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Exploiting the trifluoroethyl group as a precatalyst ligand in nickel-catalyzed Suzuki-type alkylations†

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We report herein the exploitation of the partially fluorinated trifluoroethyl as precatalyst ligands in nickel-catalyzed Suzuki-type alkylation and fluoroalkylation coupling reactions. Compared with the $[L_nNi^{II}(aryl)(X)]$ precatalysts, the unique characters of bis-trifluoroethyl ligands imparted precatalyst $[(bipy)Ni(CH_2CF_3)_2]$ with bench-top stability, good solubilities in organic media and interesting catalytic activities. Preliminary mechanistic studies reveal that an eliminative extrusion of a vinylidene difluoride (VDF, $CH_2=CF_2$) mask from $[(bipy)Ni(CH_2CF_3)_2]$ is a critical step for the initiation of a catalytic reaction.

Introduction

Transition metal catalyzed cross-coupling reactions have advanced organic synthesis in the last few decades and have become powerful tools for the generation of molecular complexity.¹ Substantial effort has been devoted to identifying general and robust transition metal catalytic systems for reaction methodology research and chemical production improvement. A prominent example is the development of Suzuki–Miyaura coupling systems, which now employ a diverse combinations of transition-metals, supporting ligands, and coupling partners to construct $C(sp^2)–C(sp^2)$ bonds.² Although Pd catalysts operate with much success in this arena,³ the development of Ni-catalyzed protocols has been of interest because of the cost efficiency and complementary reactivities.⁴ For instance, Ni-catalyzed couplings are particularly useful for constructing synthetically challenging $C(sp^2)–C(sp^3)$ linkages,^{5–7} due to the facile oxidation of low-valent nickel by $C(sp^3)$ -centered electrophiles and the suppression of undesired β -hydrogen eliminations at nickel.^{4,8} One of the most successful catalysts for nickel-catalyzed coupling reactions is derived from the $[(bipyridine)nickel]$ motif which has been widely employed for both traditional cross-coupling and photoredox catalysis.^{5–7} However, it should be noted that the conventional $[(bipyridine)$

nickel] systems characterized by a combination of Ni^0 catalysts or inorganic Ni^{II} salts with bipyridyl ligand still suffer from some unneglectable limitations: (i) commonly used Ni^0 sources for catalysis are expensive and air-sensitive, thus hindering their use out of glovebox for large-scale synthesis; (ii) the low solubility of inorganic Ni^{II} salts complicates the heteroleptic coordination of exogenous supporting ligands which could have deleterious effects on reaction outcomes.

In this context, the development of robust nickel-based precatalysts in which the metallic cores are preligated with privileged ligands is highly desirable and constitutes a viable solution to address the above-mentioned limitations.⁹ Recently, the carbon-bound nickel precatalysts have exerted powers in a variety of coupling reactions as alternatives to the conventional $[L_nNiX_2]$ precatalysts ($L = P$ or N ligands).¹⁰ Notably, many previously reported carbon-bound Ni precatalysts $[L_nNi(X)(R)]$ feature sterically bulky ligands ($R = o$ -tolyl, mesityl, 1-naphthyl), or highly stabilizing motifs ($R = \eta^3$ -allyl, η^5 -Cp) for sheltering reactive organometallic nickel cores (Fig. 1-A).¹⁰ Considering that fluoroalkyl ligands are known to confer enhanced stability to metal complexes relative to their non-fluorinated alkyl counterparts owing to fluorine's unique electronic and steric properties,^{11–13} we wondered whether the incorporation of selected fluoroalkyl moieties could support novel nickel-based precatalysts and render new catalytic activities for use in synthetic methods development (Fig. 1-B). Herein, we describe the synthesis of such a fluoroalkyl-bound nickel precatalyst and demonstrate its use in $C(sp^2)–C(sp^3)$ Suzuki-type coupling reactions.

Result and discussion

At the outset, we began the rational design of precatalyst based on the principles of utilizing short fluoroalkyl and bipyridine as supporting ligands for atomic economy and $C(sp^2)–C(sp^3)$

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A. Well-defined carbon-ligated nickel precatalysts

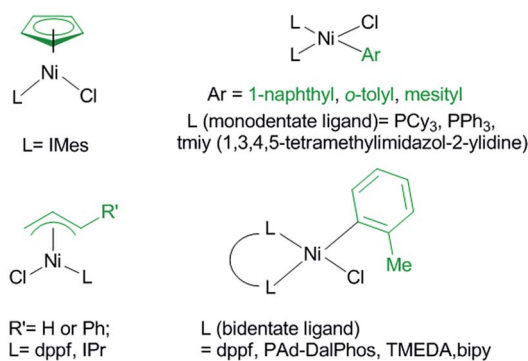
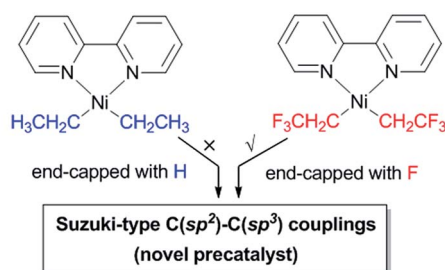
B. Fluoroalkyl-bound Ni-based precatalyst for C(sp²)-C(sp³) couplings

Fig. 1 Strategy for the development of fluoroalkyl-bound nickel precatalyst.

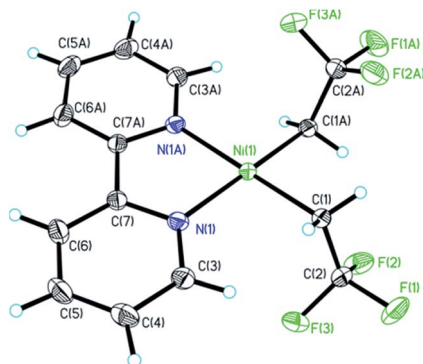
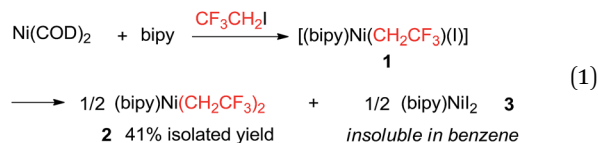


Fig. 2 ORTEP diagram of precatalyst [(bipy)Ni(CH₂CF₃)₂] 2.

coupling reaction efficiency. Specifically, the short and partially fluorinated CF₃CH₂ group was selected as supporting ligand (analogue of ethyl group but end-capped with fluorines) which was anticipated to render distinctive thermostability and reactivities *versus* both the hydrocarbonated [(bipy)Ni(CH₂CH₃)₂]¹⁴ and perfluorinated [(bipy)Ni(CF₂CF₃)₂]^{12g} counterparts (Fig. 1B, bipy = 2,2'-bipyridine). Gratifyingly, the reaction of [Ni(COD)₂], CF₃CH₂I, and 2,2'-bipyridine furnished [(bipy)Ni(CH₂CF₃)₂] 2 in 41% isolated yield (eqn (1)) presumably *via* an interesting ligand redistribution^{12c,15} of the intermediate [(bipy)Ni(CH₂CF₃)(I)] 1. The solubility of complex 2 in benzene facilitated its isolation from the nickel halide co-products 3. The ¹⁹F NMR spectra of 2 exhibited a triplet at δ = 47.98 ppm, demonstrating the presence of CH₂CF₃ groups and their bonding to Ni core. Dark red

crystals of 2 can be grown from THF/pentane and are air-stable at room temperature for several weeks.



X-ray diffraction analysis of 2 confirmed the ligation of two CH₂CF₃ groups at nickel (Fig. 2). Complex 2 featured a square planar arrangement at the Ni^{II} core with a rough linear *trans* N-Ni-C linkage (bond angle: 177.4(2) and 177.8(2)°). In contrast, more striking distortions were found in the previously reported and related complexes [(bipy)Ni(CF₃)₂] 4 (*trans* N-Ni-C bond angles: 165.1(2)° and 159.7(2)°) and [(bipy)Ni(CF₂CF₃)₂] 5 (both at 152.2)°,^{12g,12i} indicating fewer steric and electronic repulsions of the CH₂CF₃ chains in 2 compared to the perfluorinated derivatives. Interestingly, Ni-C distances of 2 (1.944(5) and 1.942(4) Å) are substantially longer than those of 4 (1.872(6) and 1.883(6) Å) and 5 (1.910(6) and 1.911(6) Å). Besides, the value of C(2)-F(3) bond length [1.366(6) Å, *trans* coplanar to C(1)-Ni bond] was clearly larger than the others two carbon-fluorine bonds [C(2)-F(1) 1.346(5) Å; C(2)-F(2) 1.342(6) Å] which implied the possible use of β-fluorine elimination for further coupling reaction development.

Although we did not obtain the one fluoroalkyl accommodated nickel complex [(bipy)Ni(CH₂CF₃)(I)] 1 which showed more structural similarities to the reported [(bipy)Ni(*o*-tolyl)Cl] precatalyst,^{14b} we presumed that β-fluorine elimination¹⁶ of [(bipy)Ni(CH₂CF₃)₂] 2 hinted by the C-F bond length analysis could be leveraged for the *in situ* generation of [(bipy)Ni(F)(CH₂CF₃)] 2a with concurrent extrusion of vinylidenedifluoride (CH₂=CF₂). Notably, the Ni-F structural motif of intermediate 2a was supposed to facilitate the transmetalation of arylboronic acids towards nickel according to a recent example of base-free Suzuki coupling.^{4g} Furthermore, the bis-trifluoroethyl structural motifs of [(bipy)Ni(CH₂CF₃)₂] 2 entails the bench-top stability and excellent solubility in organic solvents which is of vital importance for developing nickel-based precatalysts.^{9,10}

With [(bipy)Ni(CH₂CF₃)₂] 2 in hand, we initially assessed it as a precatalyst for the Suzuki-type coupling between CF₃CH₂I and arylboronic acids for C(sp²)-C(sp³) bonding. Based upon previously established Ni-catalyzed trifluoroethylation conditions,¹⁷ we were pleased to find that coupling products can be obtained in excellent yields at 80 °C with 5 mol% catalyst loading using K₃PO₄ as a base and DME as a solvent (Table 1, entry 1). Use of other solvents decreased the yields, and only polar non-protic DMSO solvent was comparatively effective. Furthermore, the use of K₃PO₄ was critical to the success of the coupling reaction and suppressing dehydrofluorination of the final products (for details, see ESI Tables S1 and S2†). The commercialized [(TMEDA)Ni(*o*-tolyl)Cl]^{10e,10f} bearing modular TMEDA was found to be less efficient (yield 35%) with using the privileged bipyridine as the leading supporting ligand (Table 1, entry 2). In contrast, the bipyridine preligated [(bipy)Ni(*o*-tolyl)



Table 1 Survey of reaction conditions^a

Entry	Variation from standard conditions	Isolated yield
1	None	93%
2	[(TMEDA)Ni(<i>o</i> -tolyl)Cl] (5.0 mol%) and 2,2-bipyridine (5.0 mol%) instead of 2	35%
3	[(bipy)Ni(<i>o</i> -tolyl)Cl] (5.0 mol%) instead of 2	78%
4	[(dppf)Ni(<i>o</i> -tolyl)Cl] (5.0 mol%) instead of 2	25%
5	[Ni(COD) ₂] (5.0 mol%) and 2,2-bipyridine (5.0 mol%) instead of 2	40%
6	[(bipy)NiEt ₂] (5.0 mol%) instead of 2	13%
7	[(MeCN) ₂ Ni(CF ₃ CF ₃) ₂] (5.0 mol%) and 2,2-bipyridine (5.0 mol%) instead of 2	55%
8	[(MeCN) ₂ Ni(CF ₃) ₂] (5.0 mol%) and 2,2-bipyridine (5.0 mol%) instead of 2	51%
6	2.5 mol% loading of precatalyst 2	91% (83% ^b)
7	1.0 mol% loading of precatalyst 2	79%

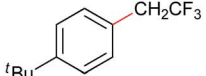
^a General conditions: 4-biphenylboronic acid (0.3 mmol), CF₃CH₂I (0.2 mmol), K₃PO₄ (0.4 mmol), 5.0 mol% precatalyst loading of **2**, DME (1.0 mL).

^b Gram-scale synthesis.

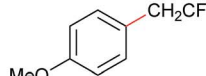
Cl]^{14b} bearing the privileged *o*-tolyl ligand improved the coupling yield to 78% but was still inferior to that of [(bipy)Ni(CH₂CF₃)₂]. These results demonstrated the importances of supporting ligation groups of both trifluoroethyl and bipyridine in the structural motif of [(bipy)Ni(CH₂CF₃)₂] (Table 1, entries 2–3). Even the classical [(dppf)Ni(*o*-tolyl)Cl]^{10c} or [Ni(COD)₂]/

Table 2 Substrate scope of (hetero)arylboronic acid partners^a

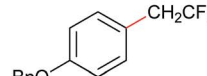
$$\begin{array}{c}
 \text{(Het)Ar-B(OH)}_2 + \text{CF}_3\text{CH}_2\text{I} \xrightarrow[\text{K}_3\text{PO}_4, \text{DME}, 80^\circ\text{C}]{2.5 \text{ mol\% of } \mathbf{2}} \text{(Het)Ar-CH}_2\text{CF}_3 \\
 \mathbf{6} \qquad \qquad \qquad \qquad \qquad \qquad \qquad \qquad \qquad \qquad \qquad \qquad \qquad \qquad \qquad \mathbf{7}
 \end{array}$$



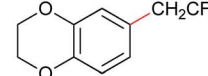
7b, 79%^b



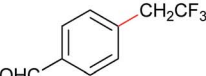
7c, 86%^b



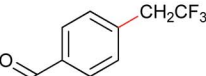
7d, 83%^b



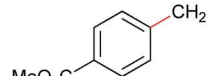
7e, 80%^b



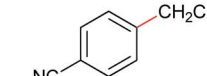
7f, 70%^b



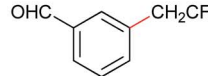
7g, 74%^b



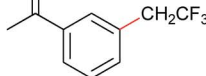
7h, 80%^b



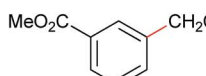
7i, 72%^b



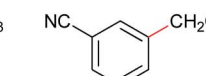
7j, 48%^b



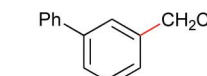
7k, 67%^b



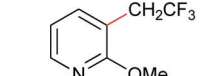
7l, 63%^b



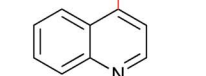
7m, 44%^b



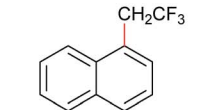
7n, 74%^b



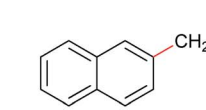
7o, 78%^b



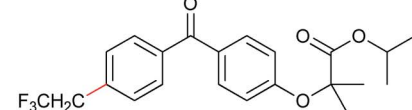
7p, 82%^b



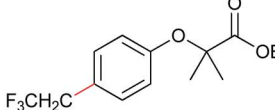
7q, 70%^c (36%)



7r, 87%^c (42%)



7s, 61%^b (Fenofibrate analogue)



7t, 44%^{b,d} (Clofibrate analogue)

^a General conditions: (hetero)arylboronic acid (0.6 mmol), CF₃CH₂I (0.4 mmol), base (0.8 mmol), 2.5 mol% precatalyst loading, DME (2.0 mL), 80 °C. ^b Isolated yield. ^c Yield determined by ¹⁹F NMR spectroscopy using PhCF₃ as an internal standard due to the volatility of naphthalene products. Data in parentheses refer to yields of isolated products. ^d 5.0 mol% precatalyst loading.



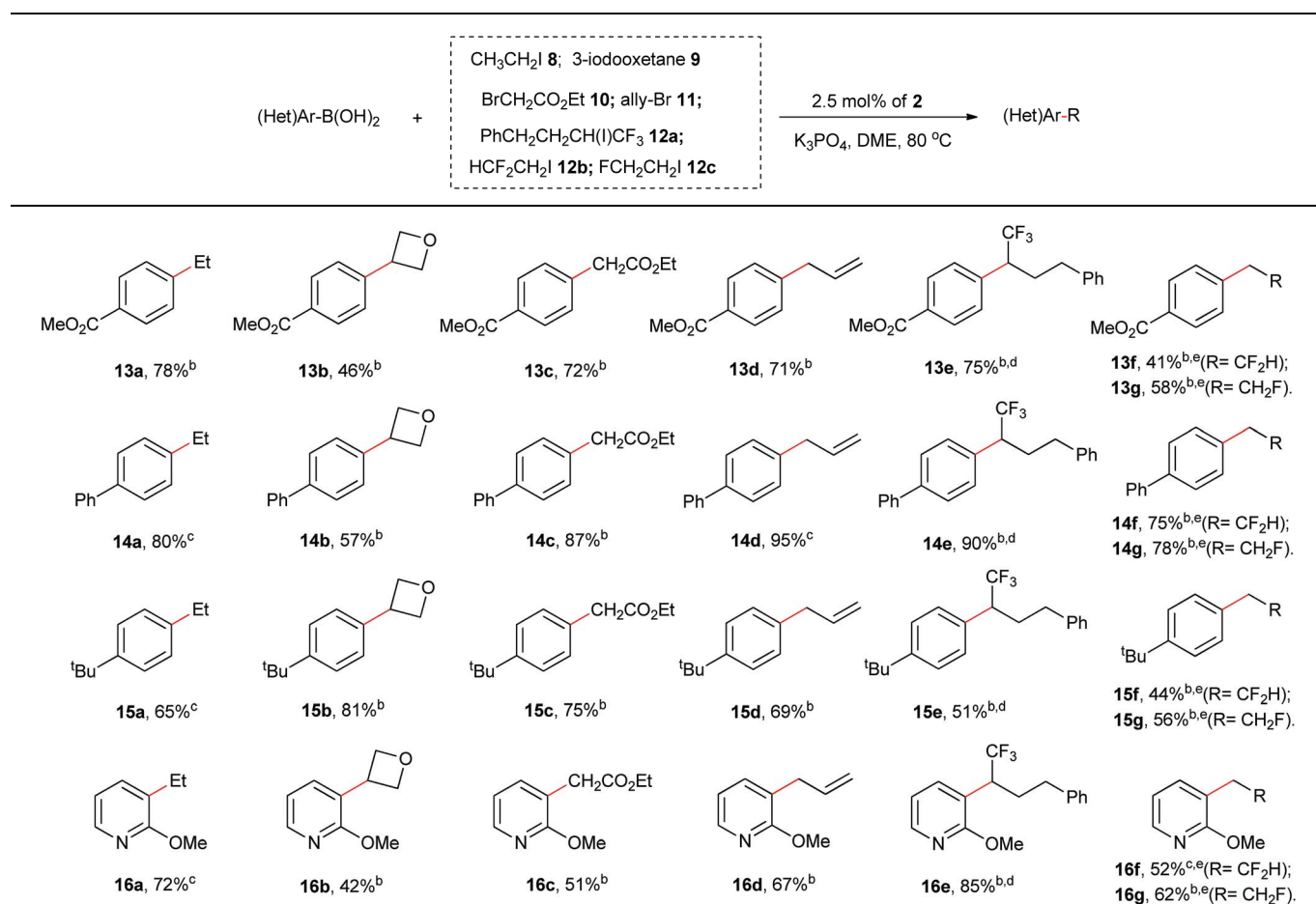
bipyridine¹⁸ combined system gave unsatisfactory yield (25–40%) compared with the use of **2** (Table 1, entries 4–5). For further demonstrating the distinctive role of partially fluorinated trifluoroethyl ligand, we compared its catalytic performance with those surrogating [(bipy)NiEt₂],^{14c} [(bipy)Ni(CF₂CF₃)₂]¹²ⁱ and [(bipy)Ni(CF₃)₂]^{12j} (Table 1, entries 6–8). It was found that these fully hydrocarbonated and fluorinated counterpart complexes can not furnish comparable catalytic outcomes. Gratifyingly, the tests of decreasing the precatalyst loading and gram-scale synthesis also provided the coupling product in comparatively good yields (Table 1, entries 6–7). These results demonstrated proof-in-principle of the excellent catalytic efficiency of precatalyst **2** for the targeted Suzuki-type couplings.

Under the optimized conditions, a wide array of arylboronic acid coupling partners were found to successfully participate in the Suzuki-type trifluoroethylation catalyzed by **2** (Table 2). Both the electron-donating and electron-withdrawing groups substituted arylboronic acids were competent substrates and gave the desired product in moderate to good yield. Broad

functional groups were well tolerated, including ethers (**7c–7e**), aldehydes (**7f**, **7j**), enolizable ketones (**7g**, **7k**), esters (**7h**, **7l**) and nitriles (**7i**, **7m**). Notably, the nitrogen-containing heterocyclic boronic acids (**7o**, **7p**) proceeded smoothly with good yields despite the potential strong binding affinity of the nitrogen atoms with Ni. To further exhibit the synthetic practicality of our precatalyst and trifluoroethylation protocol, the late-stage modifications of fenofibrate and clofibrate (drugs against cardiovascular disease) were accomplished (**7s**, **7t**). Therefore, this synthetic strategy should provide important opportunities for making more diverse biologically active molecules.

Further demonstration of the privileged catalytic utilities of precatalyst [(bipy)Ni(CH₂CF₃)₂] **2** was showcased by several types of C(sp²)–C(sp³) Suzuki-type alkylations. Iodoethane (**8**), 3-iodooxetane (**9**), ethyl bromoacetate (**10**), allyl bromide (**11**), (4,4,4-trifluoro-3-iodobutyl)benzene (**12a**), HCF₂CH₂I (**12b**) and FCH₂CH₂I (**12c**) were found to successfully couple with a series of arylboronic acids ranging from electron-poor and electron-rich types (Table 3). The encouraging results showed

Table 3 Versatility of precatalyst **2** for aryl–alkyl cross-coupling reactions^a



^a General conditions: (hetero)arylboronic acid (0.6 mmol), the indicated R–X (0.4 mmol), base (0.8 mmol), 2.5 mol% precatalyst loading, DME (2.0 mL), 80 °C. ^b Isolated yield. ^c Yield determined by ¹H NMR spectroscopy using Cl₂CHCHCl₂ as an internal standard due to the difficulties in separation of product from deboronative byproduct. ^d Using 5.0 mol% precatalyst loading and DMSO as the solvent instead of DME. ^e Using 5.0 mol% precatalyst loading.



that these primary and secondary alkyl halides were readily compatible with this catalytic alkylation, regardless of the possible β -H or β -F elimination problems.^{8a} Furthermore, the active allyl group can be coupled with the aromatic groups without detection of the migration of double bond (**13d**).⁷

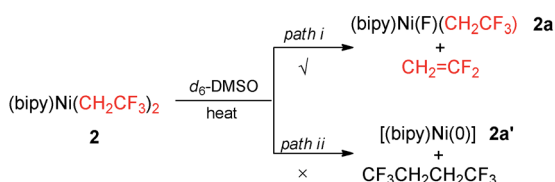
The success of precatalyst **2** for the Suzuki-type alkylations further encouraged us to investigate the reaction mechanism. At the start, we intended to determine whether the activation mode of $[(bipy)Ni(CH_2CF_3)_2]$ was consistent with our β -fluorine elimination hypothesis. The precatalyst could undergo β -fluorine elimination to afford $[(bipy)Ni(F)(CH_2CF_3)]$ **2a** with extrusion of $CH_2=CF_2$, or alternatively undergo reductive elimination like the analogous $[(bipy)Ni(CH_2CH_3)_2]$ ¹⁴ to generate a $[(bipy)Ni(0)]$ species and $CF_3CH_2CH_2CF_3$. Heating precatalyst **2** at elevated temperature indicated the clear formation of $CH_2=CF_2$ rather than $CF_3CH_2CH_2CF_3$ through continuous 1H and ^{19}F NMR monitoring (for details, see ESI Fig. S122 and S123†) which illustrated the direct formation of low-valent nickel species like $[(bipy)Ni(0)]$ from $[(bipy)Ni(CH_2CF_3)_2]$ **2** is less likely (Scheme 1-A).¹⁹ Additionally, the identification of $CH_2=CF_2$ and $Ar-CH_2CF_3$ (Ar = 4-biphenyl) in GC-MS and NMR analysis of the reactions in Table 3 (e.g. **14b**, coupling between 4-biphenylboronic acid and 3-iodooxetane in Scheme 1B) revealed the important roles of the trifluoroethyl groups bound to nickel core (for details, see ESI Fig. S124, S125 and Table S5†). These results suggested that the first trifluoroethyl group functioned as the mask of the suggested active species $[(bipy)Ni(F)(CH_2CF_3)]$ **2a** via a $CH_2=CF_2$ extrusion and the second trifluoroethyl moiety contributed as coupling partner for the formation of $Ar-CH_2CF_3$. Interestingly, the finding of byproduct $CH_3OCH(Ar)CH_2OCH_3$ and $ArCH_2OCH_2CH_2OCH_3$ (Scheme 1-B) illustrated plausible radical activation of DME

through abstraction of ethereal α -hydrogens by solvent-caged alkyl radicals.²⁰

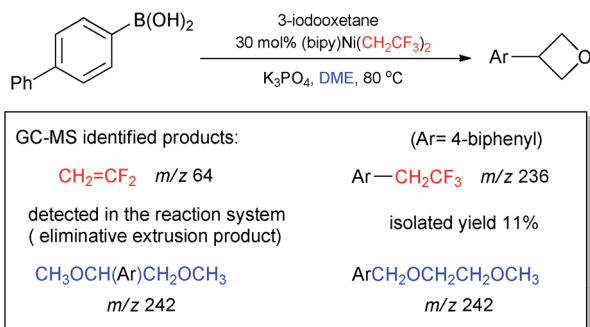
Next, a series of radical inhibition experiments were conducted to verify the possibilities of radical intermediacy (for details, see ESI Tables S8–S10†). It was found that the radical scavenger TEMPO shut down the coupling reactions completely when using the 3-iodooxetane or CF_3CH_2I as the alkyl electrophiles. Instead, TEMPO-alkyl (alkyl = 3-oxetanyl or trifluoroethyl) adducts **17** and **18** were observed in the GC-MS analysis, respectively. Also, when a radical-clock cyclopropane-based substrate was used, a few ring-opening products like the CF_3CH_2 -merged product **20** and aryl-incorporated product **21** were identified. These experimental results suggested the involvement of $CF_3CH_2\cdot$ radicals (or R radicals) as well as aryl-bound nickel intermediates in the reaction profile.

With the above clues of reaction scenarios in hand, we conducted further interrogations on whether the reactions proceeded via a Ni^0/Ni^{II} or Ni^I/Ni^{III} redox shuttle. The important findings of bis-trifluoroethyl ligands of **2** serving as $CH_2=CF_2$ mask and operational ligand for producing $Ar-CH_2CF_3$ inspired us to devise a stoichiometric reaction of complex **2** with 4-biphenylboronic acid as control experiment (Scheme 2-A). The intermediate $[(bipy)Ni(F)(CH_2CF_3)]$ **2a** could be generated *in situ* under the reaction conditions which was supposed to further undergo a facile Ni–B transmetalation^{4g} to deliver $[(bipy)Ni(Ar)(CH_2CF_3)]$ **2b** (Ar = 4-biphenyl). However, the putative $[(bipy)Ni(Ar)(CH_2CF_3)]$ intermediate did not proceed through a $Ni(II)/Ni(0)$ reductive elimination^{10e} to furnish $Ar-CH_2CF_3$. In addition, CF_3CH_2F and $CF_3CH_2CH_2CF_3$ were also not found which disfavored the scenario of Ni^0 formation from the reductive elimination of **2a** and **2**. Taken together, these divalent organonickel intermediates (**2**, **2a** and **2b**) were not productive for the corresponding Ni^{II}/Ni^0 reductive elimination under this current reaction system. Interestingly, the product

A. NMR studies on the activation of precatalyst **2**

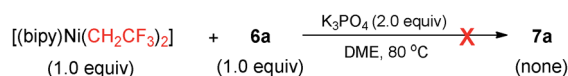


B. Probe the role of the nickel-bound trifluoroethyl as coupling partner



Scheme 1 Control experiments for identifying the role of trifluoroethyl ligands in precatalyst **2**.

A. Stoichiometric reaction of 4-biphenylboronic acid with precatalyst **2**

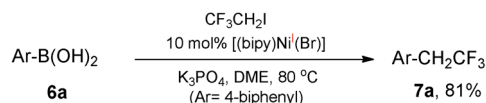


GC-MS identified products:

$CH_2=CF_2$ m/z 64	biphenyl m/z 154
detected in reaction system (evidence for eliminative extrusion)	deborylation product

Not detected:	(Ar = 4-biphenyl)
$Ar-CH_2CF_3$	CF_3CH_2F
	$CF_3CH_2CH_2CF_3$

B. Trifluoroethylation catalyzed by a putative univalent $[(bipy)Ni^I(Br)]$



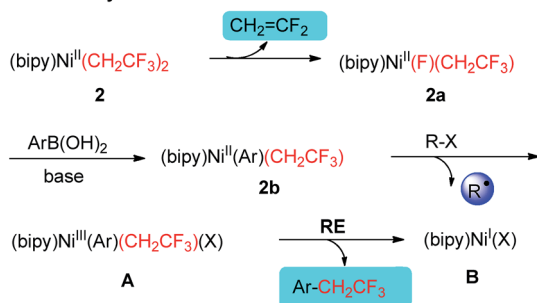
Scheme 2 Control experiments to support Ni^I/Ni^{III} redox shuttle in the catalytic cycle.



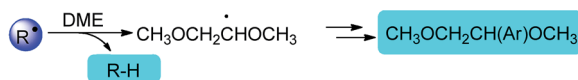
$\text{Ar-CH}_2\text{CF}_3$ was almost comparably efficiently obtained when the precatalyst **2** was replaced by the putative $[(\text{bipy})\text{Ni}^{\text{I}}(\text{Br})]$ complex²¹ for the coupling between $\text{ArB}(\text{OH})_2$ and $\text{CF}_3\text{CH}_2\text{I}$ (Scheme 2-B). These experimental results suggested that a $\text{Ni}^{\text{I}}/\text{Ni}^{\text{III}}$ catalytic cycle was highly likely to be superior to $\text{Ni}^0/\text{Ni}^{\text{II}}$ counterpart in the current reaction systems.

Based on the above-mentioned experimental results and relevant previous reports,^{5,6,13,22} a plausible mechanism was proposed for these current cross-couplings (Scheme 3). The catalysis commences with an eliminative liberation of a $\text{CH}_2=\text{CF}_2$ mask and **2a**. Intermediate **2a** is proposed to undergo transmetalation and subsequent abstraction of halogen atom from R-X to afford $(\text{bipy})\text{Ni}^{\text{III}}(\text{Ar})(\text{CH}_2\text{CF}_3)(\text{X})$ **A**. Reductive elimination of **A** has been fingerprinted by the formation of $\text{Ar-CH}_2\text{CF}_3$ and delivered a key catalytic species $(\text{bipy})\text{Ni}^{\text{I}}(\text{X})$ **B**. Upon the participation of **B** into the conventional $\text{Ni}^{\text{I}}/\text{Ni}^{\text{II}}/\text{Ni}^{\text{III}}$ catalytic cycle, the shuttles *via* transmetalation/oxidative addition²²/reductive elimination provided efficient platform for the above described Suzuki-type $\text{C}(\text{sp}^2)\text{-C}(\text{sp}^3)$ alkylation couplings.²³

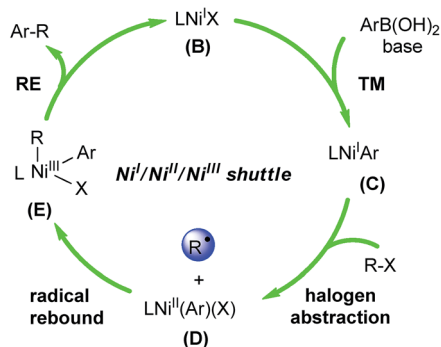
A. Precatalyst Initiation:



B. Side reactions of R radical intermediate:



C. Possible Catalytic cycle (L = bipy):



Scheme 3 Proposed reaction mechanism for Suzuki-type alkylation couplings based on control experiments. The shadow depicted species in precatalyst initiation and radical-relay side reactions were fingerprinted by GC-MS.

Conclusions

In conclusion, we demonstrated that the nickel-based precatalyst **2** $[(\text{bipy})\text{Ni}(\text{CH}_2\text{CF}_3)_2]$ can be employed in Suzuki-type coupling reactions between (hetero)arylboronic acids and a variety of alkyl halides including several typical partially fluorinated alkyl halides bearing susceptible β -fluorine atoms (2-iodo-1,1,1-trifluoroethane and **12a–12c**), leading to new $\text{C}(\text{sp}^2)\text{-C}(\text{sp}^3)$ linkages. Catalytic performance tests demonstrated the advantages of the trifluoroethyl ligand motifs in the precatalyst $[(\text{bipy})\text{Ni}(\text{CH}_2\text{CF}_3)_2]$ *versus* several sibling perfluorinated and hydrocarbonated counterparts.²⁴ The critical roles of trifluoroethyl groups of precatalyst **2** as both $\text{CH}_2=\text{CF}_2$ mask and triggering coupling-ligand in these nickel-catalyzed Suzuki-type alkylations were elucidated through mechanistic investigations. We believe that the initial success outlined here could prompt the utilization of more fluoroalkyl binding moieties for the development of new metal-based precatalysts with tailored activities. Further studies towards this endeavor and mechanistic details are underway in our laboratory, and the results will be reported in due course.

Conflicts of interest

We authors announce that there are no conflicts to declare.

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