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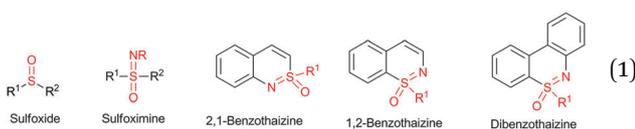
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# Ruthenium- and palladium-catalyzed consecutive coupling and cyclization of aromatic sulfoximines with phenylboronic acids: an efficient route to dibenzothiazines†

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**A ruthenium-catalyzed *ortho* arylation of aromatic sulfoximines with aromatic boronic acids followed by intramolecular cyclization in the presence of a palladium catalyst, providing dibenzothiazine derivatives in two consecutive steps, is described.**

Sulfoximine is a pivotal structural motif which is present in various biologically active molecules, pharmaceuticals and agrochemicals (eqn (1)).<sup>1</sup> The sulfoximine derivatives are also successfully used as chiral auxiliaries and ligands in asymmetric synthesis of various chiral organic molecules.<sup>2</sup> Several methods are available in the literature to synthesize linear sulfoximine derivatives.<sup>3</sup> But, the synthesis of cyclic sulfoximines is limited in the literature.<sup>4</sup> Recently, the synthesis of cyclic sulfoximines has gained much attention in organic synthesis due to their usefulness as scaffolds in drug development and as chiral ligands in enantioselective reactions.<sup>5</sup> Meanwhile, sulfoximine derivatives also serve as key synthetic intermediates in various organic transformations.<sup>5–7</sup>

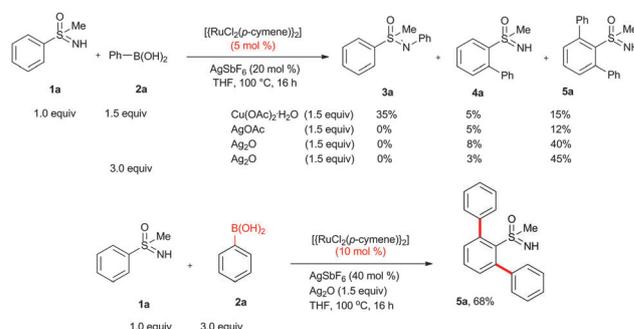


Harmata's group reported the synthesis of bicyclic sulfoximine derivatives such as 1,2-benzothiazine and 2,1-benzothiazine by palladium-catalyzed cyclization of 2-bromo benzaldehydes or 2-alkenylated aromatic bromides with sulfoximines, AlCl<sub>3</sub>-mediated cyclization of sulfonimidoyl chlorides with alkynes or alkenes and electrophilic cyclization of 2-bromophenyl substituted sulfoximines with terminal alkynes in the presence of palladium and copper catalysts.<sup>6</sup> Very recently, Bolm's group reported the synthesis of bicyclic 1,2-benzothiazine derivatives *via* rhodium-catalyzed oxidative cyclization<sup>7a</sup> of phenyl sulfoximines with

alkynes *via* a chelation-assisted C–H bond activation reaction.<sup>8–10</sup> Subsequently, sulfoximine directed *ortho* alkenylation of phenyl sulfoximines with alkenes in the presence of a metal catalyst was also disclosed.<sup>7b–e</sup> In all these reports, sulfoximine containing bicyclic benzothiazine derivatives were synthesized efficiently.

Herein, we report the synthesis of tricyclic dibenzothiazines by a ruthenium-catalyzed *ortho* arylation of phenyl sulfoximines with aromatic boronic acids followed by intramolecular cyclization in the presence of a palladium catalyst in two consecutive steps. The present reaction was compatible with various sensitive and useful functional group substituted phenyl sulfoximines and aromatic boronic acids. An enantioselective version of *ortho* arylation of phenyl sulfoximines with phenylboronic acids followed by cyclization, and this transformation leads to chiral dibenzothiazines with an excellent ee ratio of 99%.

Initially, the *ortho* arylation of phenyl sulfoximine (**1a**) with phenylboronic acid (**2a**) (1.5 equiv.) in the presence of [[RuCl<sub>2</sub>(*p*-cymene)]<sub>2</sub>] (5 mol%), AgSbF<sub>6</sub> (20 mol%) and Cu(OAc)<sub>2</sub>·H<sub>2</sub>O (1.5 equiv.) in THF at 100 °C for 16 h was investigated (Scheme 1). In the reaction, *N*-arylated phenyl sulfoximine **3a** in 35% yield, mono *ortho* arylated phenyl sulfoximine **4a** in 5% yield and bis *ortho* arylated phenyl sulfoximine **5a** in 15% yield were observed, respectively (Scheme 1). It is known that the free N–H group of **1a** is acidic in nature and smoothly undergoes *N*-arylation with aromatic electrophiles or organometallic reagents



Scheme 1 *ortho* Arylation of sulfoximine **1a** with **2a**.

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providing *N*-arylated sulfoximines **3** in the presence of metal catalysts.<sup>11</sup> To successfully carry out the *ortho* arylation reaction, the suppression of product **3** is highly important. Next, the reaction was tested with other oxidants and acetate sources such as AgOAc, NaOAc, K<sub>2</sub>CO<sub>3</sub>, CsOAc and Ag<sub>2</sub>O. Among them, silver salts such as AgOAc and Ag<sub>2</sub>O were active for the reaction and no *N*-arylated product **3a** was observed. In AgOAc, product **4a** in 5% and **5a** in 12% yields were observed, respectively. In Ag<sub>2</sub>O, product **4a** in 8% and **5a** in 40% yields were observed, respectively. Other acetate sources were not active for the reaction. Next, the coupling reaction was tested with an excess amount of phenylboronic acid **2a** (3.0 equiv.). In the reaction also, a mixture of products **4a** and **5a** were observed in 3% and 45% yields, respectively. To increase the yield of **5a**, the coupling reaction was performed in the presence of 10 mol% of catalyst and 40 mol% of AgSbF<sub>6</sub>. Interestingly, in the reaction, only bis *ortho* arylated product **5a** was observed in 68% isolated yield and no mono arylated product **4a** was observed. For the detailed optimization studies, please see ESI.†

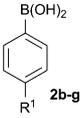
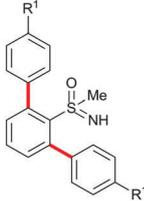
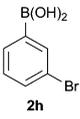
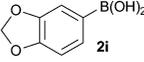
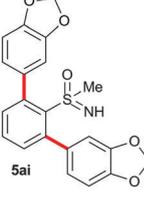
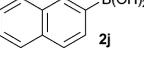
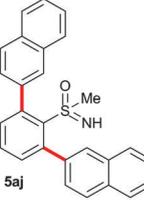
In addition to phenylboronic acid (**2a**), a wide range of aromatic boronic acids **2b–j** also readily participates in the reaction with **1a**. Table 1 summarizes the results of these reactions. Treatment of 4-phenyl substituted phenylboronic acid (**2b**) with **1a** provided *ortho* bis arylated product **5ab** in 72% yield (entry 1). Electron rich 4-methoxyphenyl boronic acid (**2c**) reacts smoothly with **1a**, yielding the corresponding product **5ac** in 66% yield (entry 2). Aromatic boronic acids having halogen groups I, Br, Cl and F **2d–g** also undergo an *ortho* arylation reaction with **1a** efficiently, giving products **5ad–ag** in 65%, 62%, 64% and 60% yields, respectively (entries 3–6). However, 3-bromo phenylboronic acid (**2h**) yielded product **5ah** only in 19% yield (entry 7). Benzo[*d*][1,3]dioxol-5-ylboronic acid (**2i**) and 2-naphthylboronic acid (**2j**) also efficiently participated in the reaction, affording coupling products **5ai** and **5aj** in 61% and 64% yields, respectively (entries 8 and 9).

The arylation reaction was examined with substituted aromatic sulfoximines **1b–g** (Scheme 2). Electron-rich, halogen and electron-deficient group substituted sulfoximines were compatible for the reaction. Methyl, Br and NO<sub>2</sub> substituted sulfoximines **1b–d** reacted with **2a**, providing products **5ba–da** in 65%, 63% and 54% yields, respectively. Similarly, Cl and F substituted aromatic sulfoximines **1e–f** reacted with **2c**, providing products **5ec–fc** in 60% and 63% yields, respectively. Likewise, (ethylsulfonylimidoyl)-benzene (**1g**) afforded **5ga** in 71% yield.

Apart from bis arylation, mono arylation of phenyl sulfoximines was also disclosed (Scheme 3). Treatment of 2-methyl phenylsulfoximine (**1h**) with **2a** or **2f** gave mono arylated sulfoximine derivatives **5ha** and **5he** in 70% and 63% yields, respectively. However, 3-methyl phenylsulfoximine (**1i**) afforded regioisomeric mono arylated products **5ia** and **5ia'** in 62% and 7% yields, respectively.

Next, we tried to couple the N–H bond of sulfoximine with one of the C–H bond of phenyl groups of compound **5** *via* chelation-assisted remote C–H activation in order to prepare tricyclic dibenzothiazine derivatives. A Pd(OAc)<sub>2</sub> catalyst along with an oxidant is the suitable condition for this type of cyclization.<sup>7c,12</sup> The intramolecular cyclization of compound **5aa**

**Table 1** Ruthenium-catalyzed *ortho* arylation of **1a** with aromatic boronic acids **2b–j**<sup>a</sup>

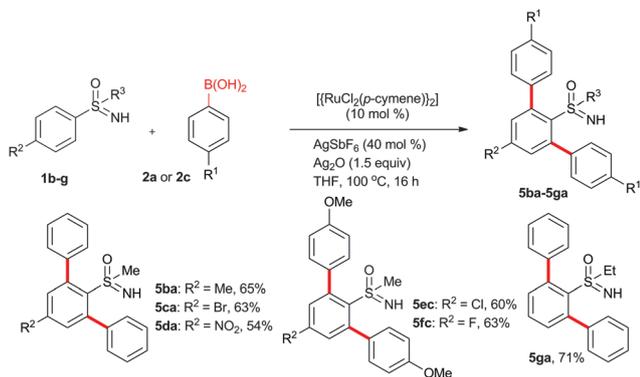
| Entry | <b>2b–j</b>  | Product <b>5</b>  | Yield <sup>b</sup> (%) |
|-------|--|---|------------------------|
| 1     |    |    | 72                     |
| 2     | <b>2c</b> : R <sup>1</sup> = OMe   | <b>5ac</b> : R <sup>1</sup> = OMe   | 66                     |
| 3     | <b>2d</b> : R <sup>1</sup> = I   | <b>5ad</b> : R <sup>1</sup> = I   | 65                     |
| 4     | <b>2e</b> : R <sup>1</sup> = Br  | <b>5ae</b> : R <sup>1</sup> = Br  | 62                     |
| 5     | <b>2f</b> : R <sup>1</sup> = Cl  | <b>5af</b> : R <sup>1</sup> = Cl  | 64                     |
| 6     | <b>2g</b> : R <sup>1</sup> = F   | <b>5ag</b> : R <sup>1</sup> = F   | 60                     |
| 7     |    |    | 19                     |
| 8     |    |   | 61                     |
| 9     |  |  | 64                     |

<sup>a</sup> All reactions were carried out using **1a** (100 mg), aromatic boronic acids (**2b–j**) (3.0 equiv.), [RuCl<sub>2</sub>(*p*-cymene)]<sub>2</sub> (10 mol%), AgSbF<sub>6</sub> (40 mol%), Ag<sub>2</sub>O (1.5 equiv.) in THF (3.0 mL) at 100 °C for 16 h. <sup>b</sup> Isolated yield.

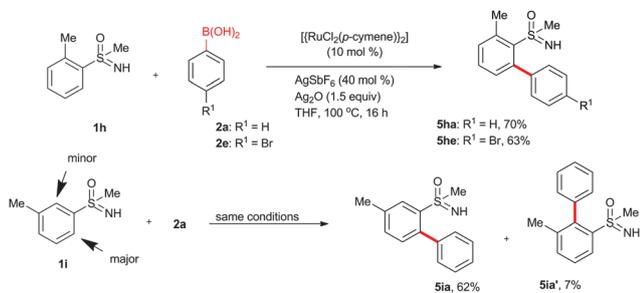
proceeded smoothly in the presence of Pd(OAc)<sub>2</sub> (10 mol%) and PhI(OAc)<sub>2</sub> (2.0 equiv.) in toluene at 120 °C for 10 h giving a tricyclic dibenzothiazine derivative **6a** in 76% yield (Table 2, entry 1). The cyclization reaction also proceeded in the presence of PhI(OAc)<sub>2</sub> without a palladium catalyst. However, product **6a** was observed in a less amount of 25% yield. Under similar reaction conditions, products **5ab**, **5ac**, **5ad**, **5ae**, **5af** and **5ag** also efficiently participated in the reaction, providing cyclization products **6b–g** in good to excellent yields (entries 2–7). Similarly, products **5ba**, **5ca**, **5da**, **5ga** and **5ha** afforded dibenzothiazines **6h–l** in 80%, 84%, 79%, 83% and 41% yields, respectively (entries 8–12). The structure of compound **6f** was further confirmed by single crystal X-ray analysis (see ESI†).

This result prompted us to explore the possibility of synthesis of chiral tricyclic dibenzothiazines by using chiral phenyl





Scheme 2 Scope of aromatic sulfoximines.

Scheme 3 Mono arylation of aromatic sulfoximines **1h-i**.

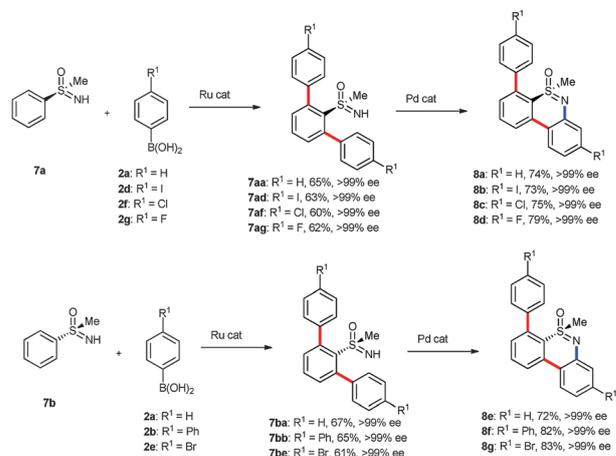
sulfoximines **7a-b** (Scheme 4). Treatment of chiral (*R*)-(-)-*S*-methyl-*S*-phenylsulfoximine (**7a**) with substituted phenyl boronic acids **2a**, **2d**, **2f** and **2g** in the presence of  $[(\text{RuCl}_2(p\text{-cymene}))_2]$ , AgSbF<sub>6</sub> and Ag<sub>2</sub>O in THF at 100 °C for 16 h gave chiral *ortho* arylated phenyl sulfoximines **7aa-ag** in 65%, 63%, 60% and 62% yields, respectively (Scheme 4). Interestingly, the enantiomeric excess (ee) of products **7aa-ag** did not drop and in all cases >99% ee ratios were observed. Later, compounds **7aa-ag** were cyclized into chiral dibenzothiazines **8a-d** in excellent 74%, 73%, 75% and 79% yields, respectively, in the presence of a palladium catalyst. In all these reactions, >99% ee ratios were observed. Furthermore, (*S*)-(-)-*S*-methyl-*S*-phenylsulfoximine (**7b**) underwent *ortho* arylation with aromatic boronic acids **2a**, **2b** and **2e** in the presence of a ruthenium catalyst, providing chiral *ortho* arylated phenyl sulfoximines **7ba-be** in 67%, 65% and 61% yields, respectively, with >99% ee ratios. Furthermore, **7ba-be** were converted into chiral dibenzothiazines **8e-g** in the presence of Pd(OAc)<sub>2</sub> in 72%, 82% and 83% yields, respectively.

A possible reaction mechanism is proposed to account for the present reaction in Scheme 5. Two different catalytic reactions were involved in the reaction. In the first catalytic cycle, AgSbF<sub>6</sub> likely removes all Cl<sup>-</sup> ligands from the ruthenium complex providing a cationic ruthenium complex **9**.<sup>13</sup> Coordination of the nitrogen atom of sulfoximine **1** into catalyst **9** followed by *ortho*-metalation provides a ruthenacycle intermediate **10**. Transmetalation of phenyl boronic acid **2** into intermediate **10** in the presence of Ag<sub>2</sub>O affords intermediate **11**. Subsequent reductive elimination of intermediate **11** in the presence of Ag<sup>+</sup> source provides product **5** and regenerates the

Table 2 Synthesis of dibenzothiazines<sup>a</sup>

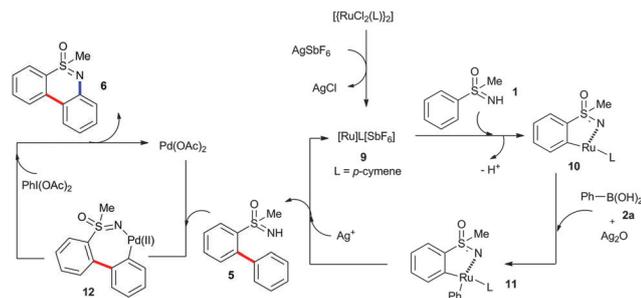
| Entry | 5 | Product 6 | Yield <sup>b</sup> (%) |
|-------|---|-----------|------------------------|
| 1     |   |           | 76                     |
| 2     |   |           | 85                     |
| 3     |   |           | 65                     |
| 4     |   |           | 77                     |
| 5     |   |           | 85                     |
| 6     |   |           | 79                     |
| 7     |   |           | 81                     |
| 8     |   |           | 80                     |
| 9     |   |           | 84                     |
| 10    |   |           | 79                     |
| 11    |   |           | 83                     |
| 12    |   |           | 41                     |

<sup>a</sup> All reactions were carried out using **5** (100 mg), Pd(OAc)<sub>2</sub> (10 mol%) and PhI(OAc)<sub>2</sub> (2.0 equiv.) in toluene at 120 °C for 10 h. <sup>b</sup> Isolated yield.



Scheme 4 Synthesis of chiral dibenzothiazines.





Scheme 5 Proposed mechanism.

active ruthenium species **9** for the next catalytic cycle. Another *ortho* arylation also takes place in a similar fashion. In the second catalytic cycle, compound **5** reacts with Pd(OAc)<sub>2</sub> giving palladacycle **12**. Reductive elimination of intermediate **12** in the presence of PhI(OAc)<sub>2</sub> provides cyclic product **6** and regenerates the active Pd(OAc)<sub>2</sub> catalyst for the next catalytic cycle. The exact role of Ag<sub>2</sub>O is not clear to us, it could be possible that the AgO<sup>-</sup> anion acts as a base to accelerate the transmetalation of boronic acid **2** into intermediate **12** and the Ag<sup>+</sup> ion acts as an oxidant to oxidize Ru(0) to Ru(II).

In conclusion, we have described a two-step synthesis of dibenzothiazines *via* a ruthenium-catalyzed *ortho* arylation of phenyl sulfoximines with phenyl boronic acids followed by intramolecular cyclization in the presence of Pd(OAc)<sub>2</sub>. Chiral dibenzothiazines were prepared efficiently by using chiral phenyl sulfoximine in a similar protocol.

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