Pd/Ni-Catalyzed Germa-Suzuki Coupling *via* Dual Ge–F Bond Activation

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Abstract:

Pd/Ni→Ge–F interactions supported by phosphine-chelation were found to trigger dual activation of Ge–F bonds under mild conditions. This makes fluoro germanes suitable partners for catalytic Ge–C cross-coupling and enables Germa-Suzuki reactions to be achieved for the first time.

Organo germanium compounds are powerful transfer reagents for catalytic cross-coupling reactions.¹ They also find applications in materials science.² The formation of C–Ge bonds is thus of importance and the advent of new synthetic methodologies is highly desirable to widen the scope of usable substrates and accessible products.

Hydrogermylation involving transition metal or radical-mediated pathways is efficient and useful, but intrinsically limited to the synthesis of alkyl- and alkenyl-substituted germanes.^{3–5} The most general way to forge Ge–C bonds is probably the ionic coupling of organometallic compounds with germanium halides or alkoxides (reactions between Ge-based nucleophiles and organic electrophiles are also known but less common).⁶ Transition metal catalysis is extremely powerful and broadly used for C–C and C–X bond formation. It has certainly great potential for Ge–C cross-coupling as well, although the field is still in its infancy with only very few recent reports (Figure 1). Xiao and co-workers reported in 2018 Pd-catalyzed couplings of a hydrogermatrane with aryl halides and pseudo-halides.^{1d} The approach was then extended by Schoenebeck *et al.* to the coupling of Et₃GeH with aryl thiathrenium salts using Pd(I) dimers as catalysts.^{1k} In addition, Oestreich *et al.* significantly advanced the use of Ge-based nucleophiles in Ge–C(sp³) cross-coupling and described in 2019 a very elegant Germa-Negishi type reaction.⁷ We envisioned a radically different approach, namely cross-coupling with a Ge-based electrophile.⁸ Our recent discovery of facile Si–F bond activation thanks to transition metal / Lewis acid cooperation opened the way to catalytic Sila-Negishi coupling from fluoro-silanes.⁹ Here we demonstrate that this concept can be extended to Germanium chemistry. The combination of Pd or Ni with a Lewis acid enables dual activation of Ge–F bonds.¹⁰ Accordingly, fluoro germanes were found to undergo Ge–C cross-couplings with organo boron reagents, providing the first examples of Germa-Suzuki reactions.

Figure 1 Transition metal-catalyzed Ge–C cross-couplings.

Fluoro germanes have been shown in previous studies to engage in M→Ge–F interactions (M = Cu, Ag, Au) when chelated by phosphines, and to be stronger π -acceptor ligands than fluoro silanes.¹¹ We thus envisioned to take advantage of P-chelation to trigger Ge–F activation and cross-coupling. Given the availability and efficiency of boron reagents as nucleophiles, we sought to develop a hitherto unknown Germa-Suzuki reaction. The fluoro germane $\{(o\text{-Ph}_2\text{P})\text{C}_6\text{H}_4\}\text{GePh}_2\text{F}$ **1^{Ge}** was used as substrate and several aryl boron derivatives were tested. Gratifyingly, quantitative Ge–Ph coupling was observed using BPh₃ with either Pd₂(dba)₃ or Ni(COD)₂ (Table 1, entries 1 and 2). The

nature of the borane is critical: no reaction occurred when the less Lewis acidic borane PhBpin or the very Lewis acidic borane B(C₆F₅)₃ were used (entries 3 and 4). The formation of F₂BPh, BF₃, and BF₄[−] by-products from BPh₃ (as apparent from ¹⁹F{¹H} NMR spectroscopy, Figure S1)[†] suggests that the borane acts both as Ph source (arylated reagent) and a F acceptor (Lewis acid) in this transformation. Of note, the corresponding fluoro silane {(o-Ph₂P)C₆H₄}SiPh₂F (**1**^{Si}) remains inert under similar conditions (entry 5), indicating that only the Ge–F bond can be activated and cross-coupled this way, probably due to stronger M→Ge–F interaction. Phenylation of the diphosphine fluoro germane {(o-Ph₂P)C₆H₄}₂GePhF (**3**)¹⁰ was also possible with both Pd and Ni catalysts using BPh₃ (entries 6 and 7). Again, no cross-coupling was observed using PhBpin or B(C₆F₅)₃ (entries 8 and 9). Varying the aryl source and Lewis acid with **1**^{Ge} as substrate, we then realized that the reaction can be most conveniently achieved using the trifluoroborate salt K(F₃BPh) along with BF₃. With this combination, modest results were obtained in THF (probably due to inhibition of the Lewis acid-assisted Ge–F activation). However, shifting to 2-methyl tetrahydrofuran (2-MTHF) as solvent drastically improved the catalytic activity and the desired Ge–Ph coupling product **2** was obtained in excellent yields (entries 10 and 11). Note that K[F₃BPh] alone is unproductive, the concomitant use of BF₃ is essential for the catalysis to proceed (entry 12). Phosphine chelation also plays an important role, no reaction being observed under the same conditions from FGePh₃.

Table 1 Germa-Suzuki cross-coupling reactions of the fluoro germanes **1**^{Ge} and **3**.

Entry	Substrate	Catalyst	Ar source / L. A.	Yield(%) ^a
1	1 ^{Ge}	Pd ₂ (dba) ₃	BPh ₃ ^b	99
2	1 ^{Ge}	Ni(COD) ₂	BPh ₃ ^b	99
3	1 ^{Ge}	Pd ₂ (dba) ₃	PhBpin ^c	0
4	1 ^{Ge}	Pd ₂ (dba) ₃	B(C ₆ F ₅) ₃ ^b	0
5	1 ^{Si}	Pd ₂ (dba) ₃	BPh ₃ ^b	0
6 ^d	3	Pd ₂ (dba) ₃	BPh ₃ ^b	79
7 ^d	3	Ni(COD) ₂	BPh ₃ ^b	47
8 ^d	3	Pd ₂ (dba) ₃	PhBpin ^c	0
9 ^d	3	Pd ₂ (dba) ₃	B(C ₆ F ₅) ₃ ^b	0
10	1 ^{Ge}	Pd ₂ (dba) ₃	K(F ₃ BPh)/BF ₃ ·OEt ₂	49 (96) ^e
11	1 ^{Ge}	Ni(COD) ₂	K(F ₃ BPh)/BF ₃ ·OEt ₂	8 (99) ^{e,f}
12	1 ^{Ge}	Ni(COD) ₂	K(F ₃ BPh)/none	0 (0) ^e
13	3	7 ^{Ge}	BPh ₃ ^b	86
14	3	8c ^{Ge}	BPh ₃ ^b	89

^aDetermined by ³¹P NMR. ^b1.5 eq. ^c5 eq. ^d10 mol% catalyst, Mesitylene, 160 °C, 20 h. ^e2-MTHF. ^fIsolated yield: 68%.

The substrate scope was then investigated employing the following optimal conditions: Ni(COD)₂, K(F₃BR) and BF₃ in 2-MTHF (Table 2). Phenylation of the fluoro germane {(*o*-Ph₂P)C₆H₄}GeMe₂F (**5**) featuring methyl substituents at Ge worked nicely. The reaction proceeded well with a variety of trifluoro(aryl)borates featuring electron-donating (OMe, NMe₂) as well electron-withdrawing (CN, CF₃, F, Cl) groups. Ge–C(sp³) coupling worked with K(F₃BBn) but not with *n*-alkyl substrates prone to β-H elimination such as K[EtBF₃] and K[*n*-BuBF₃].[‡]

Table 2 Scope of borate reagents for the Germa-Suzuki Coupling of fluoro germane **5**^a

^aSpectroscopic yields, as determined by ³¹P NMR spectroscopy (isolated yields are given in parentheses). ^b120 °C.

Competitive experiments¹² were then carried out with a 1:1 mixture of **5** and **5^{Cl}** in order to compare the reactivity of fluoro and chloro germanes towards Ge–C coupling (Figure 2). When ZnPh₂ was used as arylating reagent, only the chloro germane reacted to give **6-Ph**. An opposite chemoselectivity was achieved with Ni(COD)₂, K(F₃BR) and BF₃. Only the fluoro germane **5** was found to undergo cross-coupling under these conditions, **5^{Cl}** remained unchanged.

Figure 2 Competitive phenylations of Ge–Cl and Ge–F bonds.

By analogy with the related Sila-Negishi coupling,⁹ we propose the catalytic cycle shown in Fig. 3 to account for the Germa-Suzuki coupling. To support the feasibility and relevance of this pathway, the formation, structure and reactivity of Pd complexes deriving from the diphosphine fluoro germane **3** were investigated. Complex **7^{Ge}** was first synthesized by ligand exchange (step *i*). In contrast with the related diphosphine fluoro silane complex,⁹ **7^{Ge}** proved unstable in the absence of excess PPh₃. Notwithstanding, crystals of **7^{Ge}** suitable for X-ray diffraction analysis could be obtained (Figure 3 (a)). The Pd center is in tetrahedral environment, with the fluoro germane moiety coordinated *via* Ge (Pd...Ge 2.7518(5) Å) and C_{ipso} (Pd...C 2.638(3) Å) in addition to the three P atoms. The germane moiety behaves as a Z-type ligand in complex **7^{Ge}**.^{11,13,14} The Ge–Pd distance is only slightly longer than the sum of covalent radii (2.53 Å),¹⁵ and despite the larger radius of Ge compared to Si (1.20 vs 1.11 Å),¹⁵ it is significantly shorter than the Pd...Si distance in the Si analogue of **7^{Ge}** (2.9770(8) Å in [{(o-Ph₂P)C₆H₄}₂SiPhF][Pd(PPh₃)] (**7^{Si}**)).⁹ Additional signs for Pd→Ge–F interaction in complex **7^{Ge}** are the elongation of the Ge–F bond [1.8246(19) Å vs 1.762(2) Å in {(o-Ph₂P)C₆H₄}₃GeF¹⁶] and the trigonal pyramidal geometry around Ge [the sum of C–Ge–C angles = 358.81(22)°, Pd...Ge–F bond angle = 157.94(6)°]. Consistently, NBO analysis (DFT) shows the presence of a substantial donor-acceptor d(Pd)→σ*(Ge–F) interaction (15.8 kcal/mol) at the second-order perturbation level.

Figure 3 Catalytic cycle proposed to account for the Germa-Suzuki coupling. Molecular structures of the isolated catalytic intermediates **7^{Ge}** (a), **8^{Ge}** (b) and **9-C₆F₅** (c) complexes (the Ph substituents at phosphorus and germanium are simplified, the hydrogen atoms and counter-anion for **8^{Ge}** are omitted for clarity).

Activation of the Ge–F bond at Pd was then studied (step *ii*). Upon treatment with boranes such as BF₃, BPh₃ and B(C₆F₅), complex **7^{Ge}** readily afforded the cationic germyl complex [{(o-Ph₂P)C₆H₄]₂PhGe{Pd(PPh₃)}][FBR₃] (**8^{Ge}-FBR₃**) (R = F, Ph, C₆F₅)).¹⁷ The molecular structure of **8^{Ge}** was confirmed by X-ray diffraction analysis after counter-anion exchange (complex **8^{Ge}-BPh₄**, Figure 3 (b)). As expected, the Pd–Ge distance [2.4051(3) Å] is much shorter than in **7^{Ge}** and only marginally exceeds the sum of covalent radii. With a less Lewis acidic borane such as PhBpin, the germane **7^{Ge}**

remains intact, in line with the absence of catalytic activity observed when using PhBpin as coupling partner (entries 3 and 8, Table 1). Note that the facile Ge–F bond cleavage from **7^{Ge}** in the presence of BPh₃ markedly contrasts with the inertness of the Si–F bond of **7^{Si}** under similar conditions. The fluoro silane complex remains unchanged when treated with BPh₃ even upon heating at 160 °C (see below for a tentative explanation based on DFT calculations). Interestingly, the Ge–F bond activation is reversible. Indeed, the addition of tetrabutylammonium fluoride (TBAF) to **8^{Ge}-BPh₄** at room temperature immediately and quantitatively regenerates the fluoro germane complex **7^{Ge}**. The interconversion of Z-type and X-type ligands has recently attracted much interest and clear-cut examples have been authenticated with B,¹⁸ Si,⁹ Sb¹⁹ and Sn.²⁰ To our knowledge, the back and forth between **7^{Ge}** and **8^{Ge}** represents a first case with Z-germane and X-germyl moieties.

To gain further insight into the Ge–F bond activation from the fluoro germane **7^{Ge}** to the germyl **8^{Ge}** species, DFT calculations were performed with BPh₃ as Lewis acid (Figure 4). For comparison, similar studies were carried out on Si–F bond activation. The interaction of **7^{Ge}** with BPh₃ is slightly exergonic ($\Delta G = -0.4$ kcal/mol). It induces further elongation of the Ge–F bond (from 1.866 to 1.894 Å) while the Pd–Ge bond slightly shortens (from 2.777 to 2.741 Å). The subsequent bond cleavage is exergonic ($\Delta G = -6.3$ kcal/mol) and takes place with very small Gibbs activation energies ($\Delta G^\ddagger = 2.4$ kcal/mol) to give the cationic germyl complex **8^{Ge}-FBPh₃**. Activation of the Si–F bond of **7^{Si}** with BPh₃ proceeds similarly with a slightly higher but still low activation barrier ($\Delta G^\ddagger = 5.1$ kcal/mol). It is endergonic ($\Delta G = 3.7$ kcal/mol), why may explain why no catalytic conversion was observed for this substrate (entry 5 in Table 1). In line with this energy landscape, spontaneous fluoride transfer was observed upon mixing the fluoro germane **7^{Ge}** and silyl **8^{Si}-BPh₄** complexes to give the corresponding germyl **8^{Ge}-BPh₄** and fluoro silane **7^{Si}** species.[‡]

Figure 4 Reaction profiles computed for the Si/Ge–F bond cleavage at Pd assisted by BPh₃. Free Gibbs energy changes (kcal/mol) in THF.

To promote B-to-Pd transmetalation (step *iii*), **8^{Ge}-BPh₃F** was then thermolyzed. The corresponding phenyl complex $[(o\text{-Ph}_2\text{P})\text{C}_6\text{H}_4]_2\text{PhGe}\{\text{Pd}(\text{Ph})\}$ (**9-Ph**) was quantitatively obtained after 1 h at 160 °C. Complex **9-Ph** proved too unstable to be isolated in pure form, but the analogous pentafluorophenyl species $[(o\text{-Ph}_2\text{P})\text{C}_6\text{H}_4]_2\text{PhGe}\{\text{Pd}(\text{C}_6\text{F}_5)\}$ **9-C₆F₅** could be fully characterized (X-ray structure is shown in Figure 3 (c)).[‡] It is formed when reacting the fluoro germane complex **7^{Ge}** with B(C₆F₅)₃ followed by thermolysis. Transmetalation required prolonged heating (83 % conversion after 20 h at 160 °C) and was less clean (44% yield) in this case. The last step of the catalytic cycle, *ie* Ge–C coupling *via* reductive elimination (step *iv*), could also be achieved from **9-Ph**. Thermolysis in the presence of **3** (1 h, 160 °C) cleanly and quantitatively afforded the Ge-arylation product **4**.

along with the fluoro germane complex **7^{Ge}**. Note that heating **9-C₆F₅** led to intractable mixtures, with no detectable Ge–C₆F₅ product, in line with the much lower reactivity of C₆F₅ towards reductive elimination and the absence of catalytic coupling when using B(C₆F₅)₃ as Lewis acid / arylating reagent (Entries 4 and 9 in Table 1)

The germane **7^{Ge}** and germyl **8^{Ge}-BPh₄** complexes were engaged in catalytic coupling of the diphosphine fluoro germane **3** with BPh₃ (entries 13 and 14 in Table 1). Catalytic activities similar to that achieved with Pd₂(dba)₃ were obtained, supporting the proposed catalytic cycle. In addition, the cationic **8^{Ge}** and neutral **9-Ph** germyl complexes were detected by ³¹P NMR spectroscopy (in about 8:1 ratio) when monitoring the catalysis. This is consistent with the fact that transmetalation and reductive elimination require more forcing conditions than Ge–F bond activation.

In conclusion, Pd/Ni-catalyzed Germa-Suzuki coupling reactions have been carried out for the first time. Fluoro germanes were used as electrophilic coupling partners. The key Ge–F bond cleavage was achieved thanks to phosphine-chelated Pd/Ni→Ge–F interactions. Future work will aim to extend further the application of Z-type coordination to the activation and functionalization of strong σ-bonds.

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