

Cite this: *Chem. Sci.*, 2022, **13**, 12906

All publication charges for this article have been paid for by the Royal Society of Chemistry

Received 12th July 2022  
Accepted 14th October 2022DOI: 10.1039/d2sc03877c  
rsc.li/chemical-science

## Introduction

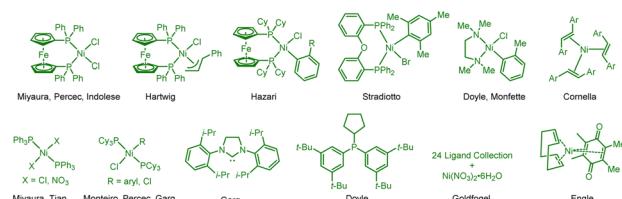
The Pd-catalyzed Suzuki–Miyaura Coupling (Pd-SMC) of aryl halides and arylboron reagents is one of the most practical catalytic methods for  $C(sp^2)$ – $C(sp^2)$  bond formation.<sup>1–3</sup> In the pharmaceutical industry, it ranks among the top 2–3 most widely practiced reactions in both medicinal and process chemistry groups.<sup>4,5</sup> Many ligands have been identified for the Pd-SMC; however, fewer ligands for catalysis by earth-abundant metals (Ni, Cu, Co and Fe) have been developed. Given the increased emphasis on greener and more sustainable catalysis, non-precious metal alternatives for the Pd-SMC are being actively sought.<sup>6</sup>

The Ni-catalyzed Suzuki–Miyaura Coupling (Ni-SMC), known since the mid-1990s, is a promising alternative to Pd-SMC and even has its advantages (Scheme 1).<sup>7–9</sup> For example, aryl electrophiles normally unreactive under Pd catalysis, such as carbamates, can be coupled with Ni.<sup>10</sup> A key challenge to the wider application of the Ni-SMC, however, is the significantly reduced scope of several key heteroarylboron nucleophiles with

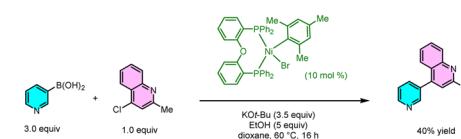
Ni *vs.* Pd.<sup>11</sup> Lewis-basic heteroarylborons such as pyridines are particularly problematic for Ni, possibly due to their coordinating ability.<sup>12–14</sup> Furthermore, aryl BPins were the least reactive of the common SMC nucleophiles examined in a study by Percec.<sup>15</sup>

Complexes of other nonprecious metals besides Ni have also been reported as catalysts for the SMC. Fe<sup>16,17</sup> and Co-

### Selected ligands, catalysts, and precursors for the Nickel Suzuki–Miyaura Coupling (Ni-SMC)



### State of the art Ni-SMC of Lewis basic arylboronic acid and heteroaryl halides (Stradiotto, 2018)



### This research: efficient Ni-SMC of Lewis-basic aryl-BPins for biheteroaryl synthesis



<sup>a</sup>Process Research and Development, AbbVie Inc., North Chicago, Illinois, 60064, USA.  
E-mail: michael.haibach@abbvie.com

<sup>b</sup>Pfizer Chemical Research and Development, Pfizer Inc., Groton, Connecticut 06340, USA

† Electronic supplementary information (ESI) available. CCDC 2109558 and 2109557. For ESI and crystallographic data in CIF or other electronic format see DOI: <https://doi.org/10.1039/d2sc03877c>



catalyzed<sup>18,19</sup> couplings of arylboronic esters with alkyl halides have recently been developed. Co catalysts have also been reported for the coupling of arylboronic esters with aryl halides.<sup>20–23</sup> These examples rely on strong bases (alkyllithiums, lithium amides, or lithium alkoxides) to activate the arylboron reagent, limiting their application with base or nucleophile-sensitive functional groups. An interesting exception to the strong base requirement is the Cu-SMC reported by Giri, which can couple arylboronic esters in the presence of CsF at 120 °C.<sup>24</sup> In the Fe, Co, and Cu systems, heterocyclic or Lewis-basic arylboron nucleophiles are also not well-precedented.

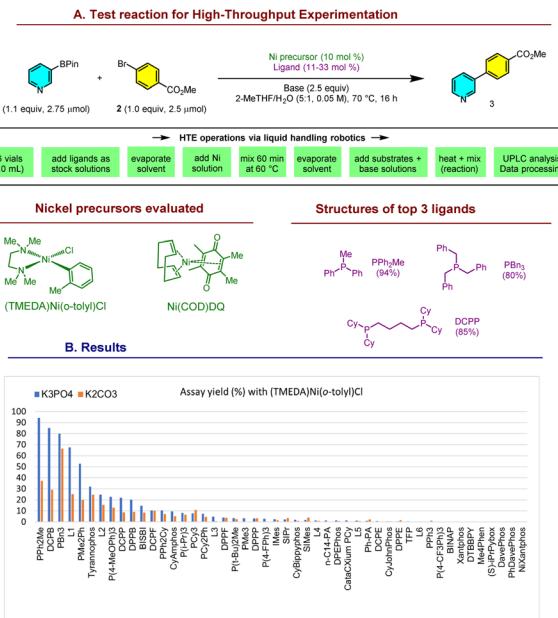
For these reasons, nonprecious metal catalyzed analogues of the Pd-SMC remain mostly limited to the preparation of biaryl compounds with limited functional groups. The Ni-SMC has rarely been used to prepare heterobiaryls, which are highly relevant in the synthesis of bioactive molecules.<sup>6</sup> Given our interest in more cost-effective and sustainable catalysts for cross-coupling reactions,<sup>25–27</sup> the authors at AbbVie and Pfizer desired to develop a general and economical Ni-SMC that addressed these drawbacks. Here we report the identification of a simple inexpensive ligand,  $\text{PPh}_2\text{Me}$ , that enables the Ni-SMC of an unprecedented range of heterocyclic and Lewis-basic BPins and trifluoroborate salts. The reaction conditions tolerate base-sensitive functional groups such as methyl esters, arylsulfonamides, and unprotected indoles. For several substrates, we observe catalyst performance on par with the best reported Pd catalysts. The origin of the high activity of this catalytic system was also investigated by experimental and computational approaches.

## Results and discussion

### Catalyst optimization

At the outset of our research, Ni-SMCs of the important pyridine-3-BPin **1** were unknown with (hetero)aryl halides, and had only been reported in a few cases with specialized electrophiles including vinyl sulfones and amides.<sup>28–30</sup> The absence of successful precedents and the limited mechanistic understanding of the Ni-SMC of pyridylborons make rational ligand design challenging. Hence, we turned to high-throughput experimentation (HTE)<sup>31–33</sup> to identify potential leads for a model reaction (Scheme 2).

Screening the coupling of **1** with methyl 4-bromobenzoate **2** against a bespoke library of 48 phosphine, nitrogen, and NHC ligands with  $(\text{TMEDA})\text{Ni}(\text{o-tolyl})\text{Cl}$ <sup>34,35</sup> as the Ni precursor afforded promising leads:  $\text{PBn}_3$  (80% yield), DCPB<sup>36</sup> (85%) and  $\text{PPh}_2\text{Me}$  (94%). Other interesting trends were apparent from the data. Reactions with  $\text{K}_3\text{PO}_4$  were typically more effective than  $\text{K}_2\text{CO}_3$ , although this difference was smaller with  $\text{PBn}_3$ . Commonly applied ligands for the Ni-SMC were ineffective for this coupling, highlighting the challenge posed by Lewis-basic arylborons such as **1** for Ni: DPPF (<5%), DPEPhos (<5%),  $\text{PCy}_3$  (11%), and  $\text{PPh}_3$  (<5%). The same screen was also conducted with  $\text{Ni}(\text{COD})\text{DQ}$ <sup>37</sup> as the Ni precursor, however this resulted in much lower assay yields (0–17%). We selected the highest yielding, readily available  $\text{PPh}_2\text{Me}$  for further development.



tolyl)Cl and (TMEDA)Ni(*o*-tolyl)Cl are commercially available, they are currently more costly than common Pd sources such as Pd(OAc)<sub>2</sub> and Pd<sub>2</sub>(dba)<sub>3</sub>. Hence, we pursued the inexpensive and easily prepared (PPh<sub>2</sub>Me)<sub>2</sub>NiCl<sub>2</sub> (**Ni-1**) as a catalyst precursor. The catalyst generated from **Ni-1**, PPh<sub>2</sub>Me and *n*-BuMgCl as an activator afforded 95% conversion of **2** to **3** after 16 h (entry 1). Omitting the *in situ* activation leads to lower conversion and the observation of significant amounts of byproducts (entries 3–4). Zn and *n*-BuLi could also be employed as *in situ* activators<sup>40–42</sup> of **Ni-1**, though conversion was slightly lower than with *n*-BuMgCl (entries 5–6).

Many reported Ni-SMC methods are either sensitive to adventitious moisture or require careful control of H<sub>2</sub>O stoichiometry. Gratifyingly, the reaction proceeded efficiently with no added H<sub>2</sub>O through 4 : 1 v/v ratio of 2-MeTHF/H<sub>2</sub>O (entries 7–9). We chose 5 : 1 v/v 2-MeTHF/H<sub>2</sub>O for subsequent evaluation due to the slightly higher conversion obtained at this ratio. The conditions with less H<sub>2</sub>O may be useful for specific applications (*i.e.*, a water-sensitive substrate). Dioxane and *t*-AmOH were also effective solvents (entries 10 and 11). Weaker bases K<sub>2</sub>CO<sub>3</sub> and K<sub>2</sub>HPO<sub>4</sub> were less effective than K<sub>3</sub>PO<sub>4</sub> (entries 13 and 14).

### Scope of the reaction

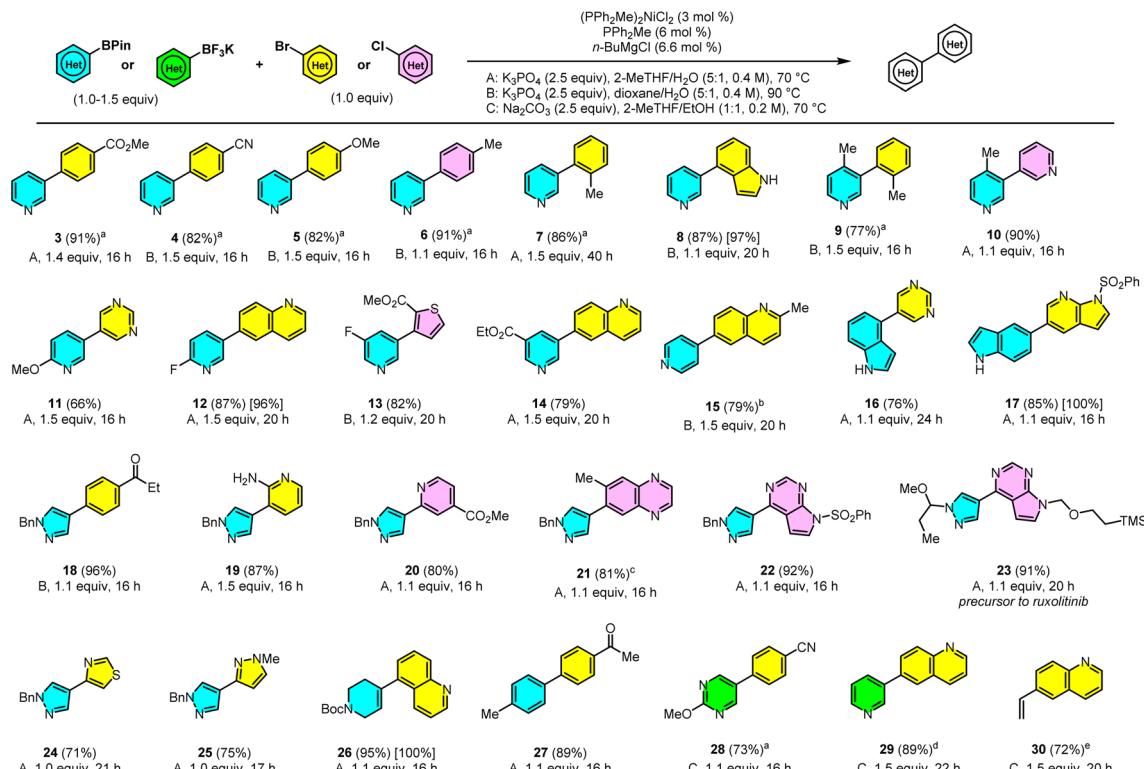
With optimized conditions in hand, we explored the scope of the reaction catalyzed by 3 mol% **Ni-1** (Scheme 3). The coupling of **1** proceeds efficiently with a range of aryl halides at 70 °C (in

2-MeTHF, bp = 80 °C) or 90 °C (in dioxane, bp = 101 °C). Electron-poor (**3** and **4**), electron-rich (**5** and **6**), and *ortho*-substituted (**7** and **9**) aryl bromides afforded the coupled products in excellent yields. Impressively, an electronically neutral aryl chloride (**6**), an unprotected bromoindole (**8**), and Lewis-basic 3-chloropyridine (**10**) also coupled in high yield.

In addition to the electron-rich pyridine-3-BPin (**11**), the previously unreported electron-poor pyridine-3-BPins coupled with heteroaryl halides (**12**–**14**). Even the less nucleophilic pyridine-4-BPin, an unknown substrate for the Ni-SMC and a challenging one even for the Pd-SMC, coupled efficiently at 90 °C using 5 mol% catalyst (**15**). A functionalized chlorothiophene was well tolerated as an electrophile (**13**).

Unprotected indoles are another class of heterocycles previously unreported in the Ni-SMC and were shown to be problematic in Watson's study of DPPF-Ni.<sup>11</sup> Examples **8**, **16** and **17** show the efficient coupling of unprotected indoles as both nucleophile and electrophile components. Compound **17** also demonstrates the viability of a derivatized 7-azaindole as an electrophile, a motif found in several pharmaceuticals.<sup>43</sup> Methyl/ethyl esters (**3**, **13**, **14**, and **20**), 2-fluoropyridine (**12**) and heterocyclic sulfonamides (**17** and **22**) were well-tolerated, due to the mildly basic conditions of the reaction.

Five-membered Lewis-basic arylborons represent another important class of nucleophiles that is typically difficult for the Ni-SMC. The Ni-SMC of (hetero)aryl halides with pyrazoleborons was previously unreported. Examples **18**–**25** show the



**Scheme 3** Scope of the Suzuki–Miyaura coupling catalyzed by (PPh<sub>2</sub>Me)<sub>2</sub>NiCl<sub>2</sub> (**Ni-1**). Reactions were run on 1.0 mmol scale unless noted, isolated yield in parentheses. Equiv. given in scheme refers to the arylboron reagent. The yield in brackets represents assay yield of reaction with 1 mol% Ni on 1 g scale. (a) Reaction run on 1.0 g scale (b) 5 mol% (PPh<sub>2</sub>Me)<sub>2</sub>NiCl<sub>2</sub>/10 mol% PPh<sub>2</sub>Me/11 mol% *n*-BuMgCl (c) 3 mol% (TMEDA)Ni(*o*-tolyl)Cl/9.9 mol% PPh<sub>2</sub>Me as catalyst (d) 90 °C (e) coupling partner was potassium vinyltrifluoroborate (1.5 equiv.).



coupling of a pyrazole-4-BPin with functionalized heteroaryl halides at 70 °C, including an unprotected aminopyridine (**18**). The pyrazolodiazaindole core of several classes of kinase inhibitors was efficiently generated in **22** and **23**.<sup>44,45</sup> Diazaindole **22** is an intermediate in an enantioselective synthesis of ruxolitinib reported by Lin and coworkers at Incyte.<sup>46</sup> Examples **24** and **25** are notable examples of coupling two five-membered rings, which is typically more challenging than 6-membered rings for both Pd and Ni.<sup>47,48</sup> An *N*-Boc containing vinyl BPin coupled to form **26** in excellent yield at 70 °C. This type of Suzuki coupling is used extensively in pharma to access 4-arylpiperidine derivatives yet was previously unknown with Ni.<sup>49</sup> Heteroaryl or Lewis-basic substrates are not required for the catalyst to operate efficiently, as shown by the coupling to form **27**.

Excitingly, **Ni-1** could also efficiently couple potassium organotrifluoroborates under slightly modified conditions. Both Lewis-basic heteroaryl and vinyl  $\text{BF}_3\text{K}$  derivatives coupled efficiently at 70–90 °C in the presence of  $\text{Na}_2\text{CO}_3$ /EtOH/2-MeTHF<sup>50</sup> (**26–28**). These examples further highlight the flexibility of this catalytic system to proceed under mild conditions.

While the scope of the reaction was explored with 3 mol% **Ni-1**, examples **8**, **12**, **17** and **26** were also conducted on gram scale with 1 mol% **Ni-1**/2 mol%  $\text{PPh}_2\text{Me}$ /2.2 mol% *n*-BuMgCl. The reactions afforded excellent assay yields in 16–24 h without any modification of the general reaction conditions, demonstrating the amenability of this system to large-scale applications. For general couplings of the arylboron nucleophile in **24**, the lowest reported catalyst loading with Pd was 1 mol%.<sup>51,52</sup> In the case of the arylboron in **11**, it was 2 mol% with Pd.<sup>53</sup> Thus, our Ni catalyst can achieve efficiency on par with Pd catalysts for these more challenging substrates! Overall, the scope of the Ni-SMC

catalyzed by **Ni-1** shows broad Lewis base and functional group tolerance, while in many cases requiring only 1.1 equiv. of the aryl-BPin to afford high yield.

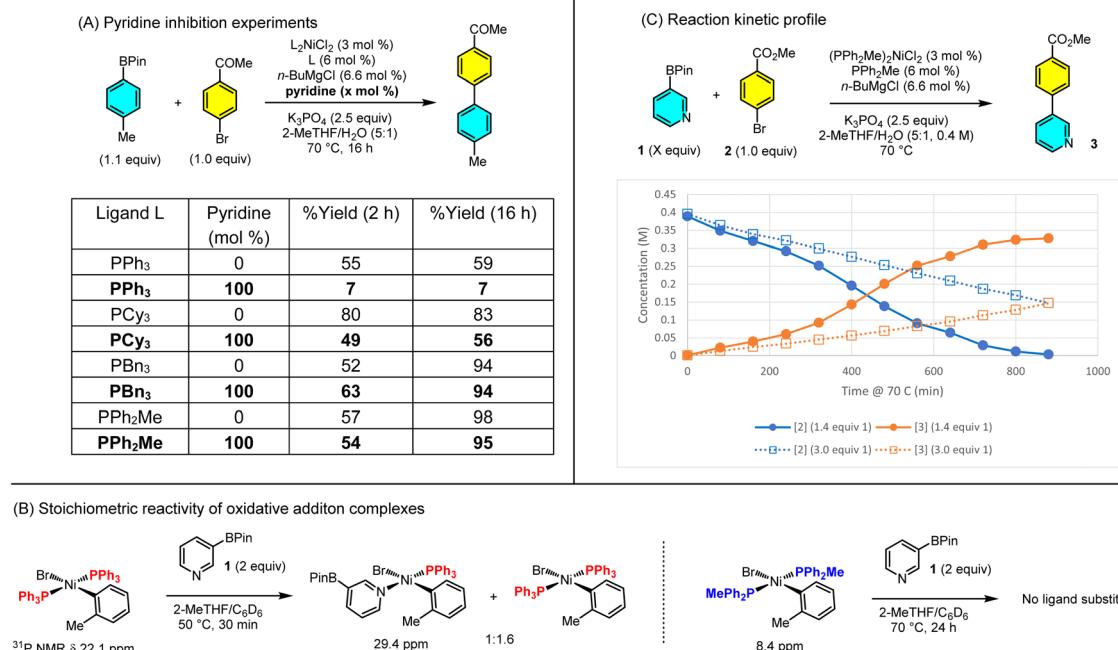
### Characterization of the activated precatalyst

As noted earlier, **Ni-1**/ $\text{PPh}_2\text{Me}$ /*n*-BuMgCl showed activity comparable to  $(\text{PPh}_2\text{Me})_2\text{Ni}(\text{o-tolyl})\text{Cl}$  or  $\text{PPh}_2\text{Me}/(\text{TMEDA})\text{Ni}(\text{o-tolyl})\text{Cl}$  in our optimization studies. NMR and X-ray analysis of **Ni-1**/ $\text{PPh}_2\text{Me}$ /*n*-BuMgCl confirmed clean and rapid formation of  $(\text{PPh}_2\text{Me})_4\text{Ni}$ , a compound known for over 50 years<sup>54</sup> but with few prior catalytic applications.<sup>55–57</sup> Isolated  $(\text{PPh}_2\text{Me})_4\text{Ni}$  was found to be a competent precatalyst in the Ni-SMC but had limited stability on the bench, thus *in situ* generation from **Ni-1** is preferred.

### Mechanistic studies

We were intrigued by the high activity afforded by the relatively simple  $\text{PPh}_2\text{Me}$  ligand in the Ni-SMC of Lewis-basic arylboron esters, a reaction where many well-established ligands for Ni did not perform well (see Scheme 2). The most obvious challenge posed by the coupling of substrates such as **1** is their ability to coordinate to Ni, potentially displacing phosphine ligands. Indeed phosphine substitution by Lewis basic amines, including pyridine, was studied by Mizoroki and Nakamura using the related  $(\text{PPh}_3)_2\text{Ni}(\text{o-tolyl})\text{Br}$  complex over forty years ago.<sup>58–60</sup> Given this precedent, we wondered if the unique ability of  $(\text{PPh}_2\text{Me})_2\text{NiCl}_2$  in the SMC stems from a reduced tendency of the  $\text{PPh}_2\text{Me}$  ligand to be substituted by Lewis basic donors.

To test this hypothesis, we first investigated the non-heterocyclic Ni-SMC shown in Scheme 4A. With  $\text{PCy}_3$ ,  $\text{PPh}_3$ ,  $\text{PPh}_2\text{Me}$ , or  $\text{PBn}_3$ , the reaction proceeded to 59–98% assay yield



Scheme 4 Mechanistic experiments on the role of Lewis bases in the Ni-SMC. <sup>a</sup>See ESI† for details.



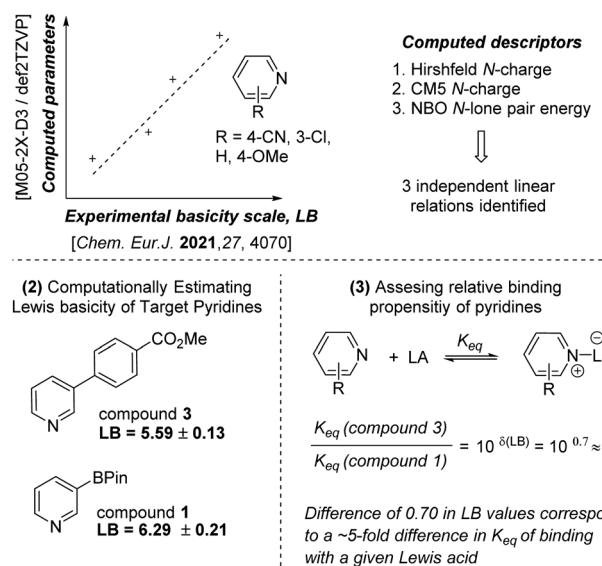
after 16 h. When 100 mol% pyridine was added, however, the reactions catalyzed by  $\text{PCy}_3$  and  $\text{PPh}_3$  (inactive with **1**) were significantly inhibited, as expected, while  $\text{PPh}_2\text{Me}$  and  $\text{PBn}_3$  (active with **1**) were unaffected.<sup>61</sup> While consistent with our hypothesis that catalyst derived from  $(\text{PPh}_2\text{Me})_2\text{NiCl}_2$  could be resistant to phosphine displacement, we explored this in more detail through  $^{31}\text{P}$  NMR studies.

Thus we prepared oxidative addition complexes  $(\text{PPh}_2\text{Me})_2\text{Ni}(o\text{-tolyl})\text{Br}$  and  $(\text{PPh}_3)_2\text{Ni}(o\text{-tolyl})\text{Br}$  to probe the extent of phosphine substitution. We chose these complexes as (1) the ortho substituent renders them isolable as well as air-stable, and (2) 2-bromotoluene is a catalytically relevant aryl halide with  $\text{PPh}_2\text{Me}$  (Scheme 3, compounds **7** and **9**). Complex  $(\text{PPh}_3)_2\text{Ni}(o\text{-tolyl})\text{Br}$  reacted with 2 equiv. **1** at 50 °C to generate a new downfield species in  $^{31}\text{P}$  NMR that is consistent with previously reported  $(\text{PPh}_3)(\text{pyridine})\text{Ni}(o\text{-tolyl})\text{Br}$  complexes (Scheme 4B).<sup>60</sup> By contrast, heating  $(\text{PPh}_2\text{Me})_2\text{Ni}(o\text{-tolyl})\text{Br}$  with 2 equiv. **1** at 70 °C for 24 h produced no new  $^{31}\text{P}$  NMR peaks. Hence, substitution of  $\text{PPh}_2\text{Me}$  by pyridines is disfavored relative to  $\text{PPh}_3$ . The extent of substitution of  $\text{PPh}_3$  is significant even with only 2 equiv. of **1** relative to the Ni complex. Under typical catalytic conditions (37–47 equiv. **1** per Ni) the substitution would be expected to proceed much further.<sup>62</sup> Taken together, these experiments show that addition of a Lewis basic donor such as **1** or pyridine leads to significant substitution of the phosphine ligands, which could lead to off-cycle species and thus slow or inhibit the desired SMC.

In Mizoroki and Nakamura's ligand substitution studies, sterics were the primary determinant of substitution equilibrium between  $\text{PPh}_3$  and other ligands. Less-hindered amines and smaller cone angle phosphines displaced  $\text{PPh}_3$  to a greater extent than larger or more hindered ones. When steric effects were held constant, however, higher basicity of the incoming ligand led to greater displacement of  $\text{PPh}_3$ . Thus, under typical reaction conditions for the SMC, we should expect a potentially different degree of substitution of the phosphine ligand as  $\text{BPin}$  **1** is converted into the biaryl product **3**. Indeed, the Lewis basicity of **1** and **3**, estimated using computed descriptors and experimental values for related pyridines (Scheme 5) show that **1** is 0.70 units more Lewis basic than **3**, which corresponds to a 5-fold greater  $K_{\text{eq}}$  in coordinating to a Lewis acid. If correct, we would expect the reaction to show a sigmoidal reaction profile consistent with conversion-dependent relief of inhibition. Thus, we profiled the coupling of **1** and **2** by **Ni-1** over time (Scheme 4C) and did observe the expected conversion profile. When the same reaction was conducted with a greater excess **1** (3.0 equiv. vs. 1.4 equiv.), the reaction showed overall zero-order behavior over the first 16 h, further evidence of inhibition by **1**.<sup>63</sup> An aliquot of the reaction with 1.4 equiv. **1** at 80 min (5.5% yield) was analyzed by  $^{31}\text{P}$  NMR, and showed 96% uncoordinated  $\text{PPh}_2\text{Me}$ , 1%  $(\text{PPh}_2\text{Me})_4\text{Ni}$ , and 3% of a  $\text{PPh}_2\text{Me}$ –Ni complex.<sup>64</sup>

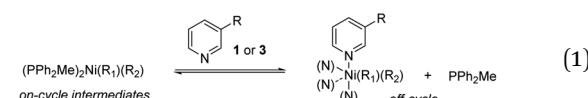
The evidence described above implies that the Lewis basic Ni-SMC is governed by substantial pre-equilibrium steps involving ligand substitution between  $\text{PPh}_2\text{Me}$ , **1** and **3** (eqn (1)). While under catalytic conditions ( $[\mathbf{1}] > [\text{PPh}_2\text{Me}]$ ) most of the equilibrium favors the off-cycle species, enough of the on-

### (1) Mapping Experimental Lewis Basicity Scale on Computed Parameters



Scheme 5 Summary of the procedure used to estimate computed Lewis basicities of pyridines **1** and **3**. See ESI† for computational details, calibration of linear models and thermodynamic data.

cycle intermediates are retained with  $\text{PPh}_2\text{Me}$  to allow the reaction to proceed efficiently. Indeed, our experiments show that the commonly employed ligands  $\text{PPh}_3$  and  $\text{PCy}_3$  are more susceptible to substitution by pyridines than  $\text{PPh}_2\text{Me}$  and  $\text{PBn}_3$ . Mizoroki and Nakamura's studies imply that this is primarily a steric effect.



Because they are minor species in eqn (1), the  $\text{PPh}_2\text{Me}$ –Ni intermediates must also be able to promote very efficient on-cycle steps, including the potentially difficult<sup>65,66</sup> trans-metallation. One implication of this result for future research is to avoid using a large excess of Lewis-basic arylboron reagents when screening Ni-SMCs.

## Conclusions

We have developed an effective and inexpensive catalyst system, *in situ* activated  $(\text{PPh}_2\text{Me})_2\text{NiCl}_2$ , for the Suzuki–Miyaura coupling of Lewis basic heteroaryl BPins and  $\text{BF}_3\text{Ks}$  with (hetero)aryl bromides and chlorides. The scope and functional group tolerance of the reaction allow important Lewis-basic and biheteroaryl compounds to be prepared with a nonprecious metal catalyst for the first time. Ligand substitution of  $\text{PPh}_2\text{Me}$  by Lewis basic heterocycles is disfavored relative to more commonly studied phosphines, a key reason for its unique catalytic activity. Future research on the precise balance of steric and electronic factors that contribute to this property should

lead to the development of even more highly active ligands for Ni-catalyzed cross couplings of Lewis-basic substrates.

## Author contributions

M. C. H. conceived and directed the study with contributions from S. S. and S. M. M. C. H., A. R. I., and S. S. conducted synthetic experiments. S. T. conducted the computational study. R. S., B. J. K., and J. W. conducted high-throughput experiments. A. L. W. and R. F. H. conducted HRMS and X-ray crystallography, respectively. M. C. H. and S. T. wrote the final draft manuscript. All authors contributed to the data interpretation, manuscript review, and have approved of the final version of the manuscript.

## Conflicts of interest

AbbVie contributed to the design, approval, and execution of this study. M. C. H., A. R. I., S. S., R. S., B. J. K., J. W., A. L. W., and R. F. H. are current or former AbbVie employees and may own AbbVie stocks.

## Acknowledgements

The authors thank the participants in the Pharma Alliance at AbbVie, Pfizer and Boehringer Ingelheim for their helpful discussions.

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63 The analogous reaction of **2** with 1.4 equiv. PhBPin showed first-order dependence on **2**, pointing to a different rate limiting step for the Ni-SMC of non-Lewis basic arylborons. See the ESI† for details.

64 By contrast, <sup>31</sup>P NMR of the coupling of PhBPin and **2** showed the expected mixture of PPh<sub>2</sub>Me-Ni complexes (52–85%) and free PPh<sub>2</sub>Me (48–15%) depending on the reaction progress. See the ESI† for details.

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