We present a highly regio- and chemoselective Csp3–H arylation of benzylamines mediated by synergy of single electron transfer (SET) and hydrogen atom transfer (HAT) catalysis. Under well preceded SET catalysis alone, the arylation reaction of N,N-dimethylbenzylamine proceeded via aminium radical cation formation and selectively targeted the N-methyl group. In contrast, addition of PhC(O)SH as a HAT catalyst precursor completely switched the regioselectivity to Csp3–H arylation at the N-benzylic position. Measurement of oxidation potentials indicated that the conjugate base of PhC(O)SH is oxidized in preference to the substrate amine. The discovery of the thiocarboxylate as a novel HAT catalyst allowed for the selective generation of the sulfur-centered radical, so that the N-benzylic selectivity was achieved by overriding the inherent N-methyl and/or N-methylene selectivity under SET catalysis conditions. While visible light-driven ω-C–H functionalization of amines has mostly been demonstrated with aniline derivatives and tetrahydroisoquinolines (THIQs), our method is applicable to a variety of primary, secondary and tertiary benzylamines for efficient N-benzylic C–H arylation. Functional group tolerance was high, and various 1,1-diarylmethylamines, including an α,ω,ω-trisubstituted amine, were obtained in good to excellent yield (up to 98%). Importantly, the reaction is applicable to late-stage functionalization of pharmaceuticals.

Introduction

The amino group is found in many kinds of molecules, including naturally occurring bioactive compounds, pharmaceutical drugs, agrochemicals, and functional materials. Therefore, direct and selective Csp3–H functionalization of amine compounds is extremely useful for rapid derivatization.1,2 In addition, the benzylamine unit is contained in various synthetic intermediates and is a core structure of many pharmaceuticals (Fig. 1A), as exemplified by donepezil (AChE inhibitor) and roxatidine (H2 blocker). To expedite structure–activity-relationship studies, regio- and chemoselective C–H functionalization of benzylamine derivatives is highly attractive. Moreover, 1,1-diarylmethylamine unit is a well-characterized pharmacophore (Fig. 1B).3 Considering the importance of such substructures in medicinal chemistry, mild and efficient methods to construct the 1,1-diarylmethylamine framework are of great interest, and benzylic Csp3–H arylation of benzylamines might be a straightforward and expedient approach.

Nucleophilic addition of an aryl nucleophile to an imine or an iminium ion is a commonly used strategy to construct the 1,1-diarylmethylamine core.4 On the other hand, recent work has explored direct benzylic Csp3–H arylation of amine derivatives (Scheme 1). Li and co-workers reported an oxidative cross-
A. Nucleophilic addition to iminium formed in situ

\[
\begin{align*}
R_2N^+ - &\rightarrow [Q] \rightarrow R_2N^+-Ar^1 \rightarrow R_2N^+-Ar^2
\end{align*}
\]

B. Benzylic deprotonation strategy

\[
R_2N^+ - \rightarrow \text{deprotonation} \rightarrow R_2N^- - \rightarrow \text{Ar}^2 - X (X = Cl, Br)
\]

C. Ru-catalyzed C–H activation

\[
\text{Scheme 1A: Previous benzylamine arylation strategies and this work.}
\]

D. Photoredox-catalyzed \( \alpha \)-C–H arylation of aniline derivatives

\[
\text{Scheme 1B: previous reports}
\]

E. Regioselectivity-controlled \( \text{Csp}^3 \)-H arylation of \( N \)-benzylamines

\[
\text{Scheme 1C: previous reports}
\]

results.

derivatives have been investigated actively as benchmark sub-

position, followed by transition-metal-catalyzed cross-coupling

employing a protocol involving deprotonation at the

addition to the oxidative approach, other researchers have

transformations under photoredox catalysis conditions.

In addition to the oxidative approach, other researchers have

employed a protocol involving deprotonation at the \( N \)-benzyl-

position, followed by transition-metal-catalyzed cross-coupling

reaction (Scheme 1B).\(^7\) Although a variety of aromatic groups

can be incorporated, these approaches normally require harsh

reaction conditions, such as strong base and high temperature.

Elegant methodologies using transition metal-catalyzed C–H

activation have also been disclosed, though in these cases, a
directing group is required for the interaction between the

transition metal and the benzylic C–H bond (Scheme 1C).\(^8\)

Photoinduced electron transfer has long been examined for

C–H functionalization adjacent to a nitrogen atom.\(^9,10\) In 2011,

MacMillan and co-workers reported a redox-neutral \( \alpha \)-C–H ary-

lation of various aniline derivatives using electron-deficient

arenes as arylation partners in the presence of a photoredox

catalyst (Scheme 1D).\(^11\) Following that work, we recently re-

ported a redox-neutral \( \alpha \)-C–H alkylation and cyanation of \( N \)-aryl-

THIQs using activated alkyl halides and tosylcyanide.\(^12\) While

visible light photoredox functionalization of amines has been

well investigated, the substrate scope is generally limited to

THIQ and aniline derivatives. On the other hand, there are some

reports on photo-mediated SET oxidation of tertiary alkyl

amines (e.g., \( \text{Et}_3\text{N} \) and \( \text{iPr}_2\text{NEt} \)) though they are used as sacrificial

reducing agents.\(^13,14\) As for benzylic C–H arylation, THIQ
derivatives have been investigated actively as benchmark sub-

strates.\(^15,16\) It should be noted that THIQs are special

 compounds in terms of both the oxidation potential and the

bond dissociation energy (vide infra), indicating that simple \( N \)-

alkyl benzyamines should be considered different from stan-
dard THIQs. Therefore, selective benzylic transformation of

simple alkylamines is still a challenging subject. To our

knowledge, there is no general and selective method available

for \( N \)-benzyl C–H arylation. Herein, we report a regio- and

chemoselective C–H functionalization of benzyamines via

photoredox-neutral SET and HAT synergistic catalysis (Scheme 1E).

The distinct features of the present reaction are listed below:

1. In addition to well-studied aniline-type compounds, various

unprotected \( N \)-alkyl benzyamines are available in this reaction.

2. Since PhC(=O)SH is easily deprotonated and converted to the

 corresponding sulfur-centered radical, single electron oxidation

of the amine substrates is blocked effectively. SET/HAT syner-
gistic catalysis enables excellent \( N \)-benzyl selectivity in prefer-
ence to inherent \( N \)-methyl and \( N \)-methylene selectivity observed

under SET catalysis conditions. (3) Due to the efficient genera-
tion of the HAT catalyst, as little as 1 mol% of PhC(=O)SH is

sufficient. In addition, an excess amount of starting amine is

not required in our reaction. (4) The chemoselectivity is high,

and the late-stage functionalization of pharmaceutical drugs

was demonstrated successfully.

Results and discussion

According to the reported reaction conditions of photoredox
catalysis via an aminium radical cation intermediate,\(^11\) we
began with the reaction of \( N, N \)-dimethylbenzylamine (1a)
with terephthalonitrile expecting that the most stable (benzylic
and \( 2^\# \)) \( \alpha \)-amino radical would be generated (Scheme 2). With
Ir(ppy)$_3$ (1 mol%) as a photoredox catalyst and K$_2$HPO$_4$ as base,
the reaction afforded two compounds, 2a and 3a, in excellent
mass balance (95%), though a long reaction time (12 h) was
required for completion. Contrary to our expectation, the major
product was not 2a but 3a, which is derived from the thermo-
dynamically less stable non-benzylic and \( 1^\# \) \( \alpha \)-amino radical.\(^17\)

While the origin of this regioselectivity is unclear at this
point, we think that the kinetic acidity in terms of stereo-
electronic and steric factors are crucial. Thus, deprotonation
of anilinium radical cations to give \( \alpha \)-amino radicals might be
accelerated by the overlap of the breaking Csp$^3$–H bond orbital
with the SOMO orbital on the aminium radical cation.\(^18,19\)

These

\[
\text{Scheme 2: Csp$^3$–H arylation of } N, N \text{-dimethylbenzylamine under photoredox catalysis.}
\]

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results suggested that well-precedented SET catalysis is not applicable and distinct reaction conditions should be devised for achieving the targeted N-benzylic transformation.

We considered that a HAT strategy might alter the regioselectivity (Scheme 3). A significant determinant of the selectivity of a HAT process is the C–H bond dissociation energy (BDE). Since BDE is directly associated with the stability of the radical products, a large difference between N-methyl C–H and N-benzylic C–H BDEs should be anticipated. According to the previous studies, we expected that sulfur-centered radicals would be HAT agents of choice for selective activation of the N-benzylic group. Importantly, however, in order to avoid direct single electron oxidation of N-benzylation 1\textsubscript{a} leading to the SET catalysis pathway (Scheme 3, right), the sulfur-centered radical precursor should have a lower anodic peak potential than that of N-benzylation 1\textsubscript{a}. For this purpose, we envisaged that the anion of a sulfur atom would undergo more facile oxidation than a typical thiol such as cysteine ($E_{1/2}$ = +0.85 V vs. SCE for cysteine in CH\textsubscript{3}CN, $p_{K_a}$ = 9.35 for cysteine methyl ester) and that a more acidic sulfur compound would be more favorable. Therefore, we selected thiocarboxylic acid 4, whose conjugate base could be generated with a weak base ($p_{K_a}$ = 3.2 for thioacetic acid). The negatively charged, electron-rich thiocarboxylate species might undergo SET oxidation much faster than the substrate amine, thereby affording the sulfur-centered radical 6 preferentially (Scheme 3, left). This scenario would lead to regioselective C–H activation at the N-benzylic position by overriding the natural N-methyl selectivity observed in Scheme 2.

To validate our working hypothesis, we measured the oxidation potentials of potassium thiobenzoate (PhC(O)SK) and three N-benzylnitramines by cyclic voltammetry (CV) in DMA (Fig. 2; see ESI† for potentials vs. SCE and further CV studies). Gratifyingly, PhC(O)SK was found to have a less positive anodic peak potential ($E_{\text{pa}}$ (6/5) = +0.80 V vs. Ag/AgCl in DMA) than all the amine compounds examined, even when compared to the aniline-type N-benzyl tetrahydroquinoline (THQ) ($E_{\text{pa}}$ = +0.97 V vs. Ag/AgCl in DMA). Therefore, thermodynamics predicted that selective oxidation of the thiobenzoate anion would occur even in the presence of the reducing amine substrates. Furthermore, CV revealed that oxidation of PhC(O)SK by [Ir\textsuperscript{IV}(ppy)\textsubscript{3}]$^+$ ($E_{1/2}$ (Ir\textsuperscript{IV}/Ir\textsuperscript{III}) = +0.96 V vs. Ag/AgCl in DMA) was exergonic, while oxidation of the amines was always endergonic. According to these data, a faster reaction profile is expected compared to the reaction shown in Scheme 2.

With the promising CV results in mind, we revisited benzylic C–H arylation of N,N-dimethylbenzylamine (1\textsubscript{a}) in the presence of PhC(O)SH (4) and other HAT catalyst precursors (Table 1). The arylation proceeded in 91% yield at the benzylic position with excellent regioselectivity ([r.r.] >> 20 : 1, entry 2), which is consistent with the idea of a mechanistic switchover from SET catalysis to SET/HAT synergistic catalysis. It should be noted that the reaction reached completion within 1 h (entry 3), compared to 12 h required when PhC(O)SH was absent (entry 1). This rate-acceleration might be due to exergonic SET oxidation of the thiocarboxylate and a favorable difference in BDE between PhC(O)S–H (87.4 kcal mol$^{-1}$) and the targeted N-benzylic C–H bond (84.9 kcal mol$^{-1}$). Precatalysts 10 and 11 gave 2\textsubscript{a} with excellent N-benzylic selectivity. However, the yields were much lower even in the presence of 20 mol% of HAT catalysts (entries 4, 5). We assume that these less acidic thiols are not deprotonated under the reaction conditions, and thus SET oxidation might not be efficient compared to the case of PhC(O)SK ($E_{1/2}$ = +0.85 V vs. SCE for cysteine in CH\textsubscript{3}CN, $E_{\text{pa}}$ (6/5) = +0.65 V vs. SCE in CH\textsubscript{3}CN; see ESI†). Considering the $p_{K_a}$ values of the HAT precatalysts, 12 was also examined as a more acidic thiol (entry 6). However, almost no reaction was observed, probably because the generated thiyl radical would be less reactive in the subsequent HAT process (S–H BDE of TolSH = 77–83 kcal mol$^{-1}$) (entry 4). Recently, quinuclidine and DABCO amino radical cations have been reported to perform HAT with alkylamines at the $\alpha$ position to the nitrogen. Given its accessible oxidation potential ($E_{\text{pa}}$ = +0.69 V vs. SCE in

Scheme 3 Working hypothesis for regioselective arylation at the N-benzylic position.
CH$_3$(CN)$_2$DABCO would undergo SET oxidation selectively over 1a. However, when PhC(O)SK was replaced with DABCO, the reaction did not proceed (entry 7). The reason is not clear at present, but back electron transfer processes such as quenching of the DABCO radical cation by arene radical anion might be sufficient to interfere with the HAT process.$^{26}$ When PhC(O)SK was replaced with quinuclidine, the product was formed in only very low yield with low regioselectivity (entry 8), probably because the strong reactivity of quinuclidine radical formed in only very low yield with low regioselectivity (entry 8), the product was less reactive than DABCO. When PhC(O)SH was replaced with DABCO, the reaction proceeded with excellent yield (entry 12), in contrast to most of the preceding examples in which a large excess amount of amine substrate (at least three equivalents of starting amines) was generally needed to achieve good yields. A control experiment confirmed that the reaction does not occur in the dark (see ESI†).

With the optimal reaction conditions identified (Table 1, entry 11), we turned our attention to the substrate scope of the SET/HAT synergistic catalysis for Csp$_3$–H arylation of N-benzylamines (Table 2). N,N-Dimethylbenzylamine derivatives with varying electronic properties were well tolerated, and the desired products (2b–f) were obtained in good yields. Electron-rich derivatives were the highest-yielding, which is consistent with the polarity effects.$^{24}$ However, para-ester-substituent retarded the reaction and the corresponding arylated product was not obtained, suggesting that strongly electron-withdrawing groups are not suitable for this reaction. Compared to the para-methyl-substituted derivative (2f), the ortho-methyl-substituted derivative gave 2g in a lower yield (75%), probably due to the steric interaction. Similarly, the reaction of the starting benzylamine was much faster than that of the dibenzylamine, so that over-arylation of the products was not observed. Nevertheless, we found that treating sterically congested 1h under the reaction conditions successfully afforded α,α,α-trisubstituted amine 2h in good yield (74%). Different substituted groups on the nitrogen atom did not prohibit the reaction, since ethyl groups (2i), aliphatic cyclic substituents (2j, 2k), and heteroatom-bearing cyclic substituents (2l, 2m) performed equally well under the reaction conditions. N-Methyl THIQ provided 2n in 94% yield as an expected regioisomer.$^{25}$ Notably, secondary and primary amines having no protective group were also applicable, affording 2o and 2p in 98% and 98% yield, respectively. These are relatively difficult substrates for SET catalysis due to their more positive anodic peak potentials, and so have been less well studied in the literature. Glycine derivative 1q underwent selective N-benzyl functionalization to afford 2q in 97% yield. Moreover, heterocyclic aromatic derivatives were tolerated, providing the corresponding arylated products 2r–2t. Aside from terephthalonitrile, 4-cyanopyridine derivatives and 1-cyanoisoquinoline reacted in a regioselective manner, providing 2u–2v in good to excellent yields. When cyanobenzenes bearing different para-electron-withdrawing groups were examined, the reaction proceeded with excellent yields.

### Table 1 Screening of conditions for benzylic Csp$_3$–H arylation of N,N-dimethylbenzylamine

<table>
<thead>
<tr>
<th>Entry</th>
<th>HAT cat.</th>
<th>P.C.</th>
<th>Time (h)</th>
<th>Yield$^b$ (%)</th>
<th>2a : 3a$^b$</th>
</tr>
</thead>
<tbody>
<tr>
<td>1$^e$</td>
<td>—</td>
<td>15</td>
<td>12</td>
<td>95</td>
<td>1 : 8.5</td>
</tr>
<tr>
<td>2</td>
<td>4</td>
<td>15</td>
<td>12</td>
<td>91</td>
<td>&gt;20 : 1</td>
</tr>
<tr>
<td>3</td>
<td>4</td>
<td>15</td>
<td>1</td>
<td>93</td>
<td>&gt;20 : 1</td>
</tr>
<tr>
<td>4</td>
<td>10</td>
<td>15</td>
<td>1</td>
<td>23</td>
<td>&gt;20 : 1</td>
</tr>
<tr>
<td>5</td>
<td>11</td>
<td>15</td>
<td>1</td>
<td>43</td>
<td>&gt;20 : 1</td>
</tr>
<tr>
<td>6</td>
<td>12</td>
<td>15</td>
<td>1</td>
<td>Trace</td>
<td>—</td>
</tr>
<tr>
<td>7</td>
<td>13</td>
<td>15</td>
<td>1</td>
<td>Trace</td>
<td>—</td>
</tr>
<tr>
<td>8</td>
<td>14</td>
<td>15</td>
<td>1</td>
<td>12</td>
<td>1 : 2</td>
</tr>
<tr>
<td>9</td>
<td>14</td>
<td>16</td>
<td>1</td>
<td>42</td>
<td>&gt;20 : 1</td>
</tr>
<tr>
<td>10</td>
<td>14</td>
<td>17</td>
<td>1</td>
<td>No reaction</td>
<td>—</td>
</tr>
<tr>
<td>11$^{e,d}$</td>
<td>4</td>
<td>15</td>
<td>2</td>
<td>90 (87)$^f$</td>
<td>&gt;20 : 1</td>
</tr>
<tr>
<td>12$^{e,d}$</td>
<td>4</td>
<td>15</td>
<td>2</td>
<td>72</td>
<td>&gt;20 : 1</td>
</tr>
</tbody>
</table>

$^a$The reactions were run on 0.2 mmol scale. $^b$Yield and regioisomeric ratio were determined by $^1$H NMR analysis using 1,1,2,2-tetrachloroethane as an internal standard. $^c$Data from Scheme 3. $^d$1 mol% of PhC(O)SH and 0.5 mol% of Ir(ppy)$_3$ were used on a 1 mmol scale. $^e$2 equiv. of N,N-dimethylbenzylamine was used. $^f$Isolated yield. $^g$1 equiv. of N,N-dimethylbenzylamine was used. P.C.: photocatalyst.
regioselectivity, though in less satisfactory yields (2x). For a comparative example, the reaction of 1j without PhC(O)SH was carried out, and a mixture of the C1-arylated 2j (36%) and the C2-arylated 3j (45%) was obtained after 12 h (Scheme 4). These results clearly indicate that the excellent N-benzyl selectivity observed in our reaction is attributed to SET/HAT synergistic catalysis that can outcompete the inherent N-methyl and/or N-methylene selectivity observed under SET catalysis conditions.

An interesting phenomenon was observed when comparing cyclic N-benzylaniline-type substrates. Treatment of N-benzyl indoline (1aa) under our reaction conditions resulted mainly in N-benzyl arylation. Nonetheless, under photocatalyst-mediated SET catalysis alone, the regioselectivity was inverted completely from the N-benzyl position to the cyclic N-methylene position (90% yield, C1/C2 = 1 : 6.7). In contrast, N-benzyltetrahydroquinoline (1ab) reacted under our conditions to give the cyclic N-methylene-arylated product 2ab with high regioselectivity (C1/C2 = 1 : 17). We postulate that the increase in steric repulsion around the N-benzyl position in moving from the 5- to the 6-membered ring system may be responsible. However, on the other hand, N-benzyltetrahydrobenzoxepine (1ac) gave a 1 : 1 mixture of the C1/C2 regioisomers. The origin of the regioselectivity remains unclear at this point, but we assume that the conformational flexibility within the nitrogen atom-containing ring is important to achieve orbital overlap of the n-orbital on the nitrogen atom with the σ* orbital of the breaking C–H bond, which should have an impact on the BDE.

Next, we evaluated the chemoselectivity of our reaction in more detail. For substrate 1ad, exclusive N-benzyl functionalization occurred in the presence of the benzyl alcohol (Scheme 5A), reflecting the difference in C–H BDEs between the benzyl alcohol side (C–H BDE of benzyl alcohol = 87.5 kcal mol\(^{-1}\)) and the N-benzyl group (C–H BDE of N,N-dimethylbenzylamine = 84.9 kcal mol\(^{-1}\)). An intramolecular competitive experiment with 1ae targeted the N-benzylamine side to produce 2ae in 71% yield (Scheme 5B). We next examined the reaction of a more challenging substrate (Scheme 5C). The C–H arylation reaction of 1af occurred only at the N-benzyl position, even though the BDEs of both N-benzyl C–H bond and N-allylic C–H bond are reported to be close (ca. 85 kcal mol\(^{-1}\) and 83 kcal mol\(^{-1}\), respectively). The yield was only moderate due to competitive reactions at the C–C double bond, but no arylation product at the N-allylic position was detected in this reaction. Moreover, we conducted an intermolecular competition experiment using 1b and 1c (1 : 1). In this case, the reaction mainly proceeded with 1b bearing an electron-donating group, which is indicative of a polarity matching effect (Scheme 5D).

To confirm the synthetic utility of our SET/HAT synergistic catalysis, we evaluated its efficiency in late-stage functionalization. When treated in a 1 : 1 ratio, donepezil and 7 underwent regio- and chemoselective coupling reaction smoothly to give 1,1-diarylmethane 2ag in good yield (74%). Further, even with more complex clopidogrel, C–H arylation at the endocyclic

---

**Table 2** Substrate scope for benzylic Csp\(^3\)-H arylation of N-benzylamines

<table>
<thead>
<tr>
<th>Substrate</th>
<th>Yield</th>
<th>Regioselectivity</th>
</tr>
</thead>
<tbody>
<tr>
<td>1aa</td>
<td>90%</td>
<td>C1/C2 = 1 : 6.7</td>
</tr>
<tr>
<td>1ab</td>
<td>1 : 17</td>
<td>C1/C2 = 1 : 17</td>
</tr>
<tr>
<td>1ac</td>
<td>1 : 1</td>
<td>C1/C2 = 1 : 1</td>
</tr>
</tbody>
</table>

*All reactions were conducted on 1 mmol scale. Terephthalonitrile was used as an arylating reagent (Ar = 4-NCCC\(_6\)H\(_4\)). The reaction was carried out for 6 h. Yield and regioisomeric ratio were determined by \(^1\)H NMR analysis using 1,1,2,2-tetrachloroethane as an internal standard.*

---

**Scheme 4** Arylation of 1j under SET conditions.
N-benzylic position proceeded regioselectively, furnishing 2ah in 96% yield. These examples clearly demonstrate that the present reaction is applicable to late-stage derivatization of structurally complex bioactive compounds, and thus should be useful to facilitate structure–activity-relationship studies. All reactions were conducted on 1 mmol scale and isolated yields are shown. Chemo- and regioselectivity were determined by 1H NMR analysis.

Finally, on the basis of the observed selectivity and CV measurements, we propose the catalytic cycle shown in Scheme 6. fac-Tris(2-phenylpyridinato)iridium(III) [Ir(ppy)3] (15) is excited by photo-irradiation (blue LED, 425 nm) and [IrIII(ppy)3]* (18) is generated. This reductant [Ir(ppy)3]⁺ (18) (E1/2 (IrIV/IrIII) = −1.73 V vs. SCE in CH3CN)33 undergoes SET to electron-deficient arene 20 to afford radical anion intermediate 21 along with [IrIII(ppy)3]⁻ (19). Subsequently, [IrIV(ppy)3]+ (19) (E1/2 (IrIV/IrIII) = +0.77 V vs. SCE in CH3CN)33 performs SET oxidation of the electron-rich PhC(O)SK (Eox (6/5) = +0.65 V vs. SCE in CH3CN, see ESI†), which outcompetes SET oxidation of the anion substrate leading to the different reaction pathway. The generated sulfur-centered radical 6 regioselectively undergoes HAT depending on C–H BDEs. As seen in the previous reports, the resulting N-benzyl radical 22 reacts with the separately formed radical anion intermediate 21 to yield the desired 1,1-diarylmethylamine 2.34

**Conclusions**

In conclusion, we have developed a highly regio- and chemoselective Csp³–H arylation of a variety of benzylamine derivatives by employing SET/HAT synergistic catalysis. Under SET catalysis alone, Csp³–H arylation of benzylamines proceeded at the N-methyl and/or cyclic N-methylene positions through an aminium radical cation intermediate. In contrast, synergistic SET and HAT catalysis inverts the regioselectivity. The reaction was completed within 2 h in the presence of as little as 0.5 mol% of the Ir complex and 1 mol% of PhC(O)SH as a HAT catalyst, and various 1,1-diarylmethylamines were obtained in good to excellent yields (56–98%). Importantly, high yields were achieved even when a stoichiometric amount of the benzylamine substrate was employed. From a mechanistic viewpoint, the use of PhC(O)SH as a HAT catalyst precursor is the key to success: SET oxidation of the 3'-amine substrates was suppressed effectively due to the favorable oxidation of PhC(O)SK, and the resulting sulfur radical abstracts C–H bonds selectively to give N-benzyl radicals. The excellent regio- and chemoselectivity enables the late-stage N-benzyl arylation of pharmaceuticals with reactive functional groups. Further studies including N-methyl selective functionalization are underway in our laboratory.
Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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Notes and references


19 The detail of the N-methyl selective arylation under SET catalysis conditions will be reported separately in due course.