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Electrochemical ABNO-Mediated Alpha-Cyanation of Secondary Piperidines for Pharmaceutical Building Block Diversification

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Abstract

Secondary piperidines are ideal pharmaceutical building blocks owing to the prevalence of piperidines in commercial drugs. Here, we report an electrochemical method for cyanation of the heterocycle adjacent to nitrogen without requiring protection or substitution of the N–H bond. The reaction utilizes ABNO (9-azabicyclononane *N*-oxyl) as a catalytic mediator. Electrochemical oxidation of ABNO generates the corresponding oxoammonium species, which promotes dehydrogenation of the 2° piperidine to the cyclic imine, followed by addition of cyanide. The low-potential, mediated electrolysis process is compatible with a wide range of heterocyclic and oxidatively sensitive substituents on the piperidine ring and enables synthesis of unnatural amino acids.

TOC Graphic

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ASSOCIATED CONTENT

Supporting Information

Experimental procedures, reaction optimization, and product characterization data. The Supporting Information is available free of charge on the ACS Publications website.

Strategic building blocks are commonly used in drug discovery to create or modify bioactive core structures, and efficient methods for diversification of commercially available building blocks could have broad impact in medicinal chemistry. Piperidines are the most common heterocycle found in FDA-approved drugs owing to their favorable pharmacokinetic properties that contribute to improved clinical success (Scheme 1A). Secondary piperidines represent ideal targets for chemical modification, and effective methods could greatly expand this pool of available building blocks. Whereas C–H functionalization adjacent to nitrogen in cyclic amines has been the focus of considerable attention, the vast majority of precedents use *N*-protected (i.e., *N*-acyl) or *N*-aryl/alkyl 3° amine derivatives that are less versatile as pharmaceutical building blocks. Amine protection/deprotection hampers efficient utilization of the former substrates, while fixed substitution of the *N*-aryl/alkyl derivatives limits incorporation of these compounds into more complex structures. Here, we report an electrochemical method for oxidative α -cyanation of diverse 2° piperidines. The method is enabled by the use of ABNO (9-azabicyclononane *N*-oxyl) as a catalytic mediator that exhibits broad functional group compatibility.

Nitriles are versatile functional groups that are readily converted into other substituents, including carboxylic acids, amides, ketones and amines, among others. This versatility underlies the extensive efforts directed toward α -cyanation of piperidine and other amine derivatives (Scheme 1B). Synthetically useful precedents, however, are limited to activated substrates (e.g., tetrahydroisoquinolines) and the 3° amine/amide derivatives noted above. No general methods are available for analogous cyanation of 2° piperidines, reflecting the susceptibility of cyclic imines to undergo decomposition.

Precedents for electrochemical oxidation of amines typically feature 3° derivatives and/or are initiated by outer sphere single-electron transfer (ET). Subsequent rapid proton and electron transfer (PT-ET) affords an iminium ion. We recently showed that use of an aminoxyl mediator bypasses this conventional ET-PT-ET sequence by undergoing electrochemical oxidation to an oxoammonium species that promotes direct hydride transfer from the substrate. The aminoxyl-mediated reactions operate at much lower electrode potentials (by > 1 V), thereby greatly expanding the functional group compatibility and substrate scope. Here, we demonstrate that analogous principles may be applied to enable efficient α -cyanation of 2° piperidines bearing diverse pharmaceutically relevant substituents (Scheme 1C).

We initiated our studies by investigating the redox behavior of 4-phenylpiperidine **1a** by cyclic voltammetry (CV) in the absence and presence of ABNO (Figure 1). Substrate **1a** exhibits an irreversible anodic CV feature at 739 mV (Figure 1, green trace; all potentials are reported relative to Fc/Fc⁺), while ABNO, which was selected as a low-potential sterically unhindered aminoxyl mediator, exhibits a reversible CV trace with $E_{1/2} = 195$ mV (Figure 1, gray trace). CV analysis of a solution containing both **1a** and ABNO reveals a significant increase in the anodic feature at the ABNO redox potential, with a corresponding decrease in the cathodic feature (Figure 1, red trace). This behavior implicates reaction of ABNO⁺ with the substrate and electrochemical regeneration of ABNO⁺ on the CV time scale (cf. Scheme 1c). Complete disappearance of the anodic feature corresponding to **1a** under these conditions is consistent with consumption of the substrate from the electrode surface via reaction with ABNO⁺. The large difference between the anodic peak potentials of ABNO and **1a** ($E_p > 500$ mV; cf. green vs red trace) highlights the lower electrode potential that arises when substrate oxidation proceeds via ABNO-mediated hydride transfer, rather than electrode-initiated electron transfer.

Efforts then shifted to bulk electrolysis studies in order to explore optimal conditions for substrate cyanation. The reactions were conducted under constant-current conditions (1–3 mA) in an undivided cell with a graphite rod working electrode and Pt wire counter electrode. TMSCN was used as an easily handled source of cyanide nucleophile. Initial attempts to perform direct oxidation of the substrate at the electrode (i.e., in the absence of a mediator) resulted in only low yield of the desired product **2a** (19%, Table 1, entry 1). Inclusion of 10 mol% ABNO in the reaction, under otherwise identical conditions, nearly doubled the yield of **2a** (37%, entry 2). One equivalent of hexafluoroisopropanol (HFIP) was included in the reaction as a proton source to facilitate production of H₂ at the cathode (cf. BH⁺ in Scheme 1C), which also generates an alkoxide base that can promote proton-coupled oxidation of ABNO-H to ABNO⁺ at the anode (cf. Scheme 1C). Consistent with this hypothesis, the yield improved considerably upon adding HFIP, 1 equiv; 54% yield, entry 3) as an additive (see below for further discussion).

Testing of other aminoxyl mediators clearly indicated the importance of steric effects as the primary indicator of mediator effectiveness. Specifically, TEMPO and 4-acetamidoTEMPO (ACT) have higher redox potentials than ABNO, but they afforded lower yields of **2a** (entries 4 and 5). KetoABNO exhibited the highest yield, but, as its higher redox potential could interfere with functional-group compatibility, we explored further optimization of the reaction with ABNO as the mediator. Increasing the electrolysis current from 1 to 3 mA led to an increase in yield from 54% to 66% (entries 3 and 7). Further improvement was achieved by replacing HFIP with MeOH as the protic additive and reducing the stoichiometry to 0.5 equiv, ultimately affording 84% yield of **2a** (entries 7–10). To ensure reproducibility, the optimized reaction conditions were directly implemented in two commercially available bulk electrolysis cells, one from BASi and the ElectraSyn 2.0 unit from IKA, with the former conducted on 1 g scale (entries 11 and 12). Both reactions afforded **2a** in good yield.

Analysis of the constant current electrolysis traces provides valuable insights into the ABNO-mediated reactions (Figure 2). Under the optimized reaction conditions (red trace)

the electrode potential needed to sustain 3 mA current is low (approx. 150–250 mV) and is stable throughout the entire reaction. In the absence of MeOH (blue trace), a low potential is observed only when the charge passed is less than that needed for electrochemical oxidation of ABNO to ABNO⁺. The increase in potential beyond this point (to approx. 600–700 mV) is attributed to the lack of an effective Brønsted base (i.e., no methoxide is available) to support proton-coupled oxidation of ABNOH to ABNO⁺. In the absence of both MeOH and ABNO (green trace), a much higher electrode potential is needed to support the 3 mA current. The initial observed potential is similar to that expected from the CV studies (cf. Figure 1); however, this potential rises significantly during the reaction, possibly arising from fouling of the electrode by reactive intermediates generated in the direct electrolysis process. The reaction yield correlates inversely with the potential needed to sustain the reaction, demonstrating the benefit of pairing a mediator with an appropriate proton shuttle in the undivided cell.

The optimized conditions were then tested on a broad collection of commercially available 2° piperidines (Table 2A). 4-Substituted piperidines are especially readily available, and good-to-excellent yields of the corresponding α -cyanopiperidines were obtained for substrates bearing a diverse array of functional groups, including aryl, aryl halide, aryl ether, amide, ester, pyridine and related heteroaryl derivatives, and azoles, among other groups (**3a–3r**). The method even tolerates an unprotected 2° alcohol (**3x**), indicating that amine oxidation is favored over the well-established aminoxyl-mediated alcohol oxidation. The products were primarily isolated as the *p*-toluenesulfonic acid salts. Analytically pure electrolyte salt (TBAPF₆) was also recovered from the reaction mixture and reused. High diastereoselectivity, often reflecting only a single diastereomer, is observed in all cases where the substituent is large enough to bias the ring conformation (**3a–3p**, **3s**). The favored anti relationship between the cyano group and substituent was confirmed by nOe measurements and X-ray crystallography. The observed stereochemistry is rationalized by axial attack of cyanide on the intermediate imine and aligns with stereochemical models that have been reported by Houk and Cieplak. A mixture is observed in two cases when the substituent is smaller (**3x**, **3y**) and in a bicyclic pyrrolidine substrate (**3ad**). The latter result demonstrates that the method is applicable to heterocycles beyond piperidines, as further exemplified by the reaction of the parent pyrrolidine, (**3ac**), an azepane (**3ab**), and morpholine (**3u**). Piperidine substitution in the 2- and 3- positions leads to highly regioselective cyanation at the less hindered position, suggesting that steric accessibility overrides thermodynamic considerations (**3s**, **3v**, **3w**). This observation is also evident from the selectivity observed with the azepane derivative **3ab**, where the benzylic position is avoided in favor of the more accessible site.

For a substrate containing both an unprotected and a Boc-protected 2° piperidine, cyanation only occurred on the unprotected piperidine ring (**3q**). On the other hand, 2° piperidine substrates bearing 5- or 6-membered cyclic 3° alkylamine substituents undergo preferential cyanation on the alkylamine ring (**3z**, **3aa**), presumably reflecting the enhanced hydricity of the C–H bonds at this site. Ineffective substrates primarily corresponded to substrates that were not soluble in the reaction medium.

The relevance of this method for medicinal chemistry is evident from a number of substrates that feature functional groups directly corresponding to active pharmaceuticals and drug candidates in Scheme 1A. Examples include risperidone (**3j**), two receptor antagonists (**3p** and **3r**) and the HIV drug, Nelfinavir (**3i**). Each of the corresponding α -cyanation products was isolated in diastereomeric purity. The low potential and mild conditions that make this method applicable to building block diversification also make it amenable to late-stage functionalization, as revealed by the cyanation of the anti-depressant Amitifadine (**3ad**), which proceeded in 48% yield.

The hydrolysis of the nitriles to carboxylic acids provides an efficient means to generate non-natural amino acids. To demonstrate this concept, several cyanation products were subjected directly to hydrolysis conditions, without isolation, to generate the corresponding carboxylic acid derivatives in excellent yields (**4a–d**, Table 2B).

In summary, we have developed a highly effective, user-friendly method for electrochemical α -cyanation of 2° piperidines. This class of molecules represents an especially important class of pharmaceutical building blocks. Use of ABNO as a hydride-transfer mediator allows the reactions to proceed at low electrode potentials, thereby tolerating a broad array of important functional groups. C–H functionalization methods and related reactions of this type that enable direct building block diversification should have considerable utility in medicinal chemistry and drug discovery.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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Figure 1.

CVs (0.1 V/s) in MeCN (5 mL) and NaClO_4 (0.1 M) of: ABNO (1 mM) (gray trace); 4-phenyl piperidine 1a (2 mM) (green trace); ABNO (1 mM) and 4-phenyl piperidine (2 mM) (red trace).

Figure 2.

Bulk electrolysis in MeCN, TBAPF₆ (0.1 M), TMSCN (1.5 equiv.) and **1a** (0.3 mmol) with/without ABNO (10 mol%) and/or MeOH (0.5 equiv.). MB = mass balance.

[A] Prominent piperidine-containing bioactive molecules

[B] Piperidine cyanation:

[C] This work:

Scheme 1.

Context and Strategy for α -Cyanation of 2° Piperidines

Table 1.**Bulk Electrolysis Optimization Data.**

Entry	Pre-catalyst	Additive (equiv.)	Current (mA)	Yield/% ^a
1	—	—	1	19
2	ABNO	—	1	37
3	ABNO	HFIP ^b (1)	1	54
4	TEMPO	HFIP (1)	1	10
5	ACT	HFIP (1)	1	24
6	KetoABNO	HFIP (1)	1	81
7	ABNO	HFIP (1)	3	66
8	ABNO	H ₂ O (1)	3	68
9	ABNO	MeOH (1)	3	79
10	ABNO	MeOH (0.5)	3	84
11	ABNO	MeOH (0.5)	3	74 ^c
12	ABNO	MeOH (0.5)	2	74 ^d

^a NMR yields against a mesitylene internal standard. Isolated yield.

^b HFIP = hexafluoroisopropanol.

^c Reaction performed in commercial BASi bulk electrolysis cell on 1 g scale using RVC working electrode.

^d Reaction performed using ElectraSyn 2.0.

Table 2.

Substrate Scope for Electrochemical α -Cyanation of 2° Piperidines (A) and Cyanation/Hydrolysis to Generate Pipecolic Acids (B).^a

Reaction performed on 0.3 mmol scale. Isolated yields are reported.

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