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Unsymmetric N-heterocyclic carbene ligand enabled nickel-catalysed arylation of bulky primary and secondary amines†

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The arylation of sterically hindered amines represents one of the long-standing challenges in synthetic chemistry. Herein, we report a highly efficient Ni-catalysed arylation of sterically hindered primary and secondary amines with aryl chlorides or phenol derivatives enabled by an unsymmetric N-heterocyclic carbene (NHC) ligand. The protocol provides general, efficient, and scalable access to various sterically demanding anilines in excellent yields under mild conditions. A wide range of functional groups and heterocycles are compatible (>50 examples), including those present in biologically relevant molecules. Computational studies suggest that the unsymmetric bulky and flexible NHC ligand was critical to balance the oxidative addition and reductive elimination elementary steps, thus promoting this challenging transformation.

Introduction

Transition-metal-catalysed C–N cross-coupling reactions of amines and aryl halides have become some of the most frequently employed and significant transformations in academic research and industrial processes. These amination methods have enabled mild and general access to various *N*-aryl structures ubiquitous in drugs, materials science, and natural products (Fig. 1A).^{1–4} Despite tremendous advances, transition-metal-catalysed arylation of sterically hindered amines remains a challenging but important task. Such transformations are attractive because of the prevalence of bulky anilines in medical agents and their applicability in drug design (Fig. 1A), as sterically demanding substituents can increase a bioactive molecule's lipophilicity or improve its stability against enzymatic degradation.⁵

Despite these attractive features of the coupling of bulky amines, specific factors have impeded the development of such reactions (Fig. 1B). The main challenges stem from the intrinsic

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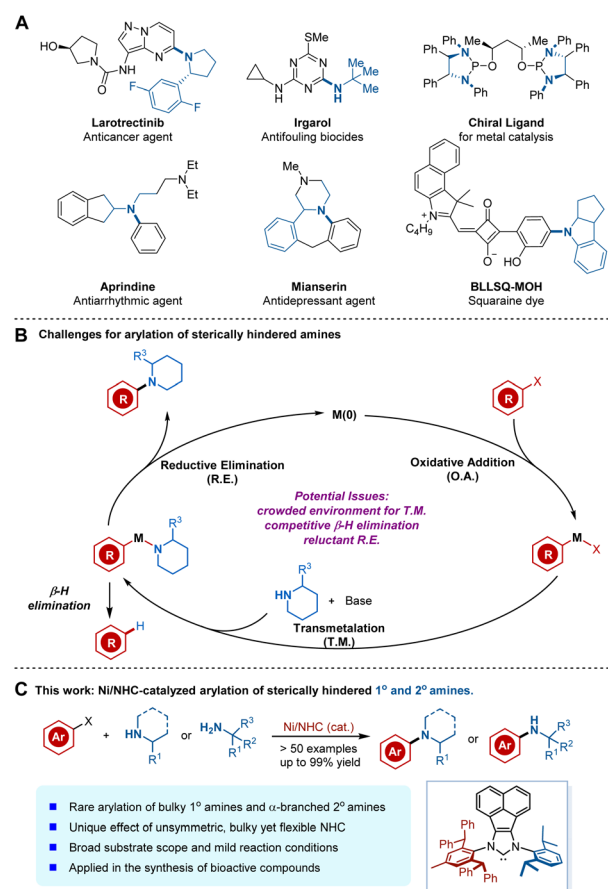


Fig. 1 (A) Examples of pharmaceuticals, ligands, and materials possessing bulky aniline motifs. (B) Potential challenges for the arylation of sterically hindered amines. (C) This work.



poor nucleophilicity of bulky amines as a consequence of steric hindrance, which results in a slower rate of coordination and transmetalation (T. M.) of bulky amines to metals.⁶ Added to this complication is the β -hydride elimination process encountered in most coupling reactions.⁷ In this regard, β -hydride eliminations from the metal-amido complex intermediate often cause the undesired reduced arene as a side product. Accordingly, such challenging yet crucial reactions have attracted chemists to search for more efficient catalytic systems. Nevertheless, very few examples of the Pd or Cu-mediated arylation of bulky amines have been reported, considering the numerous known methods for C–N cross-couplings (Fig. 1B).^{6,8} In these sporadic examples, the design and modification of the ligand architecture represent a fundamental strategy to overcome the key limitations and achieve more efficient cross-coupling.

Recently, more and more attention has been focused on nickel as a catalyst for C–N cross-coupling reactions,^{3,9} not only due to the low cost of the earth-abundant metal but also because of its unique reactivity towards aryl chloride and phenol derivatives.¹⁰ Despite these advantages, considerable limitations and challenges remain in Ni-catalysed arylation of hindered amines. In addition to challenges associated with transmetalation, Ni-catalysed *N*-arylation also suffers from more reluctant reductive elimination (R. E.) of nickel(II) amido complexes.¹¹ In addition, ancillary ligands developed for Pd catalysts are not fully applicable for Ni catalysts. Therefore, the rational ligand design or strategical application of privileged ligands for Ni-catalysed *N*-arylation is non-trivial considering the infancy of ligand design for Ni catalysis and the relative lack of general principles.^{12,13} As a consequence, efficient Ni-catalysed arylation of bulky α -branched secondary amines has not been explored, presumably due to the more difficult in balancing the elementary steps and potential deleterious β -H elimination process.

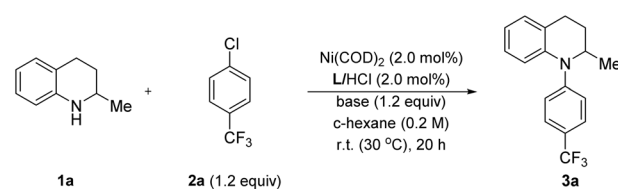
We recently developed an array of bulky yet flexible NHCs¹⁴ and applied them to a range of Ni-catalysed transformations, including challenging cross-coupling reactions.¹⁵ We felt that a sterically demanding NHC ligand could promote R. E. At the same time, the conformational flexibility, available through rotatable single bonds on the ligand, may benefit coordination to the metal center, as well as oxidative addition (O. A.) and T. M. processes. As part of our ongoing endeavors in this field, we herein report a highly efficient Ni-catalysed arylation of sterically demanding primary amines and bulky α -branched secondary amines enabled by an unsymmetric bulky yet flexible NHC ligand (Fig. 1C). Our protocol is advantageous in terms of its broad substrate scope, cost effectiveness, simple and mild reaction conditions, and the applicability of aryl chlorides or phenol derivatives when compared to previous Pd- or Cu-based methods.^{6,8} The unique ligand effect of the unsymmetric bulky yet flexible NHC on balancing the elementary catalytic steps was first observed and confirmed by DFT studies, which would inspire further rational ligand design.

Results and discussion

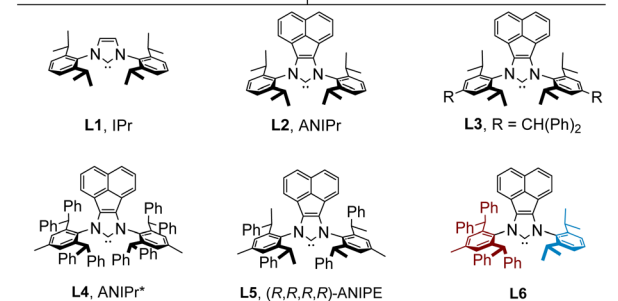
We began our study by treating commercially available 1,2,3,4-tetrahydroquinoline (**1a**) with 4-chlorobenzotrifluoride (**2a**) in the

presence of 2.0 mol% Ni(COD)₂, 1.2 equiv. *t*-BuOK, and 2.0 mol% imidazolium salts in cyclohexane at room temperature. To our surprise, the use of IPr (**L1**) and other typical ligands for C–N cross-coupling^{3a} failed to afford the desired product **3a** (Table 1, entry 1, also see the ESI† for more details). Instead, we observed significant amounts of imine by-product generated from the dehydrogenation of **1a** probably through β -H elimination. Interestingly, the bulkier NHC ligands with acenaphthoimidazolyliene skeletons (**L2**, ANIPr)¹⁶ or aniline moiety containing benzhydryl groups (**L3**)¹⁷ delivered **3a** in 26% and 24% yields, respectively (entries 2 and 3). However, a reaction using highly bulky ANIPr* (**L4**)¹⁸ resulted in no conversion to the desired product (entry 4). We felt that a suitable sterically hindered but flexible ligand would facilitate this challenging amination process. We thus used our ANIPE ligand (**L5**) to promote the C–N coupling reaction. Indeed, the yield was improved to 58% (entry 5). However, we found that further improvement in reactivity was difficult since only one of the enantiomers of substrate **1a** could react efficiently as a result of a highly selective kinetic resolution by the chiral ligand.^{15b} An achiral unsymmetric NHC (**L6**) with 2,6-diphenethylaniline and 2,6-diisopropylaniline fragments¹⁹ was then utilized as the ligand. With this change, **3a** was formed in a high yield (70%; entry 6). We attribute the low reactivity to inefficient R. E. in **L1**–**L3** and ineffective O. A., T. M., or coordination to nickel for **L4**, while the unsymmetrical NHC (**L6**) seems to strike an excellent balance in

Table 1 Reaction optimization^a

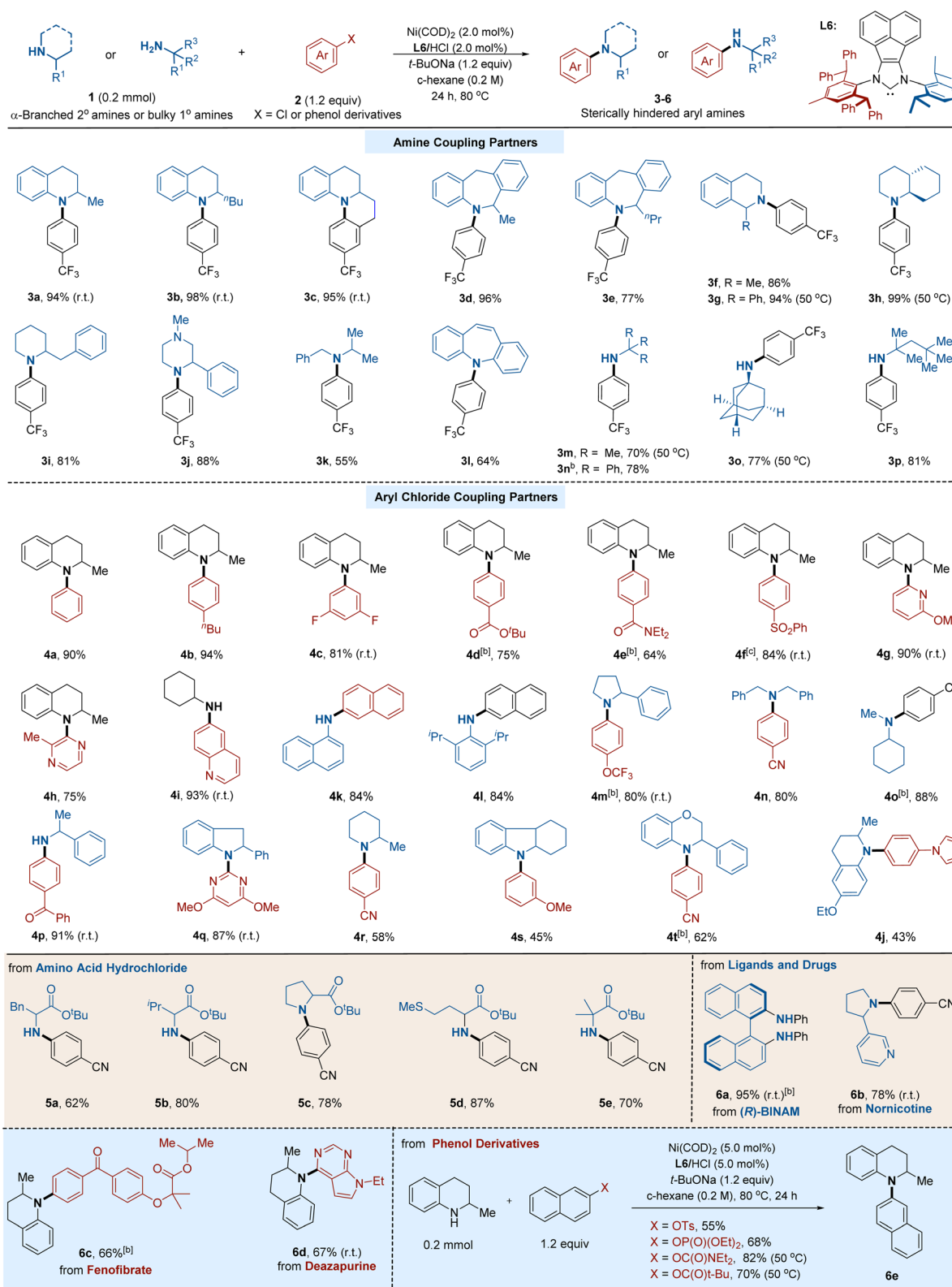


entry	ligand	base	yield (%) ^b	entry	ligand	base	yield (%) ^b
1	L1	<i>t</i> -BuOK	0	8	L6	K ₃ PO ₄ ^d	0
2	L2	<i>t</i> -BuOK	26	9	L6	DBU ^d	0
3	L3	<i>t</i> -BuOK	24	10	L6	Et ₃ N ^d	0
4	L4	<i>t</i> -BuOK	0	11	L6	<i>t</i> -BuOLi	37
5	L5	<i>t</i> -BuOK	58 (44) ^c	12	L6	<i>t</i> -BuONa	96
6	L6	<i>t</i> -BuOK	70	13 ^e	L6	<i>t</i> -BuONa	0
7	L6	CsF ^c	<2	14 ^f	--	<i>t</i> -BuONa	0



^a Reactions were performed on a 0.2 mmol scale. ^b Determined by NMR analysis using 1,3,5-trimethyl-benzene as an internal standard. ^c Using *t*-BuONa instead of *t*-BuOK. ^d 5.0 mol% *t*-BuOK was added. ^e Without Ni(COD)₂. ^f Without **L6**.



Table 2 Substrate Scope^a

^a Yields of isolated products on a 0.2 mmol scale are reported. ^b 5.0 mol% Ni(COD)₂ and L6/HCl were used. ^c Cyclohexane (0.1 M).



having favorable steric environments for different stages of the catalytic cycle (*vide infra*).

To further improve the catalytic efficiency, the effect of the base was then studied (entries 7–12). Organic bases, such as Et₃N and DBU,²⁰ were found to be ineffective in this transformation. The use of weak inorganic bases, including CsF and K₃PO₄, delivered little or no **3a**. To our delight, an excellent yield (96%) was obtained when *t*-BuONa was used (entry 12), in part because of better solubility in cyclohexane compared to *t*-BuOK (see the ESI† for solvent optimization details). As expected, control experiments revealed that Ni(COD)₂ and NHC ligands (entries 13 and 14) were both essential for the C–N cross-coupling reaction.

With the optimized reaction conditions, we focused our attention on the substrate scope of this transformation with various hindered primary and secondary amines. As depicted in Table 2, our arylation method showed an excellent efficiency profile. A diverse array of C–N cross-coupling products were prepared in good to excellent yields. A series of bulky cyclic secondary amines, including tetrahydroquinoline (**3a** and **3b**), tetrahydroisoquinoline (**3f** and **3g**), decahydroquinoline (**3h**), piperidine (**3i**), and piperazine (**3j**), all served as competent substrates. Moreover, seven-membered cyclic amines (**3d** and **3e**) could also be arylated efficiently. In addition, acyclic hindered secondary amines (**3k**)^{6b} and diarylamine (**3l**),²¹ which are non-trivial in Ni-catalyzed cross-coupling reactions due to their steric hindrance, gave the corresponding products in good yields. Notably, the arylation protocol was efficient for a wide variety of bulky α,α,α -trisubstituted primary amines, such as *tert*-butylamine (**3m**), triphenylmethylamine (**3n**), amantadine (**3o**), and *tert*-octylamine (**3p**).¹³ It is worth mentioning that room temperature was sufficient to carry on the reaction in some cases, highlighting the high efficiency of our protocol.

Encouraged by the above outcomes, we moved on to explore the scope and reactivity of aryl chlorides. As shown in Table 2, a series of aryl chlorides with electron-deficient, electron-neutral, and electron-rich substituents on the aryl group are suitable substrates, affording the corresponding products in good to excellent yields. Diverse functional groups, such as fluoride (**4c**), ester (**4d**), amide (**4e**), sulfonyl (**4f**), ether (**4g** and **4s**), cyano (**4o**), and trifluoromethoxy group (**4m**), could be well accommodated under the reaction conditions. Notably, an *ortho*-substituted aryl chloride was applicable (**4h**), which is challenging in metal-catalyzed *N*-arylation. Cross-couplings using heteroaryl chlorides containing pyridine (**4g**), pyrazine (**4h**), quinolone (**4i**), pyrrole (**4j**), and pyrimidine (**4q**) also performed efficiently. Given the wide presence of aniline and the prevalence of heterocycles in pharmaceutically relevant compounds, the current C–N cross-coupling protocol would provide a robust tool for constructing the skeleton of the biologically active molecule.

With this reliable protocol for both primary and secondary amines, we next applied this method to modify a series of ligands and bioactive compounds. We first extended this reaction to various amino acid derivatives; either primary hindered amines or secondary congeners could be arylated in good yields. Furthermore, complex ligands and drug-relevant molecules,

including (*R*)-BINAM, nornicotine, fenofibrate, and deazapurine, readily undergo amination reactions, furnishing arylated products in high yields. Considering the unique reactivity of Ni catalysts towards phenol derivatives, we then turn our attention to these challenging coupling partners. A range of phenol-derived electrophiles, including aryl triflates, aryl phosphates, aryl carbamate, and aryl carbonate, which are ineffective for Ni-bisphosphine conditions,¹³ were found to be suitable substrates under the current conditions, giving moderate to high yields of aminated products.

To demonstrate the practicality of our reaction, we conducted a 5.0 mmol-scale reaction (Fig. 2A). On this scale, product **3a** was obtained in 93% yield (1.35 g). When a combination of the air-stable precatalyst NiCl₂·DME and PhBpin was used instead of Ni(COD)₂, product **3a** was obtained with similarly high efficiency (Fig. 2B). To further show the synthetic utility of our protocol, we exemplify three applications in the synthesis of drugs, herbicides, and advanced materials. First, this Ni/NHC-catalyzed process was applied to prepare CDP-352,

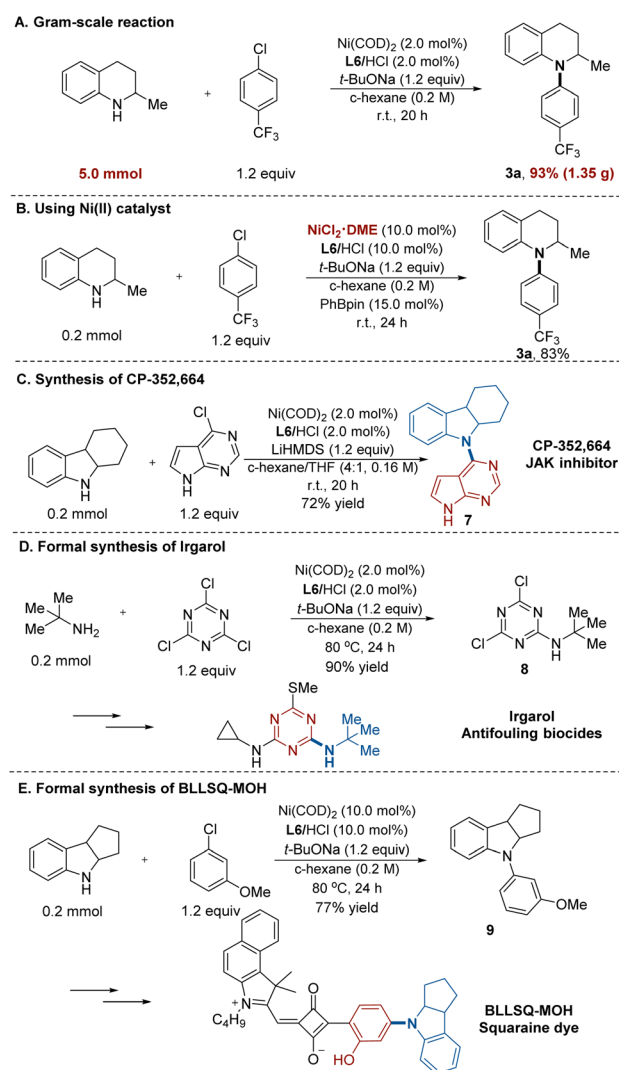


Fig. 2 Gram-scale reaction and synthetic applications.



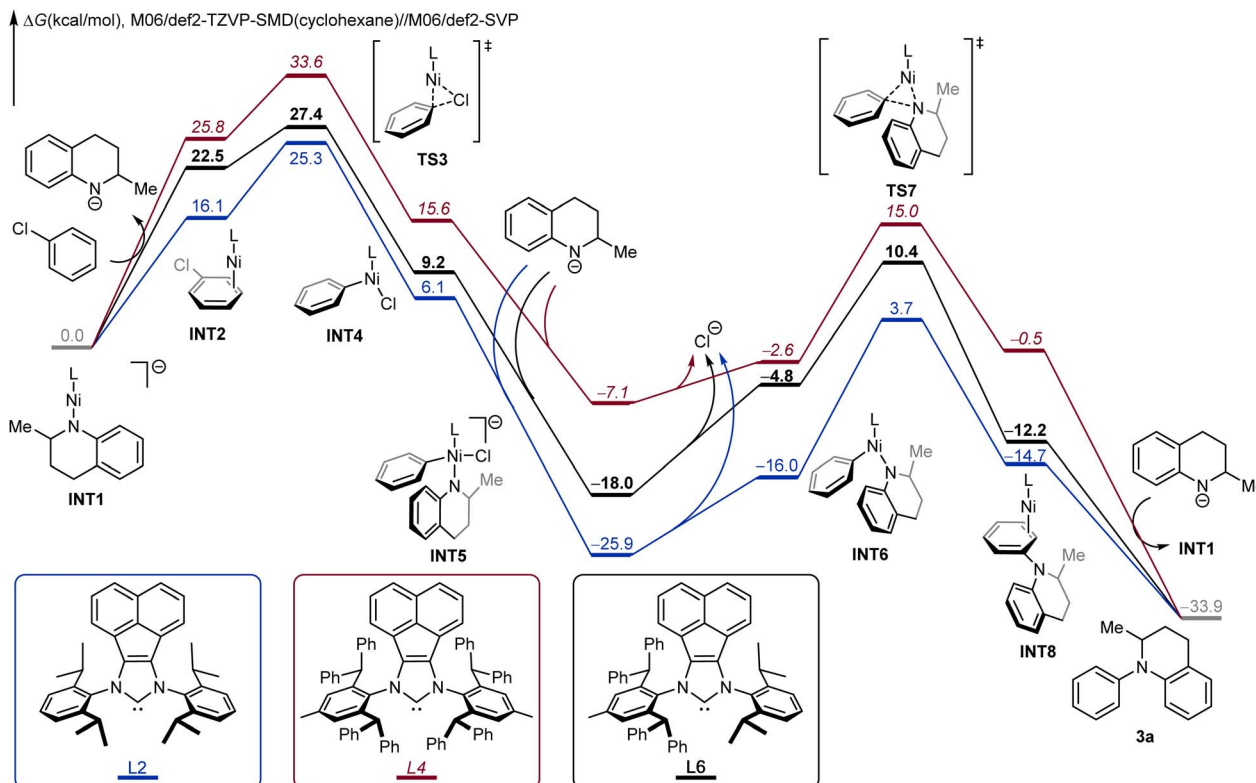


Fig. 3 DFT-computed Gibbs free profiles of Ni-catalyzed *N*-arylation between 1,2,3,4-tetrahydroquinoline and chlorobenzene under various NHC ligands.

664 (7),²² a potent JAK inhibitor. The desired product was furnished in a single transformation in 72% yield (Fig. 2C). Moreover, the intermediate for irgarol²³ (**8**) was prepared in 90% yield using *tert*-butylamine and 1,3,5-trichlorotriazine, without the formation of di- or triamination byproducts (Fig. 2D). In addition, we applied the amination protocol to synthesize the key intermediate (**9**) for BLSQ-MOH, a squaraine dye,²⁴ which was obtained in 77% yield (Fig. 2E).

We next performed density functional theory (DFT) calculations to investigate the origins of the high reactivity of the unsymmetric NHC ligand **L6**, using coupling between 1,2,3,4-tetrahydroquinoline and chlorobenzene as a model. The DFT-computed free energy profiles of the coupling catalytic cycle for **L2**, **L4** and **L6** are compared in Fig. 3. From the deprotonated tetrahydroquinoline anion-coordinated complex **INT1**, the ligand exchange with chlorobenzene leads to the substrate-coordinated intermediate **INT2**. The subsequent oxidative addition *via* **TS3** generates the $\text{LNi}(\text{phenyl})\text{Cl}$ intermediate **INT4**, which is further coordinated by the deprotonated tetrahydroquinoline anion to produce **INT5**. From **INT5**, the dissociation of the chloride anion leads to the neutral $\text{LNi}(\text{phenyl})(\text{amino})$ intermediate **INT6**. **INT6** undergoes three-membered ring C–N reductive elimination through **TS7** to produce the product-coordinated complex **INT8**. The subsequent product liberation regenerates the active intermediate **INT1** and releases the observed arylation product **3a**. For the optimal unsymmetric NHC ligand **L6** (black

pathway), the oxidative addition requires an overall barrier of $27.4 \text{ kcal mol}^{-1}$ from **INT1** to **TS3**. Due to the stable intermediate **INT5**, the C–N reductive elimination is rate-determining with an overall barrier of $28.4 \text{ kcal mol}^{-1}$.

Comparing the three free energy profiles, the computation well reproduces the trend of ligand reactivities. For **L6** (black pathway), the overall reaction barrier is $28.4 \text{ kcal mol}^{-1}$ with reductive elimination being the rate-limiting step. For the more sterically demanding ligand **L4** (red pathway), the additional steric repulsions promote reductive elimination; the C–N reductive elimination requires an overall barrier of $22.1 \text{ kcal mol}^{-1}$ from **INT5** to **TS7**. However, the same steric repulsions also disfavor the oxidative addition, and the barrier for **L4** increases to $33.6 \text{ kcal mol}^{-1}$ from **INT1** to **TS3**. For the less demanding ligand **L2** (blue pathway), the oxidative addition step requires a barrier of $25.3 \text{ kcal mol}^{-1}$, which is more efficient than **L4**; however, the reductive elimination barrier with **L2** increases to $29.6 \text{ kcal mol}^{-1}$, which is rate-determining for **L2** and is $1.2 \text{ kcal mol}^{-1}$ higher as compared to the overall reaction barrier of **L4**. Overall, our computations corroborated the reactivity trend of **L6** > **L2** > **L4**, which is in nice agreement with the experimental observations (Table 1). These calculations highlight the important role of tuning the steric repulsions by the unsymmetric NHC ligand, which is able to achieve the desired balance of the contrasting steric effects on oxidative addition and reductive elimination.



Conclusions

In conclusion, we have developed a highly efficient unprecedented Ni-catalysed *N*-arylation of sterically hindered primary and secondary amines with aryl chlorides or phenol derivatives. This protocol exhibits broad substrate scope and excellent functional group compatibility, proceeds under mild reaction conditions, and is applicable in late-stage functionalization. Computational studies reveal that the unsymmetric, bulky yet flexible NHC ligand was the key to the success of this challenging transformation, which allowed for well-balanced rates for the elementary catalytic steps. We anticipate that this Ni/NHC-catalysed protocol would offer a general and powerful platform for C–N cross-couplings in medicinal research and chemical synthesis. We expect that this intriguing discovery would inspire further design of sterically flexible ligands to promote other challenging metal-catalysed cross-coupling reactions *via* balancing the elementary steps.

Data availability

General experimental conditions, calculations related to the mechanism elucidation, and characterization of all compounds synthesized are available in the ESI.†

Author contributions

Z.-C. W. developed the catalytic system and performed the reaction optimization and scope study, Y.-Y. Li and S.-Q. Zhang conducted the DFT calculations under the supervision of X. H., S.-L. S., X. H. and Z.-C. W. wrote the manuscript with inputs from the other authors. S.-L. S. conceived and directed the project.

Conflicts of interest

There are no conflicts to declare.

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computing system at the Department of Chemistry, Zhejiang University.

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