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## Combination of $\text{RuCl}_3 \cdot x\text{H}_2\text{O}$ with PEG – a simple and recyclable catalytic system for direct arylation of heteroarenes *via* C–H bond activation†

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A simple and recyclable catalytic system for direct arylation of heteroarenes *via* C–H bond activation was developed with a relatively inexpensive  $\text{RuCl}_3 \cdot x\text{H}_2\text{O}$  as a catalyst and PEG-400 as a green medium without any additive or ligand. This system not only showed excellent functional group compatibility, but also the ratio of mono- to diarylated product was easily regulated by varying the reaction conditions. Moreover, this transformation could proceed under air and be easily scaled up to gram-scale in a low catalyst loading of 0.3 mol%. Particularly, a good yield of 85% was obtained after this catalyst was recycled six times.

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### 1. Introduction

In recent years, N-heteroaromatic scaffolds are becoming an increasing necessity, due to the structural motifs for a wide range applications in pharmaceuticals, natural products, functional materials<sup>1</sup> and ligands for highly efficient and selective catalysis.<sup>2</sup> Thus, there is an important need to develop an efficient synthetic methodology to make up the building blocks. Although transition metal catalyzed cross-coupling reaction of aryl halides with organometallic reagents to synthesize a series of heterocyclic compounds has matured to being a reliable tool,<sup>3</sup> catalytic direct arylation *via* activation of C–H bond still garners tremendous interest due to obviating prefunctionalization,<sup>4,5</sup> which addresses the issues on prolix steps of organic synthesis, formation of undesired side-product and use of auxiliary chemicals. Nevertheless, transition-metal-complexes catalyzed direct arylation *via* C–H activation still confronts various challenges. The preparation of complexes suffers from a tedious synthesis process and they are difficult to be recycled after reaction. For direct arylation in heterogeneous catalysis, polymer supported metal catalysts, such as PVP supported Pd-nanoparticle,<sup>6a</sup> Pd(*n*)@microporous polymers<sup>6b</sup> and heterogenization of homogeneous catalysts, such as polystyrene-supported palladium complex,<sup>6c</sup> polymer-supported pyridine-Pd,<sup>6d</sup> poly(ethyleneglycol)-bound SCS-Pd complexes<sup>6e</sup> and MOP-PPh<sub>3</sub>-Pd<sup>6f</sup> seem to be extremely effective ways to accomplish the

separation and recycling of the catalysts. To date, the preparation of these catalysts is not easy and leaching of metals cannot be inhibited completely. In addition, NMP or toluene as solvent<sup>7</sup> is highly toxic and is usually classified to be detrimental to the environment. To meet the requirements of green chemistry, some eco-friendly solvents such as CPME, 2-MeTHF and dialkyl carbonates have been chosen as the reaction media in direct arylation.<sup>8</sup> Although water is a better alternative, the metal complex as a catalyst must be water-soluble and insensitive to water.<sup>9</sup>

Polyethylene glycol has been the perfect reaction media, which offers a suitable strategy to meet the requirement of green chemistry.<sup>10</sup> To date, several successful examples have been reported using PEG as reaction medium, such as direct alkynylation of heteroarene,<sup>10a</sup> one-pot synthesis of propargylamines, aminoindolizine, 2,4,6-triaryl-pyridines, piperidines, and 3,4-dihydro-pyrimidinones,<sup>10b-f</sup> oxidative coupling of 2-arylbenzimidazoles with alkynes<sup>10g</sup> and hydroxylation of 2-aryl-pyridine.<sup>10h</sup> However, ruthenium catalyzed direct arylation in PEG, which was reported by Ackermann *et al.*<sup>11a</sup> and Hiebel *et al.*,<sup>11b</sup> under nitrogen atmosphere gave the arylated products with a moderate yield but in air it was inactive.<sup>11a</sup> In addition, most of the catalytic systems for direct arylation are required to be performed in oxygen-free conditions.<sup>12</sup> Hence, to develop a simple and sustainable method was highly desirable with green chemistry into consideration. Herein, in continuation to our earlier studies on developing improved methodologies for direct arylation,<sup>16</sup> we propose a simple and recyclable catalytic system for direct arylation of heteroarenes by employing  $\text{RuCl}_3 \cdot x\text{H}_2\text{O}$  as the catalyst, inexpensive aryl chlorides as the arylating reagents and PEG-400 as the “green” reaction medium without any additives or ligands. The strategy that was accomplished in air gave a satisfactory yield and a relatively high

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selectivity. Particularly, the ratio of mono- to diarylated product was easily regulated by varying the reaction conditions.

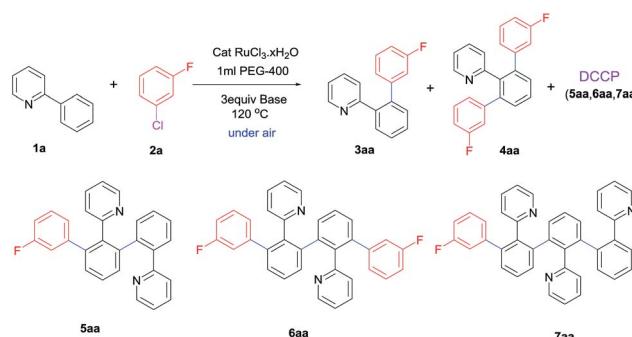
## 2. Results and discussion

### 2.1 Effect of the reaction conditions

In initial investigation, the model reaction of 2-phenylpyridine **1a** (1 mmol) with coupling partner 1-chloro-3-fluorobenzene **2a** (2.3 equiv.) was conducted in 1 mL PEG-400 to determine the optimum conditions. First, effect of various bases on this catalytic system was tested without any additive or ligand (Table 1). In the beginning, CsF, K<sub>3</sub>PO<sub>4</sub>, K<sub>2</sub>CO<sub>3</sub>, NaHCO<sub>3</sub>, Na<sub>2</sub>SO<sub>3</sub> and EtONa were substantially less effective on the transformation (Table 1, entries 1–6). Interestingly, when CH<sub>3</sub>COOK was applied in the system under similar conditions, an excellent yield of 94% for target products **3aa** and **4aa**, and relatively high selectivity towards monoarylation were obtained in 1 h (Table 1, entry 7-II). However, prolonging the reaction time resulted in a decrease of mono-arylated product **3aa** (Table 1, entries 7-IV and 7-VI), which was mainly attributed to the formation of dehydrogenative cross-coupling products **5aa**, **6aa**, **7aa**. Interestingly, this system also afforded the target products **3aa** and **4aa** in the yield of 82% even if the reaction time was shortened to 15 min (Table 1, entry 7-I). To the best of our knowledge, the activity of this transformation was unprecedented. Among the reported cases,<sup>7</sup> to obtain a satisfactory yield for direct arylation, the reaction time was at least 60 minutes,<sup>7a</sup> usually 20–40 hours<sup>7b–e</sup> under similar conditions. The ratio of monoarylation to diarylation was further improved to 5.3/1 when **1a/2a** was adjusted to 1.2/1 (see ESI†). Compared with CH<sub>3</sub>COOK, CH<sub>3</sub>COONa as a base also displayed similar activity and selectivity (Table 1, entry 8). Furthermore, it was notable that LiOH·H<sub>2</sub>O (3 equiv.) afforded a high yield of 98% and proved highly efficient to form diarylated product **4aa** (Table 1, entries 9-II and 9-III). Apart from LiOH·H<sub>2</sub>O, the introduction of KOH, NaOH, CsOH, Li<sub>2</sub>CO<sub>3</sub>, and *t*-BuOLi resulted in discontented yields (Table 1, entries 10–14). Considering the promotional effect of Li<sup>+</sup> and CH<sub>3</sub>COO<sup>–</sup> on the reaction, we were motivated to probe the role of CH<sub>3</sub>COOLi in the reaction. However, it gave the target products with a moderate yield of 50% (Table 1, entry 15). NET<sub>3</sub> was also a good base in this transformation, but it was unfavorable to improve the selectivity towards monoarylated or diarylated product (Table 1, entry 16). In the absence of a base, almost no target products were obtained, and only oligomerization products **8**, **9** and **10** were observed in this process, highlighting the importance of a base for the direct arylation process (Table 1, entry 17) (Fig. 1). Based on the above findings, it was clearly indicated that the system took advantage of high reactivity and easily regulated the ratio of diarylated to monoarylated product. The diarylation was facilitated using 5 mol% RuCl<sub>3</sub>·xH<sub>2</sub>O as a catalyst and 3 equiv. LiOH·H<sub>2</sub>O as a base at 120 °C for 24 h. Intriguingly, the system strongly favoured the monoarylation just by switching the base to CH<sub>3</sub>COOK and shortened the reaction time to 1 h. In this reaction process, the hydroxyl group of PEG-400 played the role of a reductant for the precatalyst RuCl<sub>3</sub>·xH<sub>2</sub>O.<sup>13</sup>

The further solvent screening indicated that target products could not be generated in toluene or water (Table 1, entry 18-I). It

Table 1 Effect of reaction conditions on arylation of 2-phenylpyridine with 1-chloro-3-fluorobenzene<sup>a</sup>



Entry	Base	Order	Time [h]	Conv. [%]	3aa : 4aa : DCCP [%]
1	CsF	—	17	18	28 : 50 : 22
2	K <sub>3</sub> PO <sub>4</sub>	—	17	20	11 : 54 : 35
3	K <sub>2</sub> CO <sub>3</sub>	—	17	64	50 : 33 : 17
4	NaHCO <sub>3</sub>	—	17	37	57 : 19 : 24
5	Na <sub>2</sub> SO <sub>3</sub>	—	17	Trace	—
6	EtONa	—	17	Trace	—
7 <sup>b</sup>	CH <sub>3</sub> COOK	I	0.25	82	71 : 29 : 0
		II	1	98	63 : 33 : 4
		III <sup>c</sup>	1	91	64 : 31 : 5
		IV	5	98	44 : 33 : 23
		V <sup>d</sup>	5	98	31 : 58 : 11
		VI	17	>99	22 : 43 : 35
8 <sup>b</sup>	CH <sub>3</sub> COONa	I	1	96	74 : 21 : 5
		II	17	97	51 : 24 : 25
9 <sup>b</sup>	LiOH·H <sub>2</sub> O	I	2	91	33 : 66 : 1
		II	17	>99	17 : 82 : 1
		III	24	>99	15 : 84 : 1
10 <sup>b</sup>	<i>t</i> -BuOLi	—	17	67	26 : 65 : 9
11 <sup>b</sup>	Li <sub>2</sub> CO <sub>3</sub>	I	1	60	22 : 5 : 73
		II	17	69	18 : 2 : 80
12 <sup>b</sup>	KOH	—	17	50	10 : 10 : 80
13 <sup>b</sup>	NaOH	—	17	65	2 : 6 : 92
14 <sup>b</sup>	CsOH	—	17	Trace	—
15 <sup>b</sup>	CH <sub>3</sub> COOLi	I	1	73	65 : 21 : 14
		II	17	85	46 : 13 : 41
16 <sup>b</sup>	NET <sub>3</sub>	I	1	98	47 : 51 : 2
		II	17	98	37 : 60 : 3
17 <sup>b</sup>	—	—	17	40	0 : 0 : 100
18 <sup>e</sup>	CH <sub>3</sub> COOK	I	17	Trace	—
		II	2	66	74 : 12 : 14
		III	2	83	68 : 19 : 13

<sup>a</sup> 2-Phenylpyridine (1 mmol), 3-fluoro-chlorobenzene (2.3 mmol), PEG-400 (1 mL), RuCl<sub>3</sub>·xH<sub>2</sub>O (5 mol%), and base (3 equiv.), 120 °C, under air. Yield was isolated yield. <sup>b</sup> (Entries 7–17) Yield and ratio of the mono- and diarylated products were determined by HPLC. <sup>c</sup> (3 mol%) RuCl<sub>3</sub>·xH<sub>2</sub>O. <sup>d</sup> Adding 3-fluoro-chlorobenzene (2.3 equiv.) again after 0.5 h. <sup>e</sup> I – toluene, water or MeOH as solvent; II – ethylene glycol as solvent; III – ethylene glycol as solvent and adding 18-crown-6 (1 mmol) to the reaction system.

was found that a moderate yield of 57% was also obtained in ethylene glycol, but the direct arylation proved to be ineffective in MeOH as its low boiling point led to a low reaction temperature (Table 1, entries 18-I and 18-II). Compared to ethylene glycol, PEG



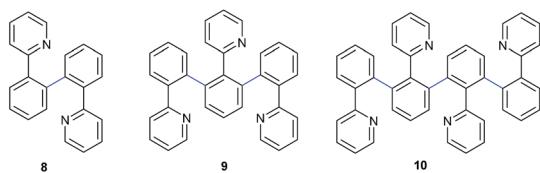


Fig. 1 The oligomerizations of 2-phenylpyridine.

Table 2 Effect of substrate to catalyst ratio on direct arylation of 2-phenylpyridine with 1-chloro-3-fluorobenzene<sup>a</sup>

Entry	S/C	Time (h)	Yield [%]	3aa/4aa [%]
1	20	1	90	77 : 23
2	40	1	91	80 : 20
3	160	1	89	81 : 19
4	300	1	50	84 : 16
		4	92	78 : 22
5	600	10	32	82 : 18
		24	34 (40)	80 : 20

<sup>a</sup> Reaction conditions: 2-phenylpyridine (*n* mmol), 3-fluoro-chlorobenzene (*n* + 0.3 mmol), PEG-400 as reaction medium (1 mL), RuCl<sub>3</sub>·xH<sub>2</sub>O (0.05 mmol), and CH<sub>3</sub>COOK (2 equiv.) and the reaction mixture was heated to 120 °C under air. The yield and ratio of mono- to diarylated product determined by HPLC.

was a better solvent. The weak interaction of PEG with ruthenium species might enhance the reactivity, analogous to the coordination of crown ether with ruthenium. In order to demonstrate it, 18-crown-6 was added to ethylene glycol, and it was found that this transformation was significantly improved (Table 1, entry 18-III). The above results suggested that the coordination of PEG with ruthenium, which was similar to the role of crown ether, promoted the performance of ruthenium in addition to the reduction role of hydroxyl group in PEG. Furthermore, the effect of reaction temperature indicated that a higher temperature of 150 °C resulted in more dehydrogenative cross-coupling products (DCCP) and a lower temperature of 90 °C caused a decrease of the catalytic activity (see ESI†). The catalytic performance of this simple system under optimum conditions was comparable to NHC-ruthenium(II) complexes,<sup>14</sup> phosphine-[RuCl<sub>2</sub>(*p*-cymene)]<sub>2</sub>,<sup>15</sup> [RuCl<sub>2</sub>(*n*<sup>6</sup>-benzene)MOTPP],<sup>16</sup> and iridium-ruthenium bimetallic complexes bearing triazolodiyliidene.<sup>17</sup>

On taking catalyst loading into consideration, when the amount of ruthenium was decreased to 3 mol%, this system gave a satisfactory yield of 86% (Table 1, entry 7-III). Gratifying, we were surprised to find that this transformation could be easily scaled up to gram level with a comparable yield of 93% when S/C was extended to 300/1 for 4 h (Table 2).

## 2.2 The scope of the proposed direct arylation and recyclability of the catalyst

With optimum reaction conditions, we explored the scope of direct arylation protocol. It was found that this system was of wide substrate suitability. Moreover, diarylated derivatives of heteroarene were predominantly achieved by employing aryl chlorides

as arylating reagents containing a variety of functional groups (Table 3). Overall, the reaction provided the desired products in high yields (77–99%). It was important to note that direct arylation of 2-phenylpyridine proceeded with high efficiency under an ambient atmosphere, highlighting the user-friendly nature of the proposed protocol. Chlorobenzene **2b** was used as arylating reagent to give an impressive yield of 92% (Table 3, entry 1). Notably, all aryl chlorides with substituents derived from electron-rich groups (–OCH<sub>3</sub>, –CH<sub>3</sub>) as well as electron-deficient groups (–CF<sub>3</sub>, –F, –COCH<sub>3</sub>, –CN) were converted into the desired products with high yields under the reaction conditions (Table 3, entries 3, 4, 6, 7, 9, 10, 12–15 and 21). Those results clearly indicated that this system was insensitive to electronic factor. At the same time, heteroaryl chlorides were also surveyed to furnish the desired products with a comparably better yield (Table 3, entries 16–18). However, for aryl chlorides containing a substituent in *ortho*-position, they afforded diarylated products with relatively low yields due to steric hindrance of *ortho*-substituent<sup>2e</sup> (Table 3, entries 2, 5 and 11). However, for 1-chloro-2-fluoro-benzene and 3-chloropyridine (Table 3, entries 8 and 19), only a trace amount of product was obtained. Furthermore, 4-chloronitrobenzene also gave a trace amount of target products in accordance with the literature<sup>2e</sup> (Table 3, entry 20). It was noteworthy that the use of (hetero)aryl chlorides bearing various electron-deficient or electron-rich groups resulted in relatively high selectivity towards diarylation in the presence of LiOH·H<sub>2</sub>O except for **2n**, **2o**, **2q**, **2r**. Even so, the relatively high yield of diarylated product was obtained with 3 equiv. CH<sub>3</sub>COOK as the base for 1 h (Table 3, entries 14, 15, 17 and 18). On using LiOH with CH<sub>3</sub>COOK (2 equiv.), high selectivity towards monoarylation was achieved (Table 3, entries 13, 15 and 18). To explore the versatility of this transformation, 1-phenylpyrazole and 2-phenyl-2-oxazoline were also surveyed to afford the desired products in moderate yields of 72% and 57%, respectively (Table 3, entries 22 and 23). Moreover, the results were comparable to catalytic systems generated from Rh(III) complex in the presence of Ag salts under nitrogen atmosphere<sup>5b</sup> and [Ru(O<sub>2</sub>CMes)<sub>2</sub>(*p*-cymene)] with *p*-TsCl<sup>5f</sup> for direct arylation of 1-phenylpyrazole and 2-phenyl-2-oxazoline. Moreover, 2-phenylpyridine derivatives bearing various substituents on the phenyl ring were well tolerated (Table 3, entries 24, 25, 27 and 28). Substrate **1f** was almost completely converted to monoarylated product due to steric hindrance of the methyl group (Table 3, entry 26). The high selectivity towards monoarylation, being caused by the steric hindrance of methyl group, was similar to the reported results.<sup>7b,18</sup>

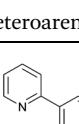
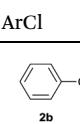
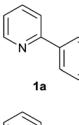
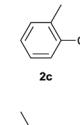
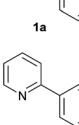
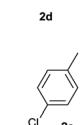
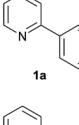
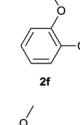
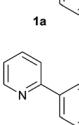
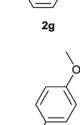
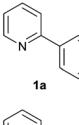
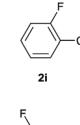
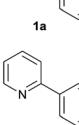
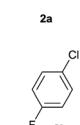
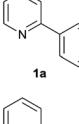
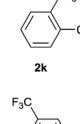
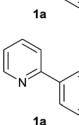
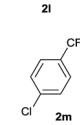
On the other hand, this system could be conveniently re-used (Table 4). A good yield of 85% was still obtained with decreased selectivity after the catalyst was recycled 6 times. To the best of our knowledge, this is the first successful example for arylation achieving both gram-scale preparation with a low catalyst loading of 0.3 mol% and maintaining its high activity after several recycles.

## 2.3 Mechanism of direct arylation

It was reported that catalytically active species were palladium nanoparticles for palladium-catalyzed direct arylation in PEG.<sup>11a</sup> In this case, the mercury poison experiment,<sup>19</sup> being an efficient



**Table 3** Ruthenium-catalyzed *ortho*-arylation of heteroarenes with (hetero)aryl chlorides<sup>a</sup>

Entry	Heteroarene	ArCl	Yield (%)	3/4 (%)	
				3 a-f, a-s	4 a-f, a-s
1			92	37 : 63 <sup>d</sup>	
2			41	54 : 46	15
3			95	28 : 72 <sup>d</sup>	16
4			84	39 : 61	17
5			44	39 : 61	18
6			77	30 : 70	19
7			90	26 : 74	20
8			Trace	—	21
9			95	22 : 78 <sup>d</sup>	22
10			88	43 : 57	23
11			34	47 : 53	24
12			94	30 : 70	25
13			90 67 <sup>c</sup>	11 : 89 73 : 27	26

**Table 3 (Contd.)**

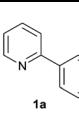
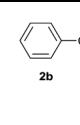
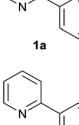
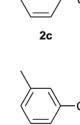
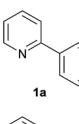
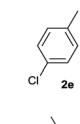
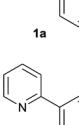
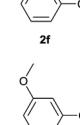
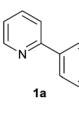
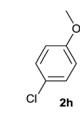
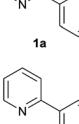
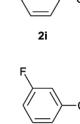
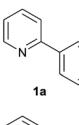
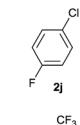
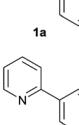
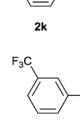
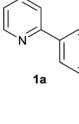
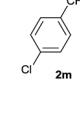
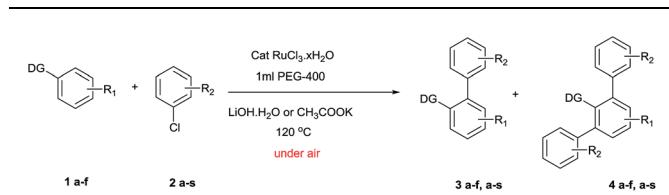
Entry	Heteroarene	ArCl	Yield (%)	3/4 (%)	
				3 a-f, a-s	4 a-f, a-s
14			92 <sup>b</sup>	58 : 42	
15			91 <sup>b</sup> 69 <sup>c</sup>	32 : 68 78 : 22	
16			72	51 : 49	
17			28 88 <sup>b</sup>	75 : 25 18 : 82 <sup>d</sup>	
18			36 83 <sup>b</sup> 68 <sup>c</sup>	67 : 33 <sup>d</sup> 31 : 69 <sup>d</sup> 75 : 25 <sup>d</sup>	
19			Trace	—	
20			Trace	—	
21			65 <sup>b</sup>	85 : 15	
22			72	58 : 42	
23			57 <sup>b</sup>	77 : 23	
24			92	22 : 78	
25			84	62 : 38	
26			80 60 <sup>b</sup>	95 : 5 100 : 0	

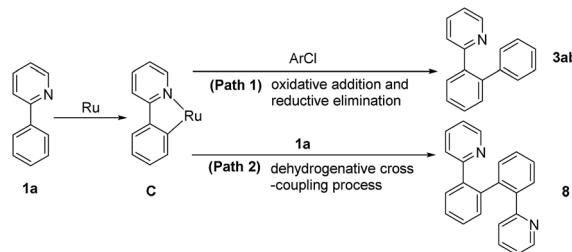


Table 3 (Contd.)



Entry	Heteroarene	ArCl	Yield (%)	3/4 (%)
27	1g	2m	80 <sup>b</sup>	75 : 25
28	1h	2m	32 <sup>b</sup>	—

<sup>a</sup> Heteroarene (1 mmol), (hetero)aryl chloride (2.3 mmol), PEG-400 as reaction medium (1 mL), RuCl<sub>3</sub>·xH<sub>2</sub>O (5 mol%), and LiOH·H<sub>2</sub>O (3 equiv.) and the reaction mixture was heated to 120 °C under air for 24 h and isolated yield of the products. <sup>b</sup> CH<sub>3</sub>COOK (3 equiv.) as base for 1 h. <sup>c</sup> Aryl chloride (1 mmol), 2-phenylpyridine (1.2 mmol), CH<sub>3</sub>COOK (2 equiv.) as base for 1 h. <sup>d</sup> Ratios determined by <sup>1</sup>H NMR spectroscopy.



Scheme 1 Two competitive paths for arylation of 2-phenylpyridine.

routes in this transformation. In the proposed catalytic system, apart from the desired products 3aa and 4aa, formation of by-product DCCP revealed that the reaction routes were in good agreement with the reported results.<sup>21</sup>

When CH<sub>3</sub>COOK was employed as the base, the reaction proceeded preferentially in path 1 (Table 1, entries 7-I and 7-II) and no DCCP was observed at the beginning of the reaction. If the reaction time was prolonged to 5 h, the expended mono-arylated product 3aa was almost completely transferred to DCCP (Table 1, entry 7-IV). This possibly resulted from the decrease in the chlorobenzene concentration, which slowed its oxidative addition rate with cyclometalated ruthenium intermediate. Moreover, the accumulated 3aa would promote the formation of DCCP. It was thought that extra 1-chloro-3-fluorobenzene (2.3 equiv.) was introduced into the system after the reaction began for 0.5 h; the yield of diarylated product 4aa was significantly improved and dehydrogenation crossing-coupling processes were greatly inhibited (Table 1, entry 7-V). The result indicated that high concentration chlorobenzene was beneficial to direct the arylated reaction in path 1. Based on concerted-metalation-deprotonation mechanism (CMD) with the assistance of carboxylate,<sup>22</sup> a mechanism in the presence of potassium acetate was proposed in Scheme 2. First, Ru(III) species was reduced to Ru(II) by hydroxyl group in PEG-400. Moreover, the electron density of Ru was increased due to the weakly coordinated effect of PEG with Ru, which resulted in enhancing the reactivity, similar to the role of a ligand. The reaction began with the cyclometalation of 2-phenylpyridine 1a to form intermediate C through acetate-assisted dehydrogenation process. Subsequently, intermediate C reacted with chlorobenzene 2b involving an oxidative-addition process to generate intermediate D, which progressed by reduction elimination to give the monoarylated product 3ab and active ruthenium species A. When most of 2b was used up, intermediate C easily reacted with 3ab or 1a to afford DCCP. Mono-arylated product 3ab could also undergo the similar process to give the corresponding diarylated product and DCCP.

When the reaction was conducted with LiOH, the rate of this reaction was dramatically decreased, but this system gave a relatively high selectivity to diarylated product and the formation of DCCP was greatly inhibited. It was suggested that the coordination of pyridyl group with Li<sup>+</sup> could slow the chelation-directed C-H activation process. Based on the same reason, it also blocked the dehydrogenation cross-coupling route owing to a need to activate a second 2-phenylpyridine

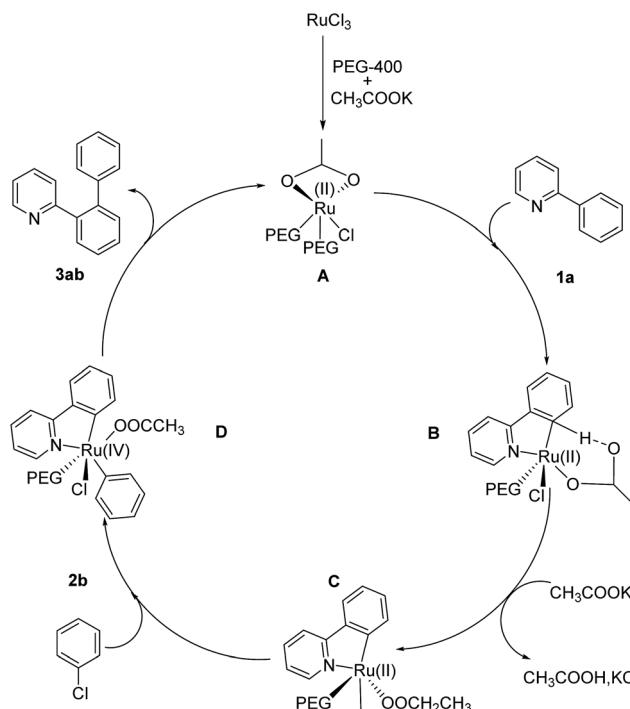
Table 4 Recyclability of the catalyst for direct arylation of 2-phenylpyridine with 1-chloro-3-fluorobenzene<sup>a</sup>

Entry	Run	Time [h]	Yield [%]	Ratio (3aa/4aa)
1	1	5	94	22 : 78
2	2	5	85	23 : 77
3	3	12	93	30 : 70
4	4	12	94	37 : 63
5	5	12	81	57 : 43
6	6	24	85	64 : 36

<sup>a</sup> Reaction conditions: 2-phenylpyridine (1 mmol), 3-fluoro-chlorobenzene (2.3 mmol), PEG-400 as reaction medium (1 mL), RuCl<sub>3</sub>·xH<sub>2</sub>O (5 mol-%), LiOH·H<sub>2</sub>O (3 equiv.), 120 °C, and under air. The ratio of mono- to diarylation product was determined by HPLC. Equivalent base needed to be added in every recycling.

method to distinguish nanoparticle catalysis and homogeneous catalysis, showed that the activity of this catalytic system did not have any loss. The result proved that the catalytically active species were not ruthenium nanoparticles. Both experimental studies and density functional theory (DFT) calculations on ruthenium or palladium catalyzed direct arylation of heteroarene *via* C-H activation have proposed that the reaction progressed *via* path 1 or path 2 (Scheme 1).<sup>20-22</sup> In both paths, formation of cyclometalated ruthenium intermediate C was a key step. The intermediate C reacted with chlorobenzene *via* oxidative addition/reductive elimination sequence to give target products or 2-arylpypyridine through the similar *ortho*-cyclometallated process to afford dehydrogenative cross-coupling products. Therefore, path 1 and path 2 were competitive





Scheme 2 Proposed mechanism for ruthenium-catalyzed direct arylation of 2-phenylpyridine.

C–H bond by a chelation-directed process, but it did not affect oxidative addition with chlorobenzene. Consequently, the system with LiOH was not only in favour of diarylation, but also greatly inhibited the formation of DCCP. To obtain some experimental evidence, PEG-400 (200  $\mu$ L), 2-phenylpyridine (1 mmol) and Li<sup>+</sup> (1 mmol) were added to a Schlenk tube and stirred under air at 120 °C for 17 h. Evidently, the coordination of the Li<sup>+</sup> with 2-phenylpyridine was clearly observed in both <sup>1</sup>H and <sup>13</sup>C NMR spectra (Fig. 2 and 3). For other lithium salts, *t*-BuOLi as base also strongly favors the diarylation of 2-phenylpyridine, analogous to the promotional role of LiOH (Table 1, entry 10). However, Li<sub>2</sub>CO<sub>3</sub> or CH<sub>3</sub>COOLi as base showed a different result with LiOH and *t*-BuOLi (Table 1, entries 11 and

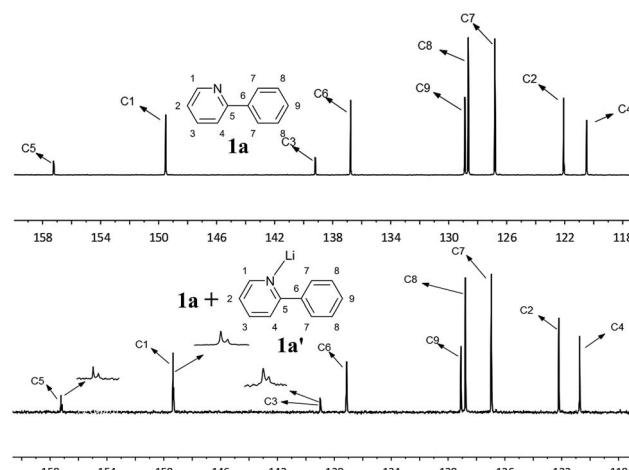


Fig. 3 <sup>13</sup>C NMR spectra for 2-phenylpyridine 1a in PEG and the mixture of 2-phenylpyridine and Li salt 1a' with 1a in PEG.

15). The promotional mechanism of LiOH was not in agreement with the CMD mechanism. The reaction possibly proceeded by a chelation-directed C–H activation with OH<sup>−</sup> to abstract proton to generate a cyclometalated ruthenium intermediate. Next, the intermediate could react with chlorobenzene *via* oxidative addition/reductive elimination sequence to give the mono- and di-arylated product or a second 2-arylpypyridine through the similar *ortho*-cyclometallated process to afford DCCP.

### 3. Conclusions

We developed a simple and recyclable catalytic system for direct arylation using RuCl<sub>3</sub>·xH<sub>2</sub>O as a precatalyst, PEG-400 as environmental benign medium and inexpensive aryl chlorides as electrophilic reagents. The reaction met the environment friendly requirements and the strategy was facile and did not require the introduction of any additive or ligand. This system showed high tolerance towards various functional groups. Particularly, selectivity towards mono- or diarylated product could be regulated by varying the reaction conditions. Furthermore, this transformation could be easily scaled up to the gram level with 0.3 mol% loading of ruthenium. A reasonable mechanism for this transformation was also proposed.

### 4. Experimental

#### 4.1 Typical experimental procedure for direct arylation of 2-phenylpyridine under air

RuCl<sub>3</sub>·xH<sub>2</sub>O (104 mg, 0.5 mmol) and PEG-400 (10 mL) was taken in an oven-dried and clean round-bottom flask. The mixture was stirred at room temperature until RuCl<sub>3</sub>·xH<sub>2</sub>O was dispersed to PEG-400 completely. Furthermore, 2-phenylpyridine (1a, 1 mmol), 1-chloro-3-fluorobenzene (2a, 2.3 mmol), LiOH·H<sub>2</sub>O (126 mg, 3 mmol) and PEG-400 (1 mL) containing RuCl<sub>3</sub>·xH<sub>2</sub>O (0.05 mmol) were added to a Schlenk tube. The mixture was stirred at 120 °C under air for 24 h. After cooling to room temperature, the reaction mixture was diluted with 10 mL

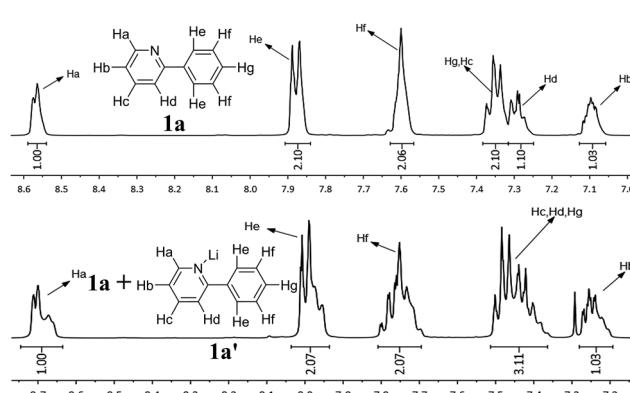


Fig. 2 <sup>1</sup>H NMR spectra for 2-phenylpyridine 1a in PEG and the mixture of 2-phenylpyridine and Li salt 1a' with 1a in PEG.



water. The aqueous layer was extracted with  $\text{Et}_2\text{O}$  (20 mL) and  $\text{EtOAc}$  (30 mL). The combined organic phase was washed with  $\text{H}_2\text{O}$  (20 mL) and brine (20 mL) and dried with  $\text{Na}_2\text{SO}_4$  and concentrated under reduced pressure. The resulting residue was separated through a flash silica gel column chromatography (petroleum ether/ $\text{EtOAc}$  = 16 : 1) to obtain arylated products (306 mg, 95%). The ratio of mono- to diarylated product was determined by  $^1\text{H}$  NMR spectroscopy. The arylated products were separated using silica gel column chromatography (petroleum ether/ $\text{CH}_2\text{Cl}_2$  = 1 : 16) to afford target products **3aa** and **4aa**, respectively. The product was characterized by  $^1\text{H}$  NMR and  $^{13}\text{C}$  NMR spectroscopy.

## 4.2 Recycling experimental procedures

In a reaction tube, 2-phenylpyridine (**1a**, 1 mmol), 1-chloro-3-fluorobenzene (**2a**, 2.3 mmol),  $\text{LiOH}\cdot\text{H}_2\text{O}$  (126 mg, 3 mmol) and PEG-400 (1 mL) containing  $\text{RuCl}_3\cdot x\text{H}_2\text{O}$  (0.05 mmol) were stirred at 120 °C under air for 24 h. At the end of the reaction, reaction mixture was cooled to room temperature and extracted by employing  $\text{Et}_2\text{O}$  (50 mL). Furthermore, 2-phenylpyridine (**1a**, 1 mmol), 1-chloro-3-fluorobenzene (**2a**, 2.3 mmol),  $\text{LiOH}\cdot\text{H}_2\text{O}$  (126 mg, 3 mmol) was added to PEG solution, the reaction and operation were repeated as mentioned above.

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