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## Silver(I)-catalyzed highly *para*-selective phosphonation of 2-aryloxazolines†

Peng-Cheng Cui<sup>a</sup> and Guan-Wu Wang<sup>ib</sup> \*<sup>abc</sup>

**A silver-catalyzed phosphonation of 2-aryloxazolines has been accomplished. This protocol provides highly regioselective access to *para*-phosphonation products with good functional group tolerance and moderate to good yields via cross-dehydrogenation coupling. Mechanistic studies have shown that *para*-phosphonation products are obtained via a radical pathway. Furthermore, the directing oxazoline group in the *para*-phosphonation products is removable and can be converted to benzoic esters.**

Benzoic acid derivatives are widely used in drugs and agrochemicals.<sup>1</sup> Hence, their chemical modifications have potential applications. In the past few decades, transition-metal-catalyzed C–H bond activation has attracted extensive attention from chemists because of its high efficiency, atom economy and good functional group tolerance.<sup>2</sup> In this regard, modifications of benzoic acid derivatives utilizing C–H activation have been significantly developed.<sup>3</sup> Remote C–H activation of benzoic acid derivatives is relatively rare compared to *ortho*-C–H functionalization. To achieve remote C–H functionalization of benzoic acid derivatives, more complex auxiliary groups need to be utilized and tend to result in regioisomeric mixtures.<sup>4</sup> Thus, it is urgent to develop new methodologies for the *para*-functionalization of benzoic acid derivatives using easily available directing groups. The conversion of benzoic acids to 2-aryloxazolines is a commonly used method.

On the other hand, phosphorus-containing compounds exhibit a wide range of applications in pharmaceuticals, materials

and catalysis.<sup>5</sup> In recent decades, transition-metal-catalyzed C–H functionalization to construct C–P bonds has been used as an important methodology for the synthesis of organophosphorus compounds.<sup>6</sup> In 2006, Ishii and coworkers published the Mn(II)/Co(II)/O<sub>2</sub> redox-couple catalyzed phosphorylation of arenes; although providing a simple approach to the preparation of organophosphorus compounds, the site-selectivity was relatively poor (Scheme 1a).<sup>7</sup> For good selectivity, functional group-directed C–H activation protocols are particularly valuable. In 2013, the Yu group reported *ortho*-phosphorylation of arenes via heterocycle-directed *ortho*-palladation (Scheme 1b).<sup>8</sup> Subsequently, the Chen and Yu groups described a process for the *ortho*-phosphorylation of arenes using an inexpensive copper catalyst (Scheme 1b).<sup>9</sup> Since phosphorus reagents strongly coordinate with metals, a slow release of phosphorus reagents is required to ensure that the reaction proceeds smoothly. In 2019, the Wen, Zhang and Xu groups demonstrated a costly rhodium-catalyzed methodology to achieve *ortho*-phosphonation of electron-rich arylamine derivatives, and the use of oxidants was avoided because of the employed electrochemical oxidation (Scheme 1b).<sup>10</sup> In 2013, the Zhu and Cheng groups realized the Ag(I)/K<sub>2</sub>S<sub>2</sub>O<sub>8</sub>-mediated *ortho*-phosphorylation of electron-deficient arenes via a radical pathway (Scheme 1c).<sup>11</sup> In 2020, the Liang group reported the visible-light-induced *para*-C<sub>Ar</sub>–H phosphonation reactions with electron-rich arenes by cross-dehydrogenation coupling (CDC) (Scheme 1d).<sup>12</sup> As part of our continuing interest in remote C–H functionalization,<sup>13</sup> herein we disclose the silver-catalyzed highly *para*-phosphonation of electron-deficient 2-aryloxazolines (Scheme 1e).

In our initial investigation, we chose 4,4-dimethyl-2-phenyl-4,5-dihydrooxazole (**1a**) and di-*p*-tolylphosphine oxide (**2a**) as model substrates to screen the reaction conditions (see Table S1 in the ESI† for details). After extensive screenings, the optimal conditions were established as follows: with 10 mol% AgNTf<sub>2</sub> as the catalyst, 3.0 equiv. of K<sub>2</sub>S<sub>2</sub>O<sub>8</sub> as the oxidant, 0.5 equiv. of pivalic acid (PivOH) as the additive and MeCN as the solvent, the reaction of **1a** and **2a** (3.0 equiv.) performed best at 120 °C for 24 h under an argon atmosphere to give **3aa** in 73% yield.

<sup>a</sup> Hefei National Research Center for Physical Sciences at the Microscale and Department of Chemistry, University of Science and Technology of China, Hefei, Anhui 230026, P. R. China. E-mail: gwang@ustc.edu.cn

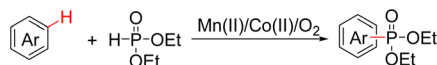
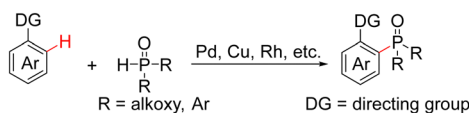
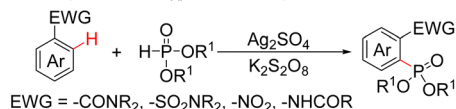
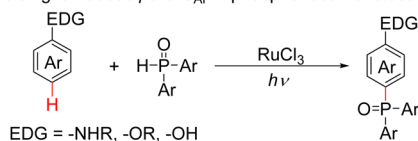
<sup>b</sup> Key Laboratory of Functional Molecular Solids, Ministry of Education, Anhui Laboratory of Molecule-Based Materials, and School of Chemistry and Materials Science, Anhui Normal University, Wuhu, Anhui 241002, P. R. China

<sup>c</sup> State Key Laboratory of Applied Organic Chemistry, Lanzhou University, Lanzhou, Gansu 730000, P. R. China

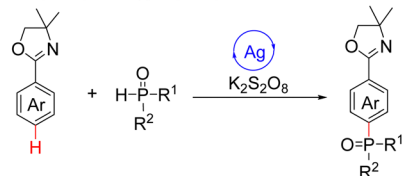
† Electronic supplementary information (ESI) available: Detailed experimental procedures and characterization data, <sup>1</sup>H NMR, <sup>13</sup>C NMR, <sup>19</sup>F NMR, <sup>31</sup>P NMR and HRMS of compounds **3** and **4**. See DOI: <https://doi.org/10.1039/d3cc06241d>



## Previous work

a) Mn(II)/Co(II)/O<sub>2</sub> redox couple catalyzed phosphorylation of arenes.b) Transition-metal-catalyzed *ortho*-C<sub>Ar</sub>-H phosphorylation/phosphonation of arenes.c) Silver-catalyzed *ortho*-C<sub>Ar</sub>-H phosphorylation of electron-deficient arenes.d) Visible-light induced *para*-C<sub>Ar</sub>-H phosphonation of electron-rich arenes.

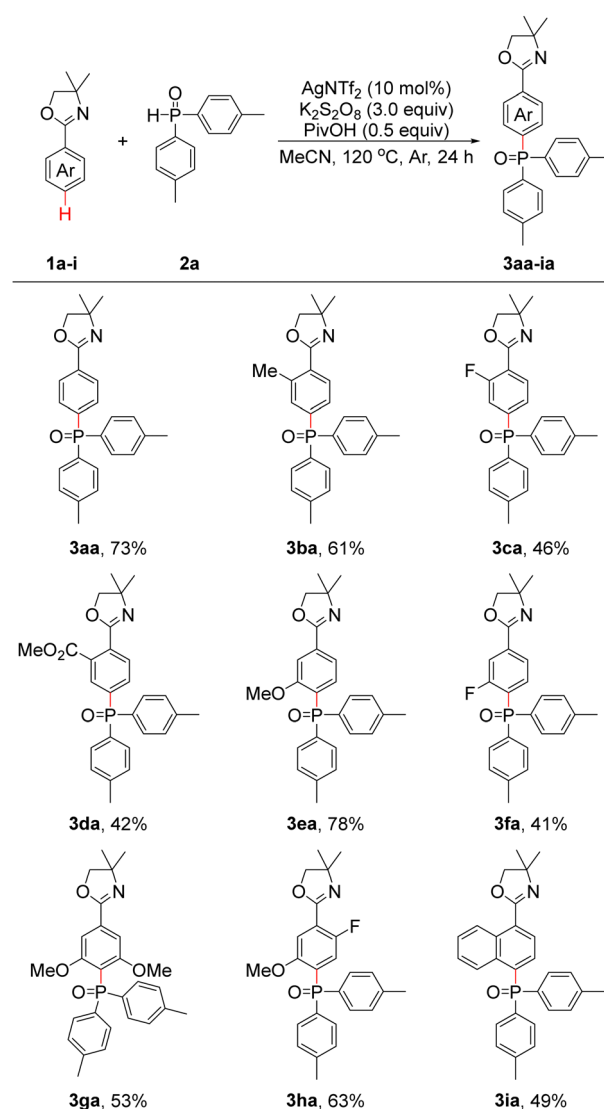
## This work

e) Silver-catalyzed *para*-C<sub>Ar</sub>-H phosphonation of electron-deficient arenes.

Scheme 1 Previous (a–d) and this (e) work on transition-metal-catalyzed C–H phosphorylation/phosphonation of arenes.

With the optimal conditions in hand, we then investigated the scope and functional group tolerance of 2-aryloxazolines **1**. As shown in Table 1, the reactions of **1a–i** with **2a** proceeded smoothly to afford *para*-phosphonation products **3aa–ia** in moderate to good yields. First, **1a**, which had an unsubstituted phenyl group, afforded **3aa** in 73% yield under optimal conditions. Substrate **1b** with an *ortho*-substituted Me group was able to provide **3ba** in 61% yield. Substrates **1c–d** with the *ortho*-substituted F and CO<sub>2</sub>Me groups could deliver **3ca** and **3da** in 46% and 42% yields, respectively. Substrates **1e–f** with the *meta*-substituted OMe and F groups also proceeded smoothly to afford **3ea–fa** in 41–78% yields. To our delight, when 3,5-dimethoxy-substituted substrate **1g** was employed, **3ga** was isolated in 53% yield. In addition, 2-(2-fluoro-5-methoxyphenyl)-4,4-dimethyl-4,5-dihydrooxazole (**1h**) could also react with **2a**, and **3ha** was obtained in 63% yield. These findings also provided a novel strategy for synthesizing multisubstituted arenes. For 4,4-dimethyl-2-(naphthalen-1-yl)-4,5-dihydrooxazole (**1i**), **3ia** could be produced in a yield of 49%.

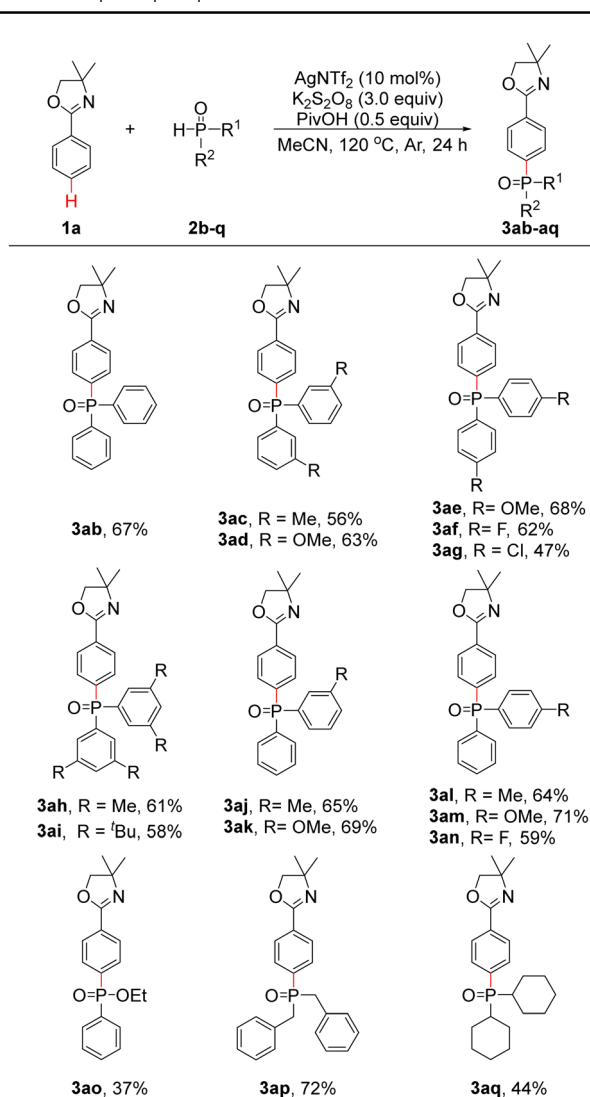
To further explore the generalizability of the substrates, the reactions of **1a** with a series of diarylphosphine oxides (**2b–n**), ethyl phenylphosphinate (**2o**) and dialkylphosphine oxides (**2p** and **2q**) were investigated under optimal conditions. As shown in Table 2, diphenylphosphine oxide (**2b**) was found to be suitable under our optimal conditions, and **3ab** was obtained in 67% yield. When bis(*m*-tolyl)phosphine oxide (**2c**)

Table 1 Scope of 2-aryloxazolines<sup>a, b</sup>

<sup>a</sup> Reaction conditions: **1a–i** (0.2 mmol), **2a** (0.6 mmol), AgNTf<sub>2</sub> (10 mol%), K<sub>2</sub>S<sub>2</sub>O<sub>8</sub> (0.6 mmol), PivOH (0.1 mmol), MeCN (1.0 mL) at 120 °C for 24 h under an argon atmosphere. <sup>b</sup> Isolated yields based on **1a–i**.

and bis(*m*-methoxyphenyl)phosphine oxide (**2d**) were employed, **3ac** and **3ad** were obtained in 56% and 63% yields, respectively. If bis(*p*-methoxyphenyl)phosphine oxide (**2e**) was used, **3ae** was obtained in 68% yield. Bis(*p*-fluorophenyl)phosphine oxide (**2f**) and bis(*p*-chlorophenyl)phosphine oxide (**2g**) proceeded well to produce **3af** and **3ag** in 62% and 47% yields, respectively. Delightedly, when bis(3,5-dimethylphenyl)phosphine oxide (**2h**) and bis(3,5-di-*tert*-butylphenyl)phosphine oxide (**2i**) were applied, **3ah** and **3ai** were generated in 61% and 58% yields, respectively. Subsequently, phenyl(tolyl)phosphine oxides (**2j** and **2l**), phenyl(methoxyphenyl)phosphine oxides (**2k** and **2m**) and phenyl(*p*-fluorophenyl)phosphine oxide (**2n**) were also applicable under optimal conditions, and **3aj–an** could be obtained in 59–71% yields. As a result, the synthesis of phosphine oxide products with three different aryl groups is made possible.



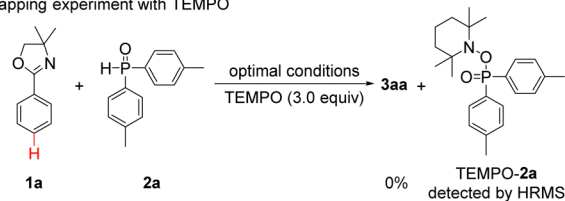
Table 2 Scope of phosphine oxides<sup>a, b</sup>

<sup>a</sup> Reaction conditions: **1a** (0.2 mmol), **2b-q** (0.6 mmol), AgNTf<sub>2</sub> (10 mol%), K<sub>2</sub>S<sub>2</sub>O<sub>8</sub> (0.6 mmol), PivOH (0.1 mmol), MeCN (1.0 mL) at 120 °C for 24 h under an argon atmosphere. <sup>b</sup> Isolated yields based on **1a**.

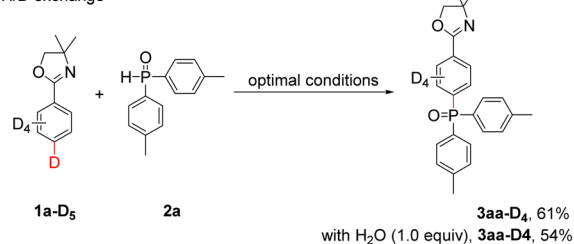
To our delight, when **2o** was investigated, **3ao** was generated in 37% yield. Products **3ap** and **3aq** could be isolated in 72% and 44% yields, respectively, when dibenzylphosphine oxide (**2p**) and dicyclohexylphosphine oxide (**2q**) were applied. Unfortunately, other substrates such as di(thiophen-2-yl)phosphine oxide and diethyl phosphonate failed to provide the desired products.

To gain insight into the reaction mechanism, additional experiments were conducted. First, **3aa** could not be obtained when 3.0 equiv. of the radical scavenger 2,2,6,6-tetramethylpiperidinoxy (TEMPO) was added under optimal conditions, and the TEMPO-**2a** adduct could be detected by high resolution mass spectrometry (HRMS) (Scheme 2a and Fig. S1, ESI<sup>†</sup>). This result showed that a radical pathway may be involved in the reaction. Second, when **1a-D<sub>5</sub>** was employed under optimal conditions, pure **3aa-D<sub>4</sub>** (Fig. S2, ESI<sup>†</sup>) was obtained in 61% yield. In addition, pure

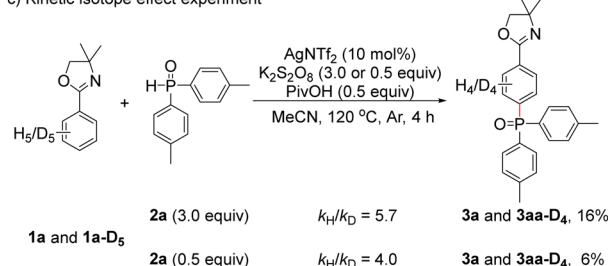
## a) Trapping experiment with TEMPO



## b) H/D exchange



## c) Kinetic isotope effect experiment

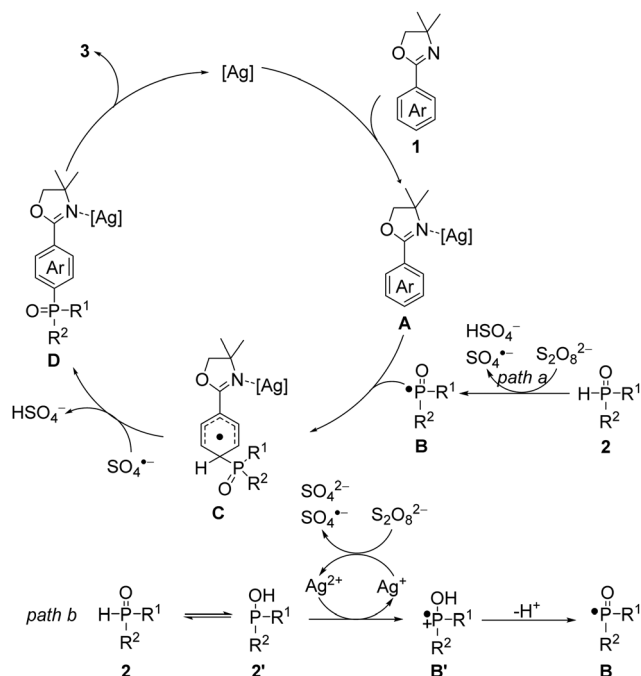


Scheme 2 Preliminary mechanistic studies.

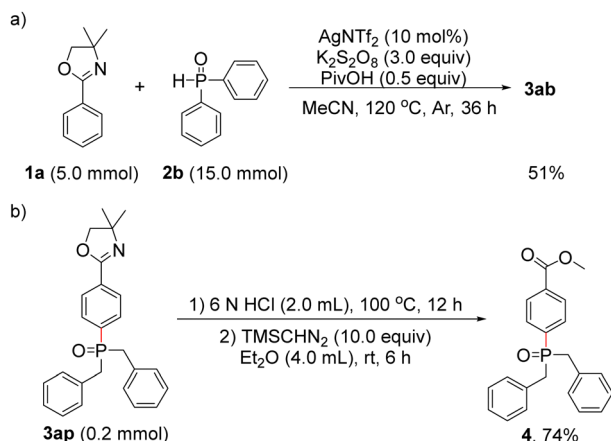
**3aa-D<sub>4</sub>** (Fig. S3, ESI<sup>†</sup>) could also be isolated in 54% yield with an additional 1.0 equiv. of H<sub>2</sub>O. These results indicated that no D/H scrambling occurred on the phenyl ring of **3aa-D<sub>4</sub>** and implied that no *ortho* C-H activation process took place (Scheme 2b). Third, an intermolecular kinetic isotope effect (KIE) experiment was performed, and the  $k_{\text{H}}/k_{\text{D}}$  value was determined to be 5.7 (Fig. S4, ESI<sup>†</sup>); when **2a** was reduced to 0.5 equiv., the  $k_{\text{H}}/k_{\text{D}}$  value was 4.0 (Fig. S5, ESI<sup>†</sup>), suggesting that C-H cleavage may be involved in the rate-determining step (Scheme 2c).

Based on the above findings of mechanistic studies and previous reports, a plausible reaction mechanism is shown in Scheme 3.<sup>11,14-18</sup> First, 2-aryloxazoline **1** interacts with the silver catalyst to form the intermediate **A**.<sup>14</sup> Similar oxazolinium ion can also be generated in the presence of PivOH and participates in subsequent steps as the intermediate **A**.<sup>15</sup> Meanwhile, phosphine oxide **2** is directly oxidized by the persulfate dianion (S<sub>2</sub>O<sub>8</sub><sup>2-</sup>) to give the key P-centered radical **B** together with a hydrogen sulfate anion (HSO<sub>4</sub><sup>-</sup>) and a sulfate radical anion (SO<sub>4</sub><sup>•-</sup>) (Scheme 3, *path a*).<sup>16</sup> Alternatively, **2** can exist in rapid equilibrium with the corresponding tautomeric isomer **2'**.<sup>17</sup> The Ag<sup>2+</sup> salt can oxidize **2'** to afford a Ag<sup>+</sup> species and cation radical **B'**, which may lead to the P-centered radical **B** after losing a proton; and the formed Ag<sup>+</sup> salt can be oxidized by a S<sub>2</sub>O<sub>8</sub><sup>2-</sup> to regenerate Ag<sup>2+</sup> together with a sulfate dianion (SO<sub>4</sub><sup>2-</sup>) and a SO<sub>4</sub><sup>•-</sup> (Scheme 3, *path b*).<sup>11,18</sup> Radical **B** then regioselectively undergoes a radical addition to the *para* position of **A** to give the intermediate **C**, which is subsequently oxidized and deprotonated by SO<sub>4</sub><sup>•-</sup> to give intermediate **D**. Finally, the *para*-phosphonation product **3** is obtained by





Scheme 3 Proposed mechanism.

Scheme 4 (a) Gram-scale synthesis of **3ab**. (b) Further synthetic transformation of **3ap**.

removal of the silver catalyst to complete the catalytic cycle. The oxidant  $K_2S_2O_8$  was mandatory and could promote the *para*-phosphonation alone, albeit in a low yield of 13% (Table S1, entries 22 and 23, ESI<sup>†</sup>). The beneficial effects of the silver salt (Table S1, entry 24, ESI<sup>†</sup>) and  $PivOH$  (Table S1, entry 25, ESI<sup>†</sup>) might originate from the activation of **1** by the formation of **A** and oxazolinium ion, respectively. Furthermore, the silver salt can also be employed as an oxidant to generate **B**.

To demonstrate the synthetic utility and importance of the present protocol, the reaction of **1a** with **2b** was carried out at the 5.0-mmol scale, and the *para*-phosphonation product **3ab** was isolated in 51% yield (Scheme 4a). We chose **3ap** as an example for further transformation, its oxazoline group could

be easily hydrolysed under acidic conditions, providing the *para*-phosphonated benzoic acid derivative **4** in 74% yield (Scheme 4b).<sup>19</sup>

In summary, a precise and efficient approach for the silver-catalyzed *para*-selective phosphonation of 2-aryloxazolines *via* cross-dehydrogenation coupling has been demonstrated. This protocol is able to provide highly regioselective *para*-phosphonation products and tolerates a wide range of functional groups. Furthermore, the directing oxazoline group in the products is removable and can be easily hydrolysed to give *para*-phosphonated benzoic esters. Preliminary mechanistic studies support that the *para*-phosphonation products are generated *via* a radical pathway. It is anticipated that the present protocol will also be valuable in pharmaceutical and materials science.

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## Conflicts of interest

There are no conflicts to declare.

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