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DOI:

[10.1021/ol702212n](https://doi.org/10.1021/ol702212n)

[Link to publication record in Manchester Research Explorer](#)

Citation for published version (APA):

Burés, J., Isart, C., & Vilarrasa, J. (2007). Efficient Preparation of N-Phenylsulfenyl Ketimines from Oximes or Nitro Compounds without Racemization of -Stereocenters. *Organic Letters*, 9, 4635-4638.
<https://doi.org/10.1021/ol702212n>

Published in:

Organic Letters

Citing this paper

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Efficient Preparation of *N*-Phenylsulfenyl Ketimines from Oximes or Nitro Compounds without Racemization of α -Stereocenters

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Received September 8, 2007

ABSTRACT

As *N*-sulfenyl imines (e.g., $\text{RR}'\text{C}=\text{N}-\text{SAr}$) can be readily transformed to their *N*-sulfenyl imines ($\text{RR}'\text{C}=\text{N}-\text{SOAr}$), *N*-sulfonyl imines ($\text{RR}'\text{C}=\text{N}-\text{SO}_2\text{Ar}$), and *N*-sulfonyl oxaziridines, the very mild procedure developed to convert ketoximes and secondary nitro derivatives to *N*-arenesulfenyl ketimines constitutes a new and efficient route to all these series of compounds. The configuration of the α -stereocenters is retained.

N-Sulfenyl imines^{1,2} and *N*-sulfonyl imines^{3,4} are enjoying an increasing number of applications in asymmetric synthesis (Mannich reactions, α -alkylation via enamine anions, hetero-Diels–Alder reactions, etc.). Chiral *N*-sulfonyl oxaziridines

are also very popular as asymmetric epoxidation and hydroxylation reagents.⁵ As known,⁶ *N*-sulfenyl imines such as sulfenimines **1** are easily oxidized with *m*-CPBA or other peroxyacids to *N*-sulfenyl derivatives (sulfenimines **2**), subsequently to their *N*-sulfonyl derivatives (sulfonimines **3**), and then to *N*-sulfonyl oxaziridines (**4**);⁶ therefore, any efficient entry to **1** would be extremely useful.⁷ This is simplified in Scheme 1, where only one stereoisomer is drawn for each species.⁸ We have focused our attention on ketimines, which have been studied much less than aldimines.

In fact, the success of many asymmetric reactions involving **2–4** relies upon their preparation as stereopure substrates and their configurational stability. Usually, *N*-sulfenyl ketimines are prepared by condensation of ketones and sulfenamides (RSONH_2 or ArSONH_2) mediated by $\text{Ti}(\text{OEt})_4$ or other

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(2) For very recent, representative papers, see: (a) Davis, F. A.; Song, M. *Org. Lett.* **2007**, 9, 2413. (b) Davis, F. A.; Zhang, Y.; Qiu, H. *Org. Lett.* **2007**, 9, 833. (c) Tanuwidjaja, J.; Peltier, H. M.; Ellman, J. A. *J. Org. Chem.* **2007**, 72, 626. (d) Pei, D.; Wang, Z.; Wei, S.; Zhang, Y.; Sun, J. *Org. Lett.* **2006**, 8, 5913. (e) Xiao, X.; Wang, H.; Huang, Z.; Yang, J.; Bian, X.; Qin, Y. *Org. Lett.* **2006**, 8, 139. (f) Peltier, H. M.; Ellman, J. A. *J. Org. Chem.* **2005**, 70, 7342. (g) Lanter, J. C.; Chen, H.; Zhang, X.; Sui, Z. *Org. Lett.* **2005**, 7, 5905. (h) McMahon, J. P.; Ellman, J. A. *Org. Lett.* **2005**, 7, 5393. (i) García-Ruano, J. L.; Topp, M.; López-Cantarero, J.; Alemán, J.; Remuñán, M. J.; Cid, M. B. *Org. Lett.* **2005**, 7, 4407.

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(6) Historical review: Davis, F. A. *J. Org. Chem.* **2006**, 71, 8993.

(7) In a very recent example, in which condensation of a sulfenamide with a carbonyl group of an avermectin failed, the authors took advantage of the oxime-to-sulfenimine conversion. See: Lamy, E.; Lüthi, P.; Paturel, C.; Winkler, T.; Jung, P. M. J. *Tetrahedron Lett.* **2006**, 47, 5657.

(8) In addition, R and/or R' may contain stereogenic centers.

Scheme 1. From Sulfenyl Imines to *N*-Sulfonyl Oxaziridines

dehydrating agents in refluxing THF⁹ but also by the above-mentioned oxidation of *N*-sulfenyl ketimines and by the reaction¹⁰ of metal iminides with sulfinates (RSOOR'). *N*-Sulfonyl ketimines are mainly obtained by condensation (with limitations) of sulfonamides and ketones,^{3,4} from oximes and sulfonyl cyanides,¹¹ and by oxidation of *N*-sulfenyl ketimines.¹²

We uncover a very mild method that gives practically quantitative yields of the desired sulfenimines **1** at room temperature (rt) from ketoximes and from secondary nitro compounds. It is a significant practical improvement with regard to the reaction of oximes with PBU₃/PhSSPh reported by Lukin and Narayanan;^{13a} these authors showed that sulfenyl ketimines are intermediates in the conversion of oximes to imines^{13b} and can be cleaved in the presence of suitable acids. Our procedure can be very useful when the direct condensation to obtain **2** and **3** fails⁷ because of the steric hindrance or when it is counter-indicated as concomitant reactions (including stereocenter inversions) take place in the R or R' chains.

When the oximes in Table 1 (usually equilibrium *Z/E* mixtures) were treated with commercially available *N*-(phenylsulfonyl)phthalimide, that is, *N*-(phenylthio)phthalimide (PhthN-SPh, a non-stinking solid) and trimethylphosphine (PMe₃) at rt, *N*-phenylsulfenyl ketimines **1a–g** were formed quickly. Direct separation and purification of the final mixtures through a short pad of alumina, with hexane as the eluent, afforded excellent isolated yields (86–97%, see Table 1), with no stench, as PhSH was not formed as a co-product or during the workup. α -Stereocenters did not epimerize (entry 5), as expected, or did not racemize at all (entries 6 and 7), as checked for **1f** and **1g** by oxidation to the known, corresponding ketones with oxone (Oxone, 2KHSO₅·KHSO₄·K₂SO₄) or with ozone;¹⁴ the enantiomeric purities of both ketones were confirmed by polarimetry and chiral HPLC (Chiralpak AD-H column).

In parallel experiments with PBU₃ (220 mol %), only 70% of **1a** was obtained after 15 h. With an excess of an aromatic

Table 1. Conversion of Ketoximes to Sulfenyl Ketimines^a

entry	ketoxime ^b	product ^b	yield (%)
1			96
2			92
3			98
4			94
5			91
6			90
7			86

^a The oxime (1.0 mmol) was added to a commercially available solution of PMe₃ in THF (1.0 M, 1.1 mL). PhthN-SPh (1.2 mmol) was added, and the mixture was stirred at rt for 30 min. ^b The *Z/E* ratios are those observed (400 Mz ¹H NMR spectra) in CDCl₃ at rt. ^c Equilibrium mixture from **5g** prepared in pyridine. When the oxime was prepared from NH₃OH⁺Cl[−] and NaHCO₃ in MeOH–H₂O, isomer *Z* largely predominated.

phosphine, such as PPh₃ or 1,1'-bis(diphenylphosphine)-ferrocene (dppf), no reaction was observed at rt after 15 h.

To achieve a total absence of starting materials in the final products, we had to use stoichiometric amounts of PhthN-SPh. For example, with 0.4 equiv of PhthN-SPh, only 40% of the oximes were converted to sulfenimines; with 0.6 equiv of PhthN-SPh, ca. 60% of conversion occurred. Turnover did not take place, even with an excess of PMe₃.

The method partially failed in one case (Scheme 2). When we applied it to oxime **5h**, the yield of sulfenimine **1h**

Scheme 2. Fragmentation Byproducts

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(13) (a) Lukin, K. A.; Narayanan, B. A. *Tetrahedron* **2002**, *58*, 215. For other related pioneering works (also with PBU₃/PhSSPh), see: (b) Barton, D. H. R.; Motherwell, W. B.; Simon, E. S.; Zard, S. Z. *J. Chem. Soc., Perkin Trans. I* **1986**, 2243 and references therein.

(14) The oxidation of sulfenyl imines with oxone or with ozone was quantitative, and no epimerization of α -stereocenters was observed.

reached a maximum of 70%. A fragmentation product (the thioacetal shown in Scheme 2) was always formed, even at 0 and -20°C , in 20–30% yields. It may come from the decomposition of the common intermediate via a benzyl-type cation, which may be trapped by PhSH. Thus, prone to fragmentation oximes (on protonation or by reaction with electrophiles, giving rise to stable carbenic or oxocarbenic cations) may not afford high yields of sulfenimines **1**.

Even in this last case (**1h**), in which the α -stereocenter position is benzylic, no racemization took place. In fact, the oxidation of **1h** with oxone gave the corresponding enantiopure ketone, as shown by chiral HPLC.

Secondary nitro groups (**6**) can also be converted to sulfenimino groups (**1**) at rt by the same procedure, using 2.2 equiv of PMe_3 instead of 1.1 equiv, as shown in Table 2. One equivalent of PMe_3 is consumed in the first step, that

Table 2. From Nitro Compounds to Sulfenyl Ketimines^a

$\text{R}-\text{CH}(\text{NO}_2)-\text{R}' \xrightarrow[\text{THF, rt, 3 h}]{\text{PMe}_3 (2.2), \text{PhthN-SPh (1.2)}} \text{R}-\text{CH}=\text{N-SPh}-\text{R}'$			
entry	nitro compd ^b	product ^b	yield (%)
1			95
2			90
3			96
4			91
5			94
6			85
7			84 ^c

^a The nitro compound (1.0 mmol) was added to a solution of PMe_3 in THF (1.0 M, 1.1 mL). PhthN-SPh (1.2 mmol) was added, and the mixture was stirred at rt for 30 min. ^b The Z/E ratios are those observed (400 Mz ^1H NMR spectra) in CDCl_3 at rt. ^c With 3 equiv of PhthN-SPh and 6 equiv of PMe_3 for 24 h.

is, the reduction of $\text{R}_2\text{CH}-\text{NO}_2$ to $\text{R}_2\text{CH}-\text{N}=\text{O}$ / $\text{R}_2\text{C}=\text{N}-\text{OH}$ catalyzed by PhthN-SPh, which is slower than the second step, the sulfenylation of the oxime group. After 1–3 h of reaction, sulfenimines **1** were isolated in excellent yields. Stereocenter α of **1m** did not epimerize.

To our knowledge, this is the first reported method for obtaining *N*-sulfenyl imines directly from nitro compounds.

Moreover, the reaction works in one pot, at rt, in a short time, and using very small amounts of commercially available reagents. The only exception was converting **6m**¹⁵ to **1m**, as 3.0 equiv of *N*-(phenylsulfonyl)phthalimide, 6.0 equiv of PMe_3 , and 24 h were needed to obtain a good yield.

To gain more insight into the mechanism, the reaction of nitrocyclohexane (**6a**) with 3 equiv of PMe_3 and 0.2 equiv of PhthN-SPh was followed by ^{13}C NMR spectroscopy in THF (Figure 1).¹⁶ For the sake of simplification, only the

Figure 1. (a) ^{13}C NMR spectra of nitrocyclohexane (**6a**) and excess of PMe_3 (3 equiv) in THF. (b–e) After successive additions of 0.2 equiv of PhthN-SPh, only the appearance of sulfenimine **1a** was observed. (f) Addition of 0.5 equiv of PhSH to (e); spectrum registered after 10 min. (g) Spectrum registered 30 min after the addition of 0.5 equiv of PhSH.

signals of the methylene carbons vicinal to the $\text{CH}-\text{NO}_2$ / $\text{C}=\text{NOH}$ / $\text{C}=\text{NSPh}$ groups are shown. The disappearance of the oxime intermediate was so quick that it could not be detected under these conditions.

However, even with large amounts of PMe_3 , sulfenimine (**1a**) was not cleaved. In fact, acidic species must be present

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(16) For an excellent description of No-D NMR, see: Hoye, T. R.; Eklov, B. M.; Ryba, T. D.; Voloshin, M.; Yao, L. *J. Org. Lett.* **2004**, 6, 953.

in the reaction media or water must be added to catalyze or mediate such a N–S bond cleavage.^{13a}

Thus, “the secret of the success” is that the phthalimide anion of [PhthN–PMe₃(SPh)⁺] traps the oxime proton, but PhthNH, in contrast to ArSH, is not acidic enough to help the conversion of sulfenimines **1** to ketimines.

It is likely that many other aromatic and heteroaromatic phthalimide derivatives (PhthN-SAr or PhthN-SHet) may behave similarly. On the other hand, PhthN-S^tBu does not work, as no reaction with nitroalkanes or oximes was observed under our conditions; thus, our method is not useful for the preparation of Ellman’s substrates.^{1c}

In summary, efficient and mild conditions (short times, rt, 84–98% yields) for the conversion of ketoximes and secondary nitro compounds to sulfenyl ketimines (**1**) have been uncovered. No epimerization of α -stereocenters takes place. The role of PMe₃ is outstanding (in relation to Bu₃P and aromatic phosphines). In principle, a plethora of aryl-sulfinyl ketimines (**2**), arylsulfonyl ketimines (**3**), and aryl-sulfonyl oxaziridines (**4**) are available via the new route.

Acknowledgment. The Ministerio de Educación y Ciencia of Spain (Madrid) is acknowledged for the grants SAF02-02728 and CTQ2006-15393 as well as for a studentship to J.B. The Universitat of Barcelona is acknowledged for a studentship to C.I. The Generalitat de Catalunya (Barcelona) contributed also partially (grant 2001SGR065, 2002–2005, Grup de Síntesi Estereoselectiva d’Antibiòtics i Antivírics). The ketone of **5f** was prepared by Carmen Jiménez, of our Department. The senior author dedicates this work to Prof. Joan Bosch (Facultat de Farmàcia, Universitat de Barcelona) on the occasion of his 60th birthday.

Supporting Information Available: Experimental procedures and NMR spectra of the new compounds. This material is available free of charge via the Internet at <http://pubs.acs.org>.

OL702212N