfinic acids onto the electron-poor double bond. All the reactions performed in neat acceptor have been repeated in DCM, still at RT, with acrylate in excess (1:6) (Table 1, entries 3, 7, 11, 15). Those experiments aimed at estimating whether the sulfinic acid could attack the double bond even when not completely surrounded by the acceptor, but only in the presence of a reasonable molar excess: the yields compare with those obtained in neat acceptor, whereas reaction times are more or less doubled.

The above results bring confirmation that the method could be applied even in the presence of more expensive or not commercially available unsaturated acceptors, and this opens the way to many more applications. As reported in Table 1, also the reactions of gave good results (entries 8 and 16, products ). These findings are consistent with the smooth generation of the four sulfinic acids and their clean addition to the double bond of both the acrylates.
A particular comment must be devoted to 3d, a case where two non-equivalent β-hydrogens are available with respect to the sulfoxide group, as already reported in our previous papers (Scheme 4).6a-c

![Scheme 4](image)

Besides self-condensation of 1d, another side reaction is then expected to happen: the generation of a different competing sulfenic acid. The same situation having appeared in our previous works,6b we had overcome the problem by using a precursor which could undergo thermolysis at 40 °C, thus minimizing formation of the undesired sulfenic acid. That was confirmed by the inspection of the 1H NMR spectra of the crudes of our reactions of 3d with different sulfenic acid acceptors, all indicating the absence of the amino ester 10 (Scheme 4).

In order to apply our method to the elaboration of new classes of three-branched molecules6d containing sulfinyl or sulfonyl moieties we thought to synthesize trissulfoxides 12 (Scheme 5). The reaction of a six-fold excess of 3d [precursor of (R)-2-(tert-butoxycarbonylamino)-2-(methoxycarbonyl)ethanesulfenic acid (1d)] with trisacrylate 1116 has been conducted in DCM at RT (Table 1, entry 17).
Scheme 5

The first results of this reaction show the formation of a mixture of diastereoisomeric sulfoxides 12, resulting from the completely regioselective addition of the sulfenic acid 1d onto the three vinyl groups in 11. In a first approach, the diastereomeric mixture 12 will be oxidized to a unique tris-sulfone, in order to rid ourselves of the sulfur stereogenicity. The next step will involve deprotection of amino and carboxyl moieties in the tris-sulfone, and evaluation of the biological activity of the enantiopure derivative obtained.

Conclusions
The Table 1 shows the good results obtained in synthesizing various sulfoxides 7-9 and 12 drawing advantage from the chemistry of sulfenic acids 1. Mild conditions required by the procedure and high yields observed support the validity of 2,2-bis(phenylsulfonyl)ethyl sulfides and their sulfoxide derivatives as efficient precursors of sulfenic acids generated by intramolecular β-syn-elimination.

Experimental Section

General Procedures. Solvents were purified according to standard procedures. Petroleum ether used refers to the fraction boiling at 30–50 °C. All reactions were monitored by TLC on commercially available precoated plates (Aldrich silica gel 60 F 254) and the products were visualized with vanillin [1 g dissolved in MeOH (60 ml) and conc. H2SO4 (0.6 ml)]; the reported Rf refer to EtOAc / petroleum ether 3:2 as eluent, unless otherwise stated. Silica gel used for column chromatography was Aldrich 60. 1H and 13C NMR spectra were recorded on a Varian Mercury 300 spectrometer at 300 and 75 MHz respectively in CDCl3 solutions with SiMe4 as internal standard: J values are given in Hz; the attributions are supported by Attached Proton Test (APT) and homodecoupling experiments.

Formation of sulfides 5a-d. General procedure. To a 2 M solution of commercial thiols 4a-d in anhydrous tetrahydrofuran (THF), at - 78 °C under argon atmosphere and continuous stirring, Triton B was added (40% wt MeOH sol., thiol 4 / Triton B molar ratio 1:0.12). After 5 min stirring a 2 M THF solution of 2 (4/2 molar ratio 1:1.2) was slowly added. The reaction was monitored by TLC (EtOAc / petroleum ether 1:3), to follow consumption of the starting product 4a-d. After completion, brine was added, the reaction mixture was extracted with DCM, and the collected extracts washed twice with brine. After drying over Na2SO4, the solution was filtered and concentrated under reduced pressure to give the reaction crude, then submitted to chromatographic purification (EtOAc / petroleum ether 1:3). All yields were almost quantitative. The characterization of sulfides 5a,b was consistent with the data reported in the literature.17

1,1-Bis(phenylsulfonyl)-2-(benzylsulfanyl)ethane (5c). Light yellow solid, m.p. 94-97 °C. TLC: Rf 0.87. 1H NMR: δ 7.8-7.2 (m, 15H, H arom), 4.40 (t, J1,2 6.0, 1H, H-1), 3.70 (s, 2H, PhCH2), 3.21 (d, 2H, H2-2). 13C NMR: δ 137.7, 137.2, 136.5, 134.9, 134.5, 129.4, 129.3, 129.1, 128.9, 128.7, 128.5, 128.3, 127.1, and 127.0 (C arom), 83.9 (C-1), 37.5 (PhC2H2), 26.8 (C-2). Anal. calcd. for C21H20O4S3 (432.58): C, 58.31; H, 4.66%. Found: C, 58.33; H, 4.63%.

1,1-Bis(phenylsulfonyl)-2-[(R)-2-(tert-butoxycarbonylamino)-2-(methoxycarbonyl)ethylsulfanyl]ethane (5d). Transparent oil. TLC: Rf 0.75. 1H NMR: δ 8.0-7.5 (m, 10H, H arom), 5.29 (br d, J vic 5.8, 1H, NH), 4.93 (br t, J1,2 5.6, 1H, H-1), 4.43 (dd, J vic 6.2 and 4.7, 1H, NCH), 3.76 (s, 3H, OMe), 3.34 (d, 2H, H2-2), 2.98 and 2.83 (split AB system, J gem 14.2, 2H, NCHCH2), 1.46 (s, 9H, CMe3). 13C NMR: δ 171.1 (CO2Me), 155.1 (OCONH), 137.6,
137.5, 134.7, 129.6, 129.1, and 129.0 (C$_{arom}$), 84.3 (C-1), 80.3 (CMe$_3$), 53.0 and 52.7 (NCH and OMe), 36.7 (NCHCH$_2$), 28.2 (CMe$_3$), 27.7 (C-2). Anal. calcd. for C$_{23}$H$_{29}$NO$_8$S$_3$ (543.67): C, 50.81; H, 5.38; N, 2.58%. Found: C, 50.84; H, 5.41; N, 2.56%.

Oxidation of sulfides 5a-d to sulfoxides 3a-d. General procedure. To a stirred 0.2 M solution of 5a-d in DCM, at – 78 °C, a DCM 0.2 M solution of an equimolar amount of m-CPBA was added. The reaction was followed by TLC (EtOAc / petroleum ether 1:1) until complete transformation of the sulfide. A 10% wt aqueous solution of Na$_2$S$_2$O$_3$ was added, and the mixture maintained under stirring until it reached RT. The inorganic layer was discarded and the organic phase was washed twice with aqueous sat. NaHCO$_3$, then twice with brine. After drying over Na$_2$SO$_4$, the solvent was removed in vacuo and the crude oil used at once in the subsequent thermolysis reaction.

1,1-Bis(phenylsulfonyl)-2-(cyclohexylsulfinyl)ethanes 3a (racemic mixture). TLC: $R_f$ 0.42. $^1$H NMR: δ 8.0-7.5 (m, 10H, H$_{arom}$), 5.29 (dd, $J_{1,2}$ 7.5 and 4.0, 1H, H-1), 3.52 (m, 2H, H$_2$-2), 2.57 (m, 1H, Cy CHS), 2.1-1.2 (m, 10H, 5 x Cy CH$_2$).

1,1-Bis(phenylsulfonyl)-2-(phenylsulfinyl)ethanes 3b (racemic mixture). TLC: $R_f$ 0.40. $^1$H NMR: δ 8.0-7.5 (m, 15H, H$_{arom}$), 5.11 (dd, $J_{1,2}$ 8.5 and 3.5, 1H, H-1), 3.57 and 3.51 (split AB system, $J_{2A,2B}$ 14.0, 2H, H$_2$-2).

1,1-Bis(phenylsulfonyl)-2-(benzylsulfinyl)ethanes 3c (racemic mixture). TLC: $R_f$ 0.38. $^1$H NMR: δ 7.9-7.3 (m, 15H, H$_{arom}$), 5.23 (dd, $J_{1,2}$ 8.3 and 3.4, 1H, H-1), 4.12 and 4.06 (AB system, $J_{gem}$ 13.0, 2H, PhCH$_2$), 3.43 and 3.38 (split AB system, $J_{2A,2B}$ 14.5, 2H, H$_2$-2).

1,1-Bis(phenylsulfonyl)-2-[((R)-2-(tert-butoxycarbonylamino)-2-(methoxycarbonyl)ethylsulfinyl]ethanes 3d (1:1 S-epimeric mixture). TLC: $R_f$ 0.38. $^1$H NMR: δ 7.9-7.5 (m, 10H, H$_{arom}$), 5.54 (m, 1H, NH), 5.24 (m, 1H, H-1), 4.65 (m, 1H, NCH), 3.80 and 3.78 (two s, 3H, OMe), 3.8-3.2 (m, 4H, NCHCH$_2$ and H$_2$-2), 1.46 and 1.45 (two s, 9H, CMe$_3$).

2-Propenoic acid 1,3,5-benzenetriyl ester (11). Prepared according to a previously reported procedure,$^{16}$ purified by very rapid flash chromatography and used immediately in the thermolysis reaction. TLC: $R_f$ (EtOAc / petroleum ether 1:1) 0.86. $^1$H NMR: δ 6.96 (s, 3H, H$_{arom}$), 6.61 (dd, $J_{(trans)_{2,3A}}$ 17.2, $J_{3A,3B}$ 1.3, 3H, 3 x H$_A$-3), 6.29 (dd, $J_{(cis)_{2,3B}}$ 10.5, 3H, 3 x H-2), 6.03 (dd, 3H, 3 x H$_B$-3).

Thermolysis of sulfoxides 3a-d in the presence of DMAD, methyl and phenyl acrylate, 2-propenoic acid 1,3,5-benzenetriyl ester (11). General procedure. A 0.2 M DCM solution of 3 was added, under continuous stirring, of an excess of alkyn or alkene acceptor (1:6 molar ratio); then the reaction was maintained under stirring at RT or under reflux (Table 1). When the reaction appeared complete by TLC (consumption of 3, EtOAc / petroleum ether 1:1, see reaction times in Table 1), the reaction crude was purified by flash column chromatography on silica gel (EtOAc / petroleum ether from 1:9 to 1:1). The yields were always > 70%.
characterization of the obtained sulfoxides 7b and 8b,c was consistent with the data reported by the literature.\textsuperscript{10-12}

\textbf{(E)-2-(Cyclohexylsulfinyl)-2-butenedioic acids dimethyl esters 7a (racemic mixture).} Transparent oil. TLC: \textit{Rf} 0.79. \textsuperscript{1}H NMR: \(\delta\) 6.69 (s, 1H, H-3), 3.87 and 3.82 (two s, 6H, 2 x OMe), 2.72 (tt, \textit{J} \text{v}ic 12.2 and 3.6, 1H, Cy CHS), 2.1-1.1 (m, 10H, 5 x CH\textsubscript{2}). \textsuperscript{13}C NMR: \(\delta\) 164.2 and 162.4 (C-1,4), 148.2 (C-2), 128.5 (C-3), 58.8 (Cy CHS), 53.2 and 52.6 (2 x OMe), 27.6, 25.9, 25.2, 25.0, and 21.5 (5 x CH\textsubscript{2}). Anal. calcd. for \(\text{C}_{12}\text{H}_{18}\text{O}_{5}\text{S}\) (274.33): C, 52.54; H, 6.61%. Found: C, 52.55; H, 6.59%.

\textbf{Transparent oil. TLC: \textit{Rf} 0.77.} \textsuperscript{1}H NMR: \(\delta\) 7.4-7.2 (m, 5H, H arom), 6.49 (s, 1H, H-3), 4.29 and 4.03 (AB system, \textit{J} \text{gem} 13.1, 2H, CH\textsubscript{2}), 3.87 and 3.79 (two s, 6H, 2 x OMe). \textsuperscript{13}C NMR: \(\delta\) 164.3 and 162.5 (C-1,4), 147.3 (C-2), 130.7, 130.2, 129.6, 129.4, 128.7, and 128.5 (C arom and C-3), 59.4 (CH\textsubscript{2}), 53.2 and 52.6 (2 x OMe). Anal. calcd. for \(\text{C}_{13}\text{H}_{14}\text{O}_{5}\text{S}\) (282.31): C, 55.31; H, 5.00%. Found: C, 55.32; H, 4.98%.

\textbf{(E)-2-[(R)-2-(tert-Butoxycarbonylamino)-2-(methoxycarbonyl)ethylsulfinyl]-2-butenedioic acids dimethyl esters 7d (1:1 S-epimeric mixture).} Transparent oil. TLC: \textit{Rf} 0.65. \textsuperscript{1}H NMR: \(\delta\) 6.95 and 6.90 (two s, 1H, H-3), 5.59 (m, 1H, NH), 4.66 (m, 1H, NCH), 3.87 and 3.79 (three s, 6H, 2 x OMe), 3.8-3.2 (m, 2H, CH\textsubscript{2}), 1.45 (s, 9H, CMe\textsubscript{3}). \textsuperscript{13}C NMR: \(\delta\) 170.4, 170.2, and 164.4 (C-1,4 and CO\textsubscript{2}Me), 147.6 and 147.5 (C-2 and OCONH), 129.7 (C-3), 80.7 (CMe\textsubscript{3}), 58.4 and 54.3 (CH\textsubscript{2}), 53.3, 53.0, 52.7, 49.7, and 49.1 (NCH and 3 x OMe), 28.2 and 28.1 (CMe\textsubscript{3}). Anal. calcd. for \(\text{C}_{15}\text{H}_{23}\text{NO}_{7}\text{S}\) (393.41): C, 45.79; H, 5.89; N, 3.56%. Found: C, 45.82; H, 5.89; N, 3.59%.

\textbf{3-(Cyclohexylsulfinyl)propanoic acids methyl esters 8a (racemic mixture).} Light yellow solid, m.p. 105-107 °C. TLC: \textit{Rf} 0.21. \textsuperscript{1}H NMR: \(\delta\) 3.70 (s, 3H, OMe), 3.0-2.8 (m, 4H, H 2-2,3), 2.56 (tt, \textit{J} \text{v}ic 11.3 and 3.4, 1H, Cy CHS), 2.2-1.2 (m, 10H, 5 x Cy CH\textsubscript{2}). \textsuperscript{13}C NMR: \(\delta\) 171.9 (C-1), 59.4 (CH), 52.1 (OMe), 43.6 (C-3), 27.1, 26.2, 25.4, 25.3, and 25.0 (C-2 and 5 x Cy CH\textsubscript{2}). Anal. calcd. for \(\text{C}_{10}\text{H}_{18}\text{O}_{3}\text{S}\) (218.31): C, 55.02; H, 8.31%. Found: C, 55.02; H, 8.33%.

\textbf{Transparent oil. TLC: \textit{Rf} 0.26.} \textsuperscript{1}H NMR: \(\delta\) 5.80 and 5.70 (two br d, \textit{J} \text{v}ic 7.8, 1H, NH), 4.69 (m, 1H, NCH), 3.80, 3.79, 3.74, and 3.73 (four s, 6H, 2 x OMe), 3.4-2.8 (m, 6H, 3 x CH\textsubscript{2}), 1.45 (s, 9H, CMe\textsubscript{3}). \textsuperscript{13}C NMR: \(\delta\) 171.6, 171.5, 170.7, and 170.4 (2 x CO\textsubscript{2}Me), 154.5 and 154.2 (OCONH), 80.5 (CMe\textsubscript{3}), 54.3 and 53.5 (NCHCH\textsubscript{2}), 53.0, 52.9, 52.2, 50.0, and 49.6 (NCH and 2 x OMe), 47.4 and 47.0 (C-3), 28.1 (CMe\textsubscript{3}), 26.7 and 23.6 (C-2). Anal. calcd. for \(\text{C}_{13}\text{H}_{23}\text{NO}_{7}\text{S}\) (337.39): C, 46.28; H, 6.87; N, 4.15%. Found: C, 46.24; H, 6.87; N, 4.17%.

\textbf{3-([R]-2-(tert-Butoxycarbonylamino)-2-(methoxycarbonyl)ethylsulfinyl]propanoic acids methyl esters 8d (1:1 S-epimeric mixture).} Transparent oil. TLC: \textit{Rf} 0.64. \textsuperscript{1}H NMR: \(\delta\) 7.5-7.0 (m, 10H, H arom), 3.4-2.8 (m, 4H, H-2,3). \textsuperscript{13}C NMR: \(\delta\) 169.9 (C-1), 150.4 and 142.7 (quaternary C arom), 131.2, 129.4, 129.3, 129.1, 126.0, 124.0, and 121.3 (CH\textsubscript{arom}), 50.8 (C-3), 26.2 (C-2). Anal. calcd. for \(\text{C}_{15}\text{H}_{14}\text{O}_{5}\text{S}\) (274.33): C, 65.67; H, 5.14%. Found: C, 65.6; H, 5.16%.

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3-[(R)-2-(tert-Butoxycarbonylamino)-2-(methoxycarbonyl)ethylsulfinyl]propanoic acids
phenyl esters 9d (1:1 S-epimeric mixture). Low melting white solid. TLC: \( R_f 0.45 \). \(^1\)H NMR: \( \delta \) 7.4-7.1 (m, 5H, \( H_{arom} \)), 5.67 (m, 1H, NH), 4.70 (m, 1H, NCH), 3.81 and 3.79 (two s, 3H, OMe), 3.4-3.0 (m, 6H, 3 x CH\(_2\)), 1.45 (s, 9H, CMe\(_3\)). \(^{13}\)C NMR: \( \delta \) 170.3 and 169.8 (C-1 and CO\(_2\)Me), 155.3 and 150.4 (quaternary Carom and OCONH), 129.5, 126.1, and 121.3 (CH\(_{arom}\)), 54.4 (NCH\(_2\)CH\(_2\)), 53.1, 53.0, 50.1 and 49.7 (NCH and OMe), 47.3 and 46.8 (C-3), 28.2 (CMe\(_3\)), 27.1 and 27.0 (C-2). Anal. calcd. for C\(_{18}\)H\(_{25}\)NO\(_7\)S (399.46): C, 54.12; H, 6.31; N, 3.51%. Found: C, 54.08; H, 6.30; N, 3.54%.

3-[(R)-2-(tert-Butoxycarbonylamino)-2-(methoxycarbonyl)ethylsulfinyl]propanoic acids
1,3,5-benzenetriyl esters 12 (diastereomeric mixture). Low melting yellow solid. TLC: \( R_f \) (EtOAc) 0.10. \(^1\)H NMR: \( \delta \) 6.90 (m, 3H, \( H_{arom} \)), 5.78 (m, 3H, 3 x NH), 4.70 (m, 3H, 3 x CHN), 3.80 (s, 9H, 3 x OMe), 3.5-3.0 (m, 18H, 9 x CH\(_2\)), 1.46 (s, 27H, 3 x CMe\(_3\)). Anal. calcd. for C\(_{42}\)H\(_{63}\)N\(_3\)O\(_2\)S\(_3\) (1042.15): C, 48.40; H, 6.09; N, 4.03%. Found: C, 48.37; H, 6.11; N, 4.01%.

Thermolysis of sulfoxides 3a-d in neat methyl acrylate. General procedure. A 0.1 M solution of 3 in methyl acrylate was maintained under stirring at RT until full consumption of the starting sulfoxide (TLC control). The reaction crude was purified by flash column chromatography on silica gel (EtOAc / petroleum ether from 1:9 to 1:1).

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References

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