

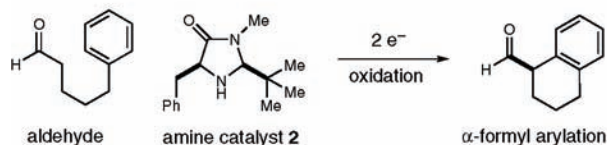
Enantioselective α -Arylation of Aldehydes via Organo-SOMO Catalysis. An Ortho-Selective Arylation Reaction Based on an Open-Shell Pathway

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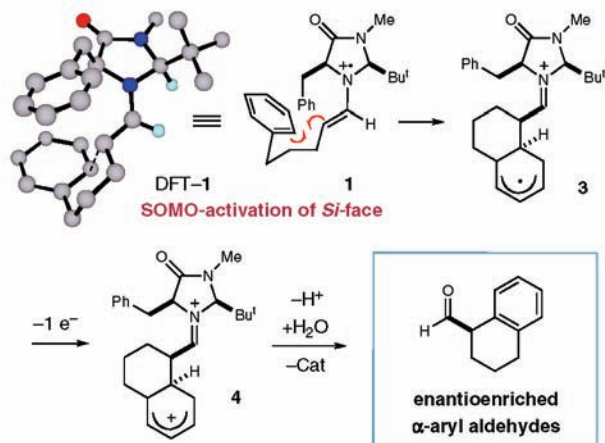
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Driven primarily by advances in transition metal chemistry, the catalytic union of enolates with aryl halides has become a mainstay transformation in asymmetric synthesis.¹ In particular, the pioneering work of Buchwald and Hartwig has provided a number of enantioselective enolate α -arylations that allow the catalytic formation of α -stereogenic carbonyls that are non-enolizable (e.g., quaternary carbon centers).^{2–4} Recently, our laboratory introduced a new mode of organocatalytic activation, termed SOMO catalysis, that is founded upon the mechanistic hypothesis that a 3π -radical cation species (e.g., **1**) can readily participate in a range of unique asymmetric bond constructions.⁵ As part of these studies, we documented the first direct and enantioselective α -allylic alkylation,^{5b} α -enolation,^{5c} and α -vinylation^{5d} of aldehydes, three protocols that were not previously known. Continuing this theme, we recently questioned whether the α -arylation of aldehydes might also be accomplished via SOMO catalysis to build and maintain enantioenriched benzylic stereocenters (eq 1). Herein, we describe the enantioselective intramolecular aldehyde α -arylation, a transformation that allows the highly efficient production of enolizable α -formyl- α -aryl substrates without postreaction epimerization or quaternary carbon formation.⁶ Moreover, this protocol is highly ortho-selective using 1,3-disubstituted aryl rings, a finding that is consistent with a radical based (open-shell) arylation mechanism.⁷

Enantioselective α -Arylation of Aldehydes via SOMO Catalysis (eq 1)



Mechanism of Organo-SOMO Catalyzed Aldehyde α -Arylation (eq 2)



Design Plan. In accord with our previous SOMO-catalysis studies,⁵ we reasoned that exposure of an aryl-tethered aldehyde to amine catalyst **2** in the presence of a suitable oxidant would render the radical cation **1** (DFT-1). Rapid and enantioselective addition of the pendent aryl ring system to the 3π -electron species would then provide a

Table 1. Development of Organo-SOMO Aldehyde α -Arylation

entry	oxidant	catalyst (Ar)	% yield ^a	% ee ^b
1	CAN	Ph, 2 (30 mol %)	63	84
2	[Fe(phen) ₃]·(PF ₆) ₃	Ph, 2 (30 mol %)	60	92
3 ^c	[Fe(phen) ₃]·(PF ₆) ₃	Ph, 2 (30 mol %)	85	92
4 ^c	[Fe(phen) ₃]·(PF ₆) ₃	Ph, 2 (20 mol %)	61	90
5 ^c	[Fe(phen) ₃]·(PF ₆) ₃	1-Naphth, 5 (20 mol %)	70	97
6 ^{c,d}	[Fe(phen) ₃]·(PF ₆) ₃	1-Naphth, 5 (20 mol %)	80	98

^a Isolated yields of the corresponding alcohol. ^b Enantiomeric excess determined by chiral HPLC. ^c 30 mol % pivalic acid. ^d 50 mol % H₂O.

bicyclic radical cation **3** that upon further oxidation would generate a dienyl cation **4** (eq 2). At this stage, we presumed that rearomatization via proton loss would be facile while hydrolysis of the iminium species would reconstitute the catalyst and deliver the desired formyl α -arylation product. Given the predilection of electron-rich aryl rings to undergo ortho-selective addition to radical cations,⁷ we presumed that high levels of regiochemical control should be possible if a traditional SOMO catalysis mechanism is operative. Moreover, we presumed that the activated radical DFT-1 would position the 3π -electron system away from the bulky *tert*-butyl group, while the benzyl group on the imidazolidinone framework effectively shields the *Re*-face leaving the *Si*-face open to enantioselective arylation.

Our intramolecular aldehyde α -arylation was first evaluated using 3-anisole pentanaldehyde, catalyst **2** or **5**, and a selection of oxidants (Table 1). Initial experiments with ceric ammonium nitrate (CAN) revealed that the proposed transformation was possible; however reaction efficiency and enantioselectivity levels were found to be moderate. Not satisfied with the general utility of these reaction conditions, we sought to find a superior catalyst/oxidant combination. As revealed in Table 1, use of [Fe(phen)₃]·(PF₆)₃ as a single electron oxidant allowed for a significant increase in enantiocontrol while application of a recently designed imidazolidinone catalyst **5** enabled optimal reaction efficiency and a further increase in stereocontrol (with 20 mol % catalyst). High-throughput experiments using a robotic platform (see Supporting Information) revealed that tightly controlled quantities of additives such as pivalic acid and water could also be employed to boost efficiency (entries 3–6). Given the notable selectivities achieved with both amines **2** and **5**, we selected both catalysts for further exploration.

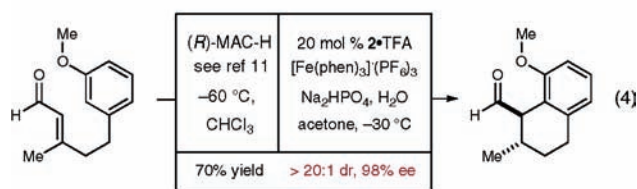
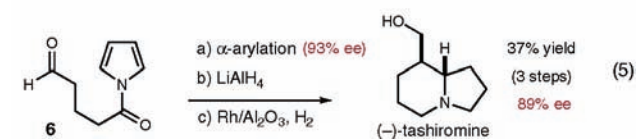
As highlighted in Table 2, a wide array of aryl and heteroaryl ring systems readily participate as SOMOphiles in this intramolecular α -aldehyde coupling. For example, anisole, naphthylene, indole, pyrrole, thiophene, and furan ring systems all function well⁹ in this new enantioselective ring forming reaction. Moreover, aldehydes that incorporate nitrogen, oxygen, or carbonyl containing tethers worked well in this protocol (entries 2, 4, 5, and 10, catalyst **5**: 93–96% ee).

Table 2. Scope of the Organo-SOMO Intramolecular α -Arylation

entry ^a	product	cat: yield, ^b ee ^c	entry ^a	product	cat: yield, ^b ee ^c
1		2: 61% yield 92% ee 5: 80% yield 98% ee	2		2: 84% yield 74% ee 5: 86% yield 95% ee
3		2: 83% yield 93% ee 5: 73% yield 96% ee	4		2: 72% yield 92% ee 5: 70% yield 95% ee
5		2: 70% yield 97% ee 5: 71% yield 96% ee	6		2: 86% yield 92% ee 5: 84% yield 96% ee
7		2: 89% yield 58% ee 5: 96% yield 90% ee	8		2: 84% yield 91% ee 5: 82% yield 96% ee
9		2: 86% yield 76% ee 5: 96% yield 94% ee	10		2: 72% yield 81% ee 5: 72% yield 93% ee

^a Conditions, entries 1–2: [Fe(phen)₃](PF₆)₃, NaHCO₃, HOPiv, –20 °C MeCN; entries 3–5: [Fe(phen)₃](PF₆)₃, Na₂HPO₄, –30 °C acetone; entries 6–10: CAN, NaHCO₃, Na₂O₂CCF₃, –30 °C acetone. ^b Isolated yields after NaBH₄ reduction. ^c Determined by chiral HPLC of corresponding alcohols.

As is clearly shown in Table 2, the imidazolidinone catalyst **5** is far superior to amine **2** with respect to enantioselectivity and cyclization efficiency. It is also important to note that these electron-rich aryl ring systems are completely tolerant to the mild oxidative conditions employed in this study. Furthermore, substrates that have the potential for aromatic substitution at two regiochemical positions (ortho or para) were in fact highly selective for ortho-functionalization (entries 1–5, ortho-substitution only).¹⁰ Such regioselectivity provides further evidence that a radical-mediated addition pathway is operative, as proposed in eq 2. It should be noted that a cationic Friedel–Crafts mechanism was recently proposed for a similar arylation reaction using our imidazolidinone catalyst **2**,⁸ a mechanistic interpretation based on the exclusive observation of para-selective products. However, the authors have subsequently issued a correction^{8b} that several of their adducts were in fact formed with ortho-selectivity (Table 2, entries 1 and 3). As such, given (i) our previous mechanistic studies into SOMO-activation, wherein an open-shell pathway was definitively shown using a radical-clock probe,^{5a} and (ii) the observation of only ortho-selective products using 1,3-disubstituted aryl rings, we presume that a radical cyclization is operative for the substrates described in this manuscript.^{9,10}

Sequential Organocatalytic Conjugate Reduction–SOMO Arylation**Enantioselective Synthesis of (–)-Tashiromine via Formyl α -Arylation**

We have been able to combine our novel SOMO α -arylation reaction into a two-step organocatalytic sequence that first involves a Hantzsch ester hydride (HEH) reduction¹¹ of an anisole tethered α,β -unsaturated aldehyde followed by the protocol described herein. As shown in eq 4, this sequence provides the resulting bicyclic ring system as a single (*anti*) diastereomer in 70% yield and 98% ee.

We have also been able to apply this new carbonyl α -arylation technology toward the total synthesis of (–)-tashiromine.¹² As outlined in eq 5, exposure of the pyrrole amide tethered aldehyde **6** to this new catalytic procedure provides the desired [6,5]-bicyclic ring system in 72% yield and in 93% ee. The resulting amide was then reduced in the presence of AlCl₃ with LiAlH₄ in 83% yield prior to exhaustive pyrrole hydrogenation using catalytic Rh/Al₂O₃ to provide (–)-tashiromine in 62% isolated yield.

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Supporting Information Available: Experimental procedures and spectral data are provided. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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