

Metal-Free Solvent Promoted Oxidation of Benzylic Secondary Amines to Nitrones with H₂O₂

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Cite This: *J. Org. Chem.* 2021, 86, 13817–13823

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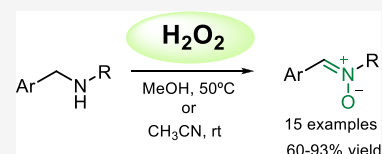
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ABSTRACT: An environmentally benign protocol for the generation of nitrones from benzylic secondary amines via catalyst-free oxidation of secondary amines using H₂O₂ in MeOH or CH₃CN is described. This methodology provides a selective access to a variety of C-aryl nitrones in yields of 60 to 93%. Several studies have been performed to shed light on the reaction mechanism and the role of the solvent.



The development of highly efficient and environmentally friendly methodologies for the preparation of nitrones is of great importance since this kind of compounds are valuable synthetic intermediates and useful scaffolds in drug discovery.¹ Nitrones are found in numerous natural products² and many studies have demonstrated the interest of benzylic nitrones as therapeutic agents for several pathologies including atherosclerosis, septicaemia, stroke, and Alzheimer³ (Figure 1).

In addition, nitrones have been used as ligands in organometallic chemistry⁴ and as spin traps in biological studies.⁵ The diastereo- and enantioselective nucleophilic additions to nitrones is a fundamental tool in organic synthesis.⁶ Furthermore, the 1,3-dipolar cycloaddition reaction of nitrones with alkenes has become one of the methods of choice for the preparation of isoxazolidines, and a wide variety of natural products has been prepared using this reaction as key step.⁷

Among the available methodologies for the preparation of nitrones, the condensation of carbonyl compounds with hydroxylamines has arguably been the most used.⁸ However, this procedure presents several limitations, such as the availability of the hydroxylamines and the low reactivity observed using ketones as carbonyl partner. Otherwise, nitro compounds can be used as an alternative to hydroxylamines under reductive conditions.⁹

Another main process for the synthesis of nitrones consists of the oxidation of secondary hydroxylamines,¹⁰ imines (either preformed¹¹ or generated in situ from primary amines and aldehydes¹²), isoxazolidines,¹³ *N*-alkyl- α -amino acids¹⁴ (via regioselective decarboxylative oxidation), and secondary amines. The great availability of secondary amines has made the last option one of the most convenient. In fact, this methodology has been used for preparative scale and as key step in the synthesis of several natural products.¹⁵

Among all the oxidants used for nitrone synthesis from amines, hydrogen peroxide is one of the most attractive for the development of environmentally friendly processes since water is the only by product of its reduction. In 1984, Murahashi and co-workers reported the first example of catalytic direct

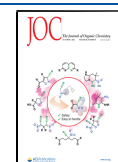
oxidation of secondary amines to nitrones with H₂O₂ in the presence of Na₂WO₄/H₂O.¹⁶ Since then, several general and efficient procedures have been developed using hydrogen peroxide or its urea complex (UHP) as oxidant in combination with different catalysts such as SeO₂,¹⁷ methyltrioxorhenium,¹⁸ and titanium¹⁹ or platinum²⁰ complexes. In addition, several heterogeneous catalysts have also been used.²¹ Alternatively, the reaction can be also carried out in the presence of alkyl hydroperoxides,²² oxone,²³ dimethyldioxirane,²⁴ *m*-CPBA,²⁵ Davis oxaziridine,²⁶ or molecular oxygen²⁷ as oxidant.

To the best of our knowledge, most of the procedures described to date for the preparation of nitrones from amines using H₂O₂ as oxidant require the presence of a metal catalyst, which is usually expensive and present environmental problems. Herein, we report a facile and clean catalyst-free oxidation protocol for the efficient preparation of nitrones from benzylic secondary amines using hydrogen peroxide as oxidant.

1,2,3,4-Tetrahydroisoquinoline was selected as the model substrate for the oxidation process. This substrate is useful for comparative purposes since its oxidation to nitrone with the combination of different oxidants and catalytic systems has been extensively studied.¹ Initially, the reaction was performed in the presence of four equivalents of H₂O₂ 30% v/v in MeOH at room temperature, leading to the formation of the nitrone with a 27% of conversion after 24 h (Table 1, entry 1). An increase of the equivalents of hydrogen peroxide leads to complete conversion (entries 2 and 3). Higher reactivity was observed at 50 °C allowing reducing the amount of H₂O₂ from 10 to 4 equiv and the reaction time to 12 h (entry 4). Further decreasing the amount of hydrogen peroxide revealed that the

Received: August 7, 2021

Published: September 16, 2021



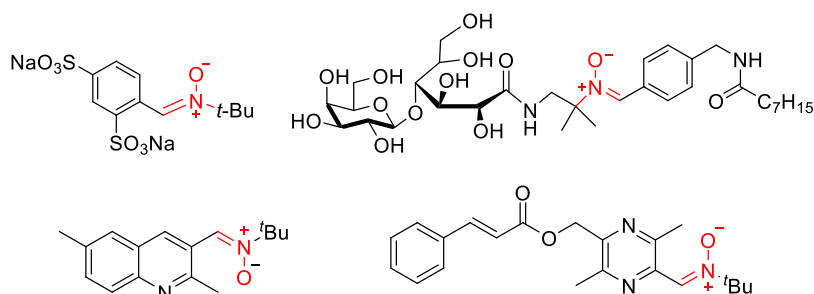


Figure 1. Examples of biologically active nitrones.

Table 1. Optimization of the Reaction Conditions

entry	equiv	T (°C)	solvent	time (h)	conversion (%) ^a
1	4	rt	MeOH	24	27 ^c
2	8	rt	MeOH	24	60
3	10	rt	MeOH	24	>99
4	4	50	MeOH	12	>99 (91) ^b
5	2	50	MeOH	12	45 ^c
6	3	50	MeOH	12	73 ^c
7	4	50	EtOH	12	>99
8	4	50	CH ₂ Cl ₂	24	
9	4	50	toluene	24	
10	4	rt	CH ₃ CN	2	>99 (93) ^b
11	2	rt	CH ₃ CN	2	>99 (90) ^b
12	2	rt	DMF	12	

^aDetermined by ¹H NMR in the crude reaction mixture. ^bYield after chromatographic purification. ^cNitrone is the only observed product in the ¹H NMR spectra.

reaction proceeded with a significant erosion of the reactivity (entries 5 and 6). Similar results were obtained using EtOH as solvent (entry 7). On the other hand, using less polar aprotic solvents, such as CH₂Cl₂ or toluene, no conversion was observed (entries 8 and 9). Interestingly, a very significant improvement of the reactivity was observed using CH₃CN as solvent, complete conversion was achieved in only 2 h at room temperature (entry 10). A similar outcome was observed using only 2 equiv of H₂O₂ (entry 11). Finally, no reaction was observed using a solvent with similar dielectric constant such as DMF (entry 12).

With the optimized reaction conditions on hands, we then investigated the scope of this oxidation reaction (Table 2). A remarkably broad range of benzylic secondary amines could be converted into the corresponding nitrones with good yields using either MeOH (conditions A) or CH₃CN (conditions B) as solvent.

In general, better reactivity and slightly superior yields were observed using CH₃CN as solvent in most of the examples studied. The influence of electronic character of the substituents was first evaluated. Tetrahydroisoquinolines **1b** and **1c** with strong electron donating substituents like methoxy afforded the corresponding nitrones with high yields both using conditions A or B (Table 2, entries 1 and 2). The reaction also tolerates electron withdrawing groups. For instance, 6-nitrotetrahydroisoquinoline effectively provides the desired nitrone **2d** in high yield (75% conditions A or 81% conditions B; entry 3). 1,2,3,4-Tetrahydroisoquinolines

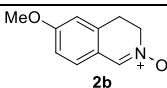
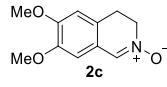
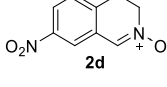
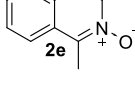
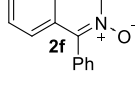
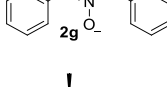
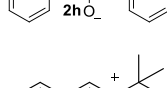
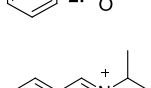
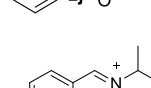
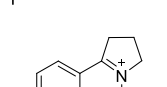
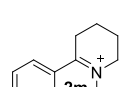
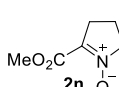
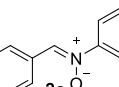
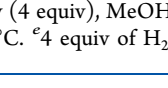
with alkyl (**2e**) or aryl (**2f**) substituents at position 1 were selectively oxidized in the benzylic more substituted position to the corresponding nitrone derivatives under both conditions (Table 2, entries 4 and 5). Dibenzylamine **1g** was cleanly converted to nitrone **2g** in 71% (MeOH) or 73% (CH₃CN) yield (entry 6). Interestingly, this methodology also allowed the straightforward preparation of chiral nitrone **2h**,²⁸ which has been extensively employed in diastereoselective 1,3-dipolar cycloadditions (entry 7). In this example, under both conditions, the reaction takes place in the less hindered site, suggesting kinetic control. Acyclic *N*-benzyl-*N*-alkyl substituted amines **1i,j,k** were selectively oxidized only on benzylic position to nitrones **2i,j,k** in good yields (entries 8, 9, and 10), although it is required to carry out the reaction at 50 °C in both solvents. The oxidation of 2-phenylpyrrolidine **1l** and 2-phenylpiperidine **1m** also proceeded efficiently, leading to nitrones **2l** and **2m** in comparable yields (entries 11 and 12). Benzylic secondary amine **1n** was also a suitable substrate, albeit the process occurred with a somewhat lower yield. No formation of nitrone **2o** was observed when less nucleophilic *N*-benzylaniline was tested under the same reaction conditions and most of the starting material was recovered unaltered. Dialkylamines are not suitable substrates for this transformation, the reaction did not occur with cyclic (piperidine) or acyclic (dioctylamine) substrates. In these examples, complex reaction mixtures were obtained under optimized reaction condition using MeOH or CH₃CN as solvent.

To demonstrate the robustness and the synthetic utility of the method, we scaled up the oxidation reaction either in CH₃CN or MeOH using 15 mmol of tetrahydroisoquinoline **1**. In both cases, the desired nitrone **2** was isolated in excellent yields (Scheme 1, eq A). The reaction can also be carried out using the urea-hydrogen peroxide adduct (UHP), a safe source of hydrogen peroxide. UHP is cheap, easy to handle, and can be stored for long periods without any change of the oxygen content^{29,11a} (Scheme 1, eq B).

Hydrogen peroxide has been extensively used as primary oxidant in tertiary amine oxidations under either heterogeneous or homogeneous catalytic conditions.²⁰ Nevertheless, the reaction of tertiary amine **3** or electron richer trialkylamine **4** with H₂O₂ in MeOH at 50 °C did not show the *N*-oxide formation (Scheme 2, eq A). Taking advantage of this chemoselectivity, a secondary amine could be selectively oxidized to nitrone in the presence of a tertiary amine. Thus, oxidation of tetrahydroisoquinoline **1o** exclusively afforded nitrone **2o** in 79% yield (Scheme 2, eq B).

Next, to gain some insights into the reaction mechanism some experiments were performed. It is well established that hydrogen peroxide could be activated toward nucleophilic attack by the formation of a hydrogen bond.³⁰ We

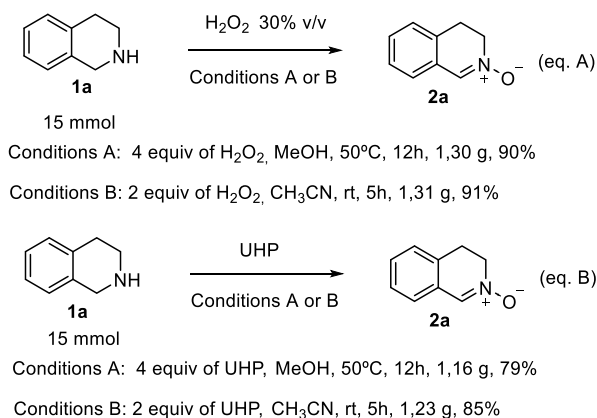
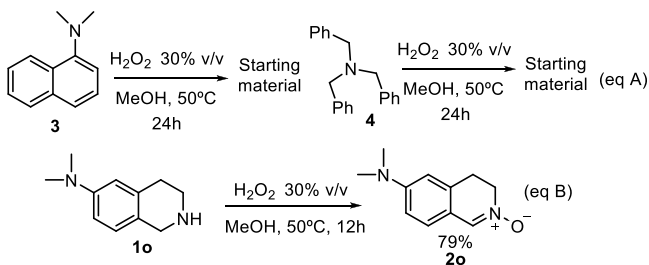
Table 2. Scope of Oxidation of Benzylic Secondary Amines to Nitrones

Entry	Product	Conditions A ^a Yield (%) ^c	Conditions B ^b Yield (%) ^c	Time (h)
1		82	85	2
2		75	76	2
3		75	81	2
4		71	77	4
5		79	83	4
6		71	73 ^{d,e}	12 ^{d,e}
7		69	71 ^{d,e}	16 ^{d,e}
8		70	73 ^d	16 ^d
9		83	87 ^d	16 ^d
10		82	89 ^d	16 ^d
11		60	75 ^e	12 ^e
12		62	73 ^e	12 ^e
13		30	62	12 ^e
14		--	--	12 ^{d,e}

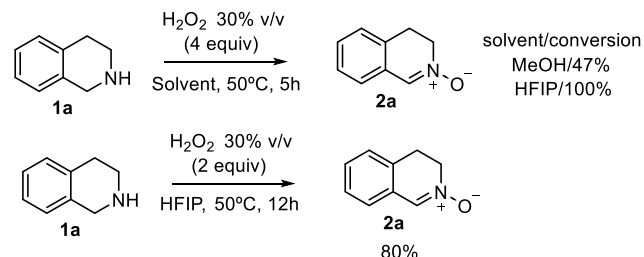
^aConditions A: H₂O₂ 30% v/v (4 equiv), MeOH, 50 °C, 12 h. ^bConditions B: H₂O₂ 30% v/v (2 equiv), CH₃CN, rt. ^cYield after chromatographic purification. ^dReaction at 50 °C. ^e4 equiv of H₂O₂ is used.

hypothesized that H₂O₂ could be electrophilically activated by MeOH, the OH moiety of the solvent forms a hydrogen bond with H₂O₂ increasing the electrophilic character of the oxygen.

Accordingly, H₂O₂ did not oxidize secondary amines in aprotic solvents such as CH₂Cl₂ or toluene. However, using UHP as hydrogen peroxide source the reaction can be performed in

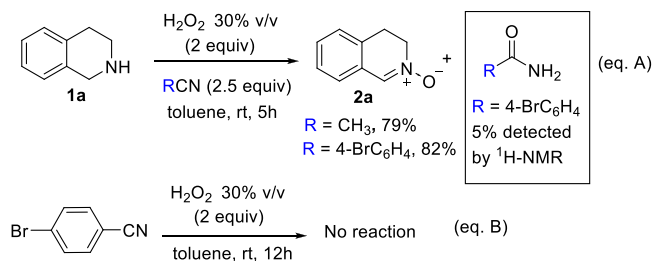
Scheme 1. Scale-up of the Oxidation of 1a and Use of UHP as the Source of Hydrogen Peroxide**Scheme 2. Selective Oxidation of Secondary Amines in the Presence of Tertiary Amines**

toluene probably because the urea is able to activate H₂O₂ by hydrogen bonding formation. Furthermore, reactions in hexafluoro 2-propanol (HFIP) are faster than in MeOH since the hydroxyl proton of HFIP forms a stronger hydrogen bond because of the electron-withdrawing character of CF₃ group.³¹ Interestingly, the use of HFIP as solvent allowed the reduction of the number of equivalents of hydrogen peroxide from 4 to 2 without erosion in reactivity. However, only 40% of conversion was observed when the reaction was performed at room temperature (Scheme 3).

Scheme 3. Reaction Using HFIP as Solvent

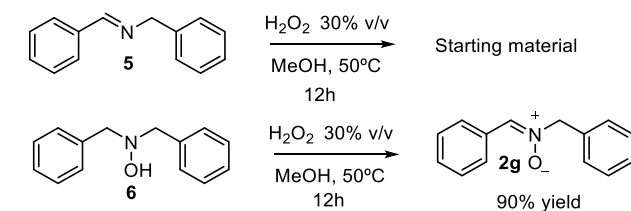
On the other hand, it has been reported that the rate of several oxidation reactions using aqueous H₂O₂ is significantly increased in the presence of a nitrile in basic media via the formation of a peroxyimide intermediate which rapidly reacts with the secondary amine to afford the corresponding nitron and acetamide.³² The oxidation of 1a in acetonitrile as solvent was monitored by ESI-MS detecting small amounts of the ion [M + H⁺] (60.0446 *m/z*) that correspond to acetamide. This oxidation process provides similar results using only 2.5 equiv of acetonitrile or 4-bromobenzonitrile in

toluene as solvent (Scheme 4, eq. A). However, only 5% of conversion of the 4-bromobenzonitrile into the 4-bromophe-

Scheme 4. ESI-MS Experiment and Reaction with 2.5 equiv of Nitrile

nylacetamide was observed in the crude ¹H NMR. In addition, 4-bromobenzonitrile was recovered unaltered after 12 h of reaction with 2 equiv of H₂O₂ in toluene at room temperature (Scheme 4, eq. B). These results suggest that the peroxyacetimidic acid is not the major oxidizing species in these reactions.

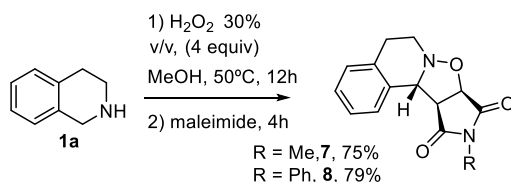
It has been proposed that the oxidation of secondary amines to nitrones is a two-step sequence involving an initial formation of a hydroxylamine followed by oxidation of the latter to nitron.²⁰ Alternatively, nitrones can also be prepared by oxidation of imines.^{11,12} In our case, the reaction of imine 5 under the optimized oxidation conditions did not give the corresponding nitron, recovering the starting material together with degradation products. On the other hand, the oxidation of commercially available dibenzylhydroxylamine 6 took place with complete conversion to the corresponding nitron 2g. These results suggested that the hydroxylamine and not the imine is the intermediate in the reaction pathway (Scheme 5).

Scheme 5. Control Experiments

As mentioned before, 1,3-dipolar cycloaddition of nitrones is one of the most straightforward methodologies for the preparation of isoxazolidines. We next studied the possibility of carrying out a one pot 1,3-dipolar cycloaddition of the obtained nitrones with alkenes. Tetrahydroquinoline 1a was treated with H₂O₂ in MeOH at 50 °C for 12h, subsequent addition of *N*-methyl or *N*-phenyl maleimide to the reaction mixture afforded the corresponding cycloadduct *exo*-7 or *exo*-8, as a single diastereomer, in high yield, after 4 h.³³ (Scheme 6).

We have developed a novel procedure for the selective oxidation of benzylic secondary amines to nitrones using H₂O₂ as the sole oxidant in MeOH or CH₃CN. An important advantage of this methodology is that the reaction can be performed under mild reaction conditions without any catalyst or additive. It is also possible to carry out the reaction using UHP as a safe source of anhydrous hydrogen peroxide. Remarkably, the system allows the selective oxidation of

Scheme 6. 1,3-Dipolar Cycloaddition



secondary amines in the presence of tertiary amines. Several studies were performed in order to shed light on the ability of MeOH and CH_3CN to activate H_2O_2 .

■ EXPERIMENTAL SECTION

General Methods. Dichloromethane and toluene were dried over the PureSolv MD purification system. Reactions were monitored by thin-layer chromatography carried out on 0.25 mm silica gel plates (230–400 mesh). Flash column chromatographies were performed using silica gel (230–400 mesh). NMR spectra were recorded on AU-300 MHz instrument and calibrated using residual undeuterated solvent (CDCl_3) as internal reference. MS spectra were recorded on a VG AutoSpec mass spectrometer.

All the chromatographic columns were carried out using deactivated silica gel. *Deactivated silica gel preparation:* Et_3N (5 mL) was added to a suspension of 300 g of silica gel in cyclohexane; the mixture was stirred for 1 h, filtered, and dried under reduced pressure on a rotary evaporator.

Nitrones **2a**,²³ **2b**,³⁴ **2c**,³⁵ **2d**,³⁶ **2e**,³⁷ **2f**,³⁸ **2g**,²³ **2h**,³⁹ **2i**,⁴⁰ **2j**,⁴¹ **2l**,⁴² **2m**,⁴³ and **2n**⁴² and cycloadducts **6** and **7**³³ have been previously described. Spectroscopic data match those previously reported.

General Procedure 1 (Conditions A). To a stirred solution of amine (1 mmol) in MeOH (3 mL) H_2O_2 30% v/v (4 mmol, 453 μL) was added. The resulting solution was stirred at 50 °C (oil bath) for 12 h, after cooling at room temperature CH_2Cl_2 (10 mL) and water (10 mL) were added. The organic layer was separated, and the aqueous phase was extracted with dichloromethane (10 mL). The combined organic layers were washed with brine, dried over MgSO_4 , and evaporated under reduced pressure. The crude mixture was purified by flash column chromatography over deactivated silica gel to afford the corresponding nitrone.

General Procedure 2 (Conditions B). To a stirred solution of amine (1 mmol) in CH_3CN (3 mL) H_2O_2 30% v/v (2 mmol, 227 μL) was added. The resulting solution was stirred at room temperature for the time indicated in Table 2, and CH_2Cl_2 (10 mL) and water (10 mL) were added. The organic layer was separated, and the aqueous phase was extracted with dichloromethane (15 mL). The combined organic layers were washed with brine, dried over MgSO_4 , and evaporated under reduced pressure. The crude mixture was purified by flash column chromatography over deactivated silica gel to afford the corresponding nitrone.

N-(4-Fluorobenzylideneisopropylamine N-Oxide (2k). Following the general procedure A, the reaction of *N*-(4-fluorobenzyl)-2-propanamine (**1k**) (217 mg, 1.30 mmol) with H_2O_2 (5.2 mmol, 554 μL) in MeOH (4 mL) at 50 °C (oil bath) afforded after purification by silica gel flash chromatography (EtOAc) the nitrone **2k** (193 mg, 82%, yellow oil). Following the general procedure B, the reaction of *N*-(4-fluorobenzyl)-2-propanamine (**1k**) (250 mg, 1.50 mmol) with H_2O_2 (3 mmol, 340 mL) in CH_3CN (4 mL) at rt, afforded after purification by silica gel flash chromatography (EtOAc) the nitrone **2k** (241 mg, 89%, yellow oil). ^1H NMR (300 MHz, CDCl_3): δ 8.34–8.32 (m, 2H), 7.49 (s, 1H), 7.25–7.23 (m, 2H), 4.24 (sep, J = 7.1 Hz, 1H), 1.57 (d, J = 7.1 Hz, 6H). $^{13}\text{C}\{^1\text{H}\}$ NMR (75 MHz, CDCl_3): δ 164.8, 161.5, 130.7, 130.6, 115.6, 115.4, 67.7, 20.9. HRMS (TOF MS EI+): calculated for $\text{C}_{10}\text{H}_{12}\text{NOF}$, 181.0903; found, 181.0902 ($[\text{M}]^+$, 56).

6-Dimethylamino-3,4-Dihydroisoquinoline N-Oxide (2o). Following the general procedure A, the reaction of 6-(dimethylamino)-1,2,3,4-tetrahydroisoquinoline (174 mg, 1 mmol) with H_2O_2 (4 mmol, 453 μL) in MeOH (4 mL) at 50 °C (oil bath) afforded after

purification by silica gel flash chromatography ($\text{CH}_2\text{Cl}_2/\text{MeOH}$ 9/1) the nitrone **2o** (150 mg, 79%, yellow oil). ^1H NMR (300 MHz, CDCl_3): δ 7.68 (s, 1H), 7.12–7.08 (m, 1H), 6.57–6.54 (m, 2H), 4.12 (t, J = 7.1 Hz, 2H), 3.15 (t, J = 7.1 Hz, 2H), 3.01 (s, 6H). $^{13}\text{C}\{^1\text{H}\}$ NMR (75 MHz, CDCl_3): δ 151.1, 135.5, 131.5, 127.3, 116.0, 110.6, 57.2, 39.8, 28.2. HRMS (ESI+): Calculated for $\text{C}_{11}\text{H}_{15}\text{N}_2\text{O}$, 191.1181; found, 191.1179 ($[\text{M} + \text{H}]$, 100).

Typical Procedure for the Cycloaddition Reaction. Cycloadduct (7). To a stirred solution of tetrahydroisoquinoline **1a** (0.5 mmol, 66 mg) in MeOH (2 mL) H_2O_2 30% v/v (2 mmol, 277 μL) was added. The resulting solution was stirred at 50 °C (oil bath) for 12 h and *N*-methyl maleimide (0.5 mmol, 56 mg) was added. The reaction was stirred at 50 °C (oil bath) for 4 h, and CH_2Cl_2 (10 mL) and water (10 mL) were added, the organic layer was separated, and the aqueous phase was extracted with dichloromethane (10 mL). The combined organic layers were washed with brine, dried over MgSO_4 , and evaporated under reduced pressure. The crude mixture was purified by flash column chromatography ($\text{CH}_2\text{Cl}_2/\text{MeOH}$ 99/1) to afford nitrone **7** (97 mg, 75%, yellow oil). Spectroscopic data match those previously reported.³³

■ ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/acs.joc.1c01888>.

^1H and ^{13}C NMR and ESI/MS spectra for all compounds and solvent specifications (PDF)

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Notes

The authors declare no competing financial interest.

■ ACKNOWLEDGMENTS

We thank FEDER/Ministerio de Ciencia, Innovación y Universidades–Agencia Estatal de Investigación (Grant PGC2018-098660-B-I00) and CAPES (Finance Code 001), CNPq (310514/2018-5), FAPEMIG for financial support.

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