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Transformation of *N,N*-Dimethylaniline *N*-Oxides into Diverse Tetrahydroquinoline Scaffolds via Formal Povarov Reactions

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Abstract

A one-pot protocol for the assembly of diversely functionalized tetrahydro-, hexahydrofuro-, hexahydropyrano-, and tetrahydrobenzofuroquinolines from *N,N*-dimethylaniline *N*-oxides and various electron-rich olefins in a tandem Polonovski–Povarov sequence is reported. Following activation of the N–O bond with Boc_2O , an exocyclic iminium ion is unveiled upon exposure to tin(IV) chloride. A formal inverse-electron-demand aza-Diels–Alder cyclization generates the tetrahydroquinoline core of 29 examples in up to 92% yield.

Graphical Abstract

The tetrahydroquinoline scaffold lies at the heart of countless natural products, biologically active materials, medicinal compounds, ligands, catalyst frameworks, and a myriad of other natural and synthetic materials.¹ Convenient access to this special structure is impactful in a variety of fields and disciplines, and a number of approaches to the tetrahydroquinoline structure have been described.²

Modern strategies toward the synthesis of highly substituted tetrahydroquinolines include the hydrogenation of quinolines,³ tandem hydroamination–hydrogenation of alkynyl anilines,⁴ and the classical Povarov reaction⁵ which combines an aniline, an aldehyde, and some alkene component in a formal inverse-electron-demand aza-Diels–Alder⁶ reaction (Scheme 1, eqs 1–3). Each of these approaches is limited by the nature of the reactions and

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Author Contributions

All experimental procedures were executed by T.B. X-ray crystallographic data was collected by G.Y.

The authors declare no competing financial interest.

Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/acs.orglett.8b02318.

Detailed experimental procedures, spectral data, and X-ray crystallographic data (PDF)

Accession Codes

CCDC 1847432–1847434 and 1847436–1847441 contain the supplementary crystallographic data for this paper. These data can be obtained free of charge via www.ccdc.cam.ac.uk/data_request/cif, or by emailing data_request@ccdc.cam.ac.uk, or by contacting The Cambridge Crystallographic Data Centre, 12 Union Road, Cambridge CB2 1EZ, UK; fax: +44 1223 336033.

the reagents required. For example, hydrogenation of quinolines can be achieved under the action of various transition metals⁷ or organocatalysts;⁸ however, the reaction conditions are often quite harsh, and the preparation of quinoline starting materials can be laborious (Scheme 1, eq 1). The tandem hydroamination–hydrogenation of alkynyl anilines is achieved under gold catalysis⁴ (and often relayed with an organocatalytic hydrogenation), which necessarily limits the scope of the available products to 2-substituted materials (Scheme 1, eq 2, substituent R₂). The Povarov reaction relies upon the protic or Lewis acid mediated condensation of an aniline and an aldehyde followed by capture of the resultant aza-diene with an alkene (Scheme 1, eq 3); thus, the Povarov reaction is often limited to electron-rich anilines and alkene components. There are successful asymmetric variants of each of these strategies that further impinge upon the scope of the available substrates (e.g., by requiring structural features such as Lewis basic groups that facilitate catalyst binding).^{3,4,7–9}

We report here a practical synthesis of the tetrahydroquinoline scaffold as part of our broader research program investigating the special reactivity of *N,N*-dialkylaniline *N*-oxides (Scheme 1, eq 4).¹⁰ The aniline *N*-oxides undergo *O*-acylation events with a host of electrophiles, and we have developed chemistry by which the weak N–O bond is excised in a Polonovski-type reaction to give an iminium ion.¹¹ Following that event, the imine is captured by an electron-rich alkene, and the resultant cation is then attacked by the pendant aniline. Thus, the reaction constitutes the stepwise formal [4 + 2] cycloaddition (aza-Diels–Alder reaction) of the Povarov reaction. In the case of aniline *N*-oxides, *O*-acylation requires an electrophile that will not undergo the N → C group transfer we have previously described;^{10,11} elimination can be achieved efficiently under the action of a Lewis acid to give a stable iminium ion (Scheme 2). The elimination pathway is favored over group transfer by employing di-*tert*-butyl dicarbonate (Boc₂O) as the acylating agent (which does not undergo N → C group transfer at a significant rate, Scheme 2b, **2** → **3b**), followed by treatment with tin(IV) chloride as a Lewis acid. The Lewis acid assisted elimination reaction to generate iminium ions proceeds cleanly on a variety of substituted *N,N*-dimethylaniline *N*-oxides (Scheme 2a, **2** → **3a**). We have successfully captured this electrophile with a variety of electron-rich vinyl ethers, thus restoring the electron-rich anilines (**3a** → **4**). The resultant cations are sufficiently stable to undergo intramolecular attack by the aniline (**4** → **5**) to give a variety of polycyclic scaffolds.

After the intramolecular attack by the aromatic ring, loss of a proton restores aromaticity (**5** → **6**). We extensively screened Lewis acidic reaction conditions (BF₃·OEt₂, BCl₃, FeCl₃, ZnBr₂, CuBr, CuI, InCl₃, TiCl₄, SnCl₄, and LiClO₄) and temperature profiles to support the cationic cyclization sequence and determined that tin(IV) chloride (0.25 equiv) in cold dichloromethane routinely afforded clean cyclized materials. As part of our study, we found these *N*-oxides to be far easier to handle and manipulate as complexes with *m*-chlorobenzoic acid. Following oxidation of the anilines with *m*-CPBA,¹⁰ the *N*-oxides cocrystallize with the acid byproduct to give robust crystalline solids that are bench stable and nonhygroscopic (see the Supporting Information, SI). Moreover, the *m*-chlorobenzoic acid does not interfere with the capture of iminium ions by exogenous nucleophiles, provided that excess activating agent is employed. For example, upon treatment of 4-acetoxy-*N,N*-dimethylaniline *N*-oxide

(**1a**) with di-*tert*-butyl dicarbonate (Boc₂O) and 4-(*N,N*-dimethyl-amino)pyridine (DMAP) at 0 °C, followed by cooling to –94 °C and addition of tin(IV) chloride and ethyl vinyl ether, the cationic cascade reaction afforded the substituted tetrahydroquinoline **7a** in 78% yield (Scheme 3).¹² The reaction is tolerant of both electron-rich and electron-poor substituents on the aromatic ring, giving 4-ethoxy-*N*-methyl-tetrahydroquinolines in 42–78% yields in 5–6 h at low temperature.

We then employed cyclic vinyl ethers in the Polonovski–Povarov reaction sequence and found these to be similarly reactive. By subjecting 2,3-dihydrofuran and 3,4-dihydro-2*H*-pyran to the same reaction conditions (Scheme 4), we obtained the polycyclic scaffolds **8** and **9** in 42–74% yields. The ring junctions of the furanoquinoline and pyranoquinoline products are formed exclusively as the *cis*-fused products as determined by ¹H NMR analysis, and we later confirmed these structural assignments by X-ray crystallography (**8a**, **8e**, **9a**, and **9b**, see SI).

Moreover, benzofurans are also sufficiently nucleophilic to participate in the cationic cyclization sequence, giving rise to tetracyclic products **10** in 41–87% yields (Scheme 5) and offering the opportunity to generate materials with two distinctly substituted aromatic residues (e.g., **10g** and **10h**). The regiochemistry of attack utilizing benzofuran nucleophiles is inverted relative to their dihydrofuran equivalents, as was expected.¹³ The products form exclusively as the *cis*-fused ring junction, and these assignments were also confirmed crystallo-graphically (**10a–c** and **10e**, see the SI).

In an effort to further expand the structural motifs accessible via the cationic cyclization sequence, we are exploring other less electron-rich nucleophiles, and we have identified allylic silanes¹⁴ as viable reaction participants (Scheme 6).¹⁵ The allylic silanes can engage the *N*-aryliminium ions, giving stabilized secondary carbocation intermediates, which undergo rapid ring closure. The allylic silanes offer the opportunity to generate diversely functionalized products **11**, which were formed in 31–92% yields under reaction conditions identical to those previously identified. Interestingly, the stereochemical outcome of these reactions with respect to the ring fusion appears to correlate with the olefin geometry of the starting materials, though our work with allylic silanes is ongoing. For example, when we employed *E*-aryl allylic silanes in the reaction sequence, we obtained the tetrahydroquinoline products **11c** and **11d** in 92% and 90% yields, respectively, as the *trans*-substituted products. We view these reactions as stepwise ring closure events, and we are further investigating the lifetime of the secondary cation intermediates under these reaction conditions.

We are working to further explore similar reactions with other allylic silanes and will report these and other new cycloaddition reactions in due course. Importantly, if the secondary cation intermediate is too stable, the reaction does not proceed beyond the initial capture of the iminium ion. For example, a silyl ketene acetal can capture the iminium ion in a Mannich-type¹⁶ reaction (Scheme 7a) directly analogous to our previous work in the synthesis of indolines (Scheme 7b).¹¹

We have described a practical and convenient synthetic protocol for the rapid assembly of various tetrahydroquinoline scaffolds via Lewis acid assisted tandem Polonovski–Povarov cyclizations. The products are obtained in up to 92% yield in 5–6 h at low temperature, and these reactions represent a novel use of the special reactivity of *N,N*-dimethylaniline *N*-oxides. Owing to the abundance of commercially available anilines and electron-rich alkenes, this methodology will allow for the facile construction of numerous natural products, pharmaceutically relevant molecules, and other useful polycyclic scaffolds.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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Scheme 1.
Representative Synthetic Approaches to Tetrahydroquinolines

Scheme 2.

(a) Mechanism of the Tandem Polonovski–Povarov Reactions with Aniline *N*-Oxides. (b) Group-Transfer Reactions Avoided by Use of Boc_2O

Scheme 3. Tandem Polonovski–Povarov Reactions Joining Aniline *N*-Oxides with Ethyl Vinyl Ether^a^aYields of isolated products; reactions were performed on a 1 mmol scale.

Scheme 4. Tandem Polonovski–Povarov Reactions Joining Aniline *N*-Oxides with Cyclic Vinyl Ethers^a^aYields of isolated products; reactions were performed on a 1 mmol scale.

Scheme 5. Tandem Polonovski–Povarov Reactions Joining Aniline *N*-Oxides with Benzofuran.^a^aYields of isolated products; reactions were performed on a 1 mmol scale.

Scheme 6. Reactions with Allylic Silanes as Nucleophiles^a^aYields of isolated products; reactions were performed on a 0.6 mmol scale.

Scheme 7.
 Mannich-Type Reactivity with Aniline *N*-Oxides