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A Glove-box- and Schlenk-line-free Protocol for Solid-state C–N Cross-coupling Reactions Using Mechanochemistry

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ABSTRACT: Carbon–nitrogen (C–N) bond-forming cross-coupling reactions catalyzed by palladium-based catalysts, the so-called Buchwald–Hartwig aminations, have been widely employed for the synthesis of pharmaceuticals and aryl-amine-based organic materials in academic and industrial settings. However, in solution, these reactions usually require glove-box and Schlenk-line techniques, which greatly reduces their practical utility. Here, we report the development of operationally simple mechanochemical C–N cross-coupling reactions in the solid-state. Intensive investigations of various ball milling parameters revealed that the air-stable ligand tri(1-adamantyl)phosphine can be used to achieve solid-state coupling reactions between aryl halides and diarylamines with high efficiency. Notably, all experimental operations of the developed protocol can be carried out in air, thus providing a more convenient, industrially attractive, and sustainable alternative to conventional solution-based palladium-catalyzed C–N coupling reactions.

INTRODUCTION

The development of carbon–nitrogen (C–N) bond-forming coupling reactions catalyzed by transition-metals is of great importance in the synthesis of nitrogen-containing pharmaceuticals and organic electro and optical materials.[1-3] Palladium-based catalysts have been widely used for the cross-coupling of aryl halides with amine nucleophiles, the so-called Buchwald–Hartwig aminations.[1-9] However, these reactions usually necessitate synthetic techniques that involve glove boxes and/or high-vacuum Schlenk lines, which are costly and require special training.[10,11] Additionally, significant amounts of dry and degassed organic solvents are used in these reactions. Thus, the development of operationally simple, solvent-free palladium-catalyzed C–N coupling reactions would greatly improve the practicality of the desired aryl amine derivatives.

In this context, the use of ball-milling techniques for solvent-free palladium-catalyzed C–N bond-forming coupling reactions that can be carried out in air has attracted attention.[12-41] In 2018, Su and co-workers reported the first example of the use of ball milling to carry out solvent-free mechanochemical C–N coupling reactions in air in the presence of Pd(OAc)$_2$/XPhos as a catalyst.[31] Almost simultaneously, Browne and co-workers developed robust mechanochemical C–N coupling reactions in air using Pd-PEPPSI-iPENT as the catalyst.[32] Although these pioneering studies are remarkable, they are mainly focused on liquid substrates that serve as both reactants and the reaction

solvent. As most aryl-amine-based organic materials exhibit extended polyaromatic structures and are prepared from solid starting materials,[42-44] the development of mechanochemical cross-coupling reactions of solid aryl halides with solid amine nucleophiles in air would be highly desirable.
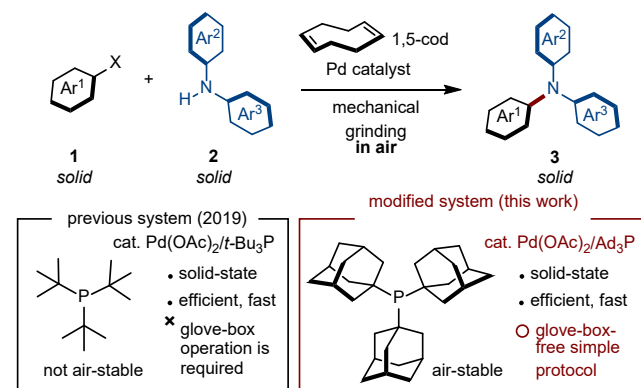
We have previously reported the first general solid-state palladium-catalyzed C–N cross-coupling reactions using mechanochemistry (Scheme 1A).[33] The key to achieving this mechanochemical transformation was the addition of small amounts of olefins such as 1,5-cyclooctadiene (1,5-cod). These additives can act as dispersants and stabilizers to facilitate the C–N cross-coupling in the solid-state reaction environment.[28] However, this procedure still requires glove-box techniques, as highly air-sensitive tri-*tert*-butylphosphine (*t*-Bu$_3$P) is used as the ligand.[45] This restriction greatly reduces the practical utility of the mechanochemical protocol.

Herein, we report a comprehensive investigation of various ball-milling parameters, which revealed that the air-stable tri(1-adamantyl)phosphine (Ad$_3$P)[46] can be used successfully in mechanochemical solid-state C–N coupling reactions under ambient conditions (Scheme 1A). Compared to the previous mechanochemical reaction using *t*-Bu$_3$P as the ligand,[45] the newly developed system shows comparable substrate generality. But more importantly, all experimental operations of the modified protocol can be carried out under ambient conditions, i.e., the chemicals, reagents, and catalysts can be added to the milling jar in air (Scheme 1B). Thus, this operationally simple protocol provides a more convenient, industrially attractive, and sustainable alternative to conventional solution-based palladium-catalyzed C–N coupling reactions.
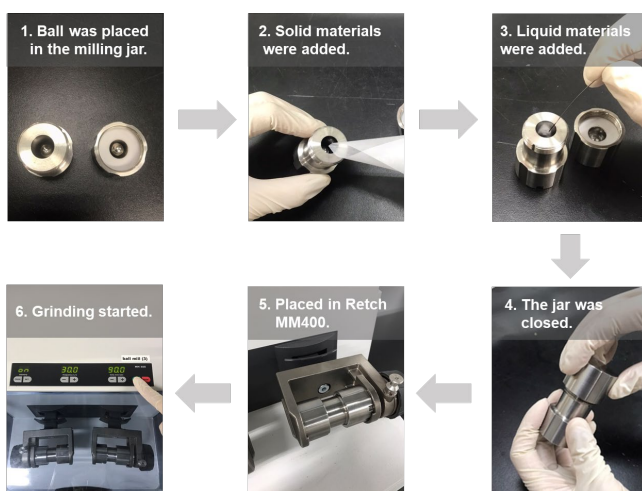
Mechanistic studies to elucidate the effect of varying the mechanochemical conditions on this solid-state C–N cross-coupling reaction are also described.

Scheme 1. Glove-box- and Schlenk-line-free Protocol for Solid-State C–N Cross-Coupling Reactions.

A Solid-state C–N cross-coupling using mechanochemistry

B Reaction set-up

1. Ball was placed in the milling jar.
2. Solid materials were added.
3. Liquid materials were added.
4. The jar was closed.
5. Placed in Retch MM400.
6. Grinding started.

RESULTS AND DISCUSSION

In light of our previous studies, we have been interested in identifying monodentate ligands that exhibit catalytic properties similar to or better than those of t-Bu₃P. To this end, we began our investigation by exploring the ligand effect in the solid-state mechanochemical C–N cross-coupling in the presence of 1,5-cod as a liquid-assisted-grinding (LAG) additive.[33] Reactions were conducted in a stainless-steel milling jar (SUS420J2, 1.5 mL) with one stainless-steel ball (SUS420J2, diameter: 5 mm) using a Retsch MM400 mill at 30 Hz. First, we conducted the mechanochemical palladium-catalyzed cross-coupling between a solid aryl bromide (**1a**) and diphenylamine (**1b**) in the presence of tri-*tert*-butylphosphonium tetrafluoroborate (*t*-Bu₃P•HBF₄), which is an air-stable precursor of *t*-Bu₃P. Although the use of *t*-Bu₃P efficiently promoted the C–N cross-coupling to form **3a** in high yield (87%) in our previous study,[33] the reaction using *t*-Bu₃P•HBF₄ showed poor conversion and furnished **3a** in 14% yield under similar mechanochemical conditions (Table 1, entry 1). We then turned our attention to air-stable tri(1-adamantyl)phosphine (Ad₃P), which was first synthesized by Carrow and co-workers.[46-50] This ligand is sterically similar to
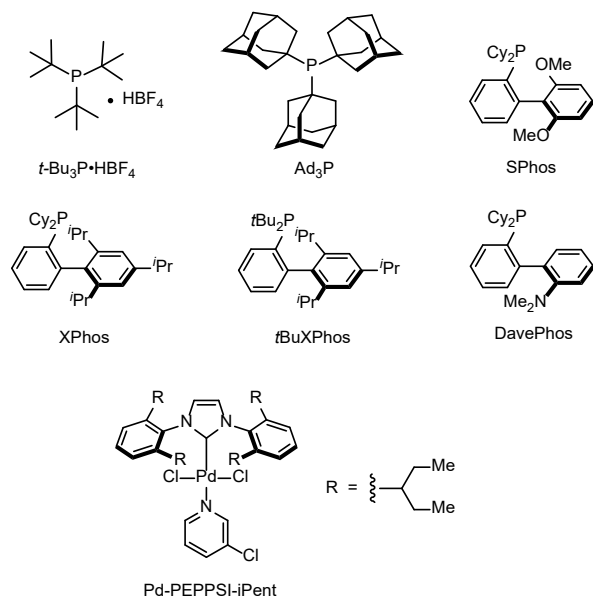
t-Bu3P but has been discovered to be more electron-donating.[46-50] We thus reasoned that Ad₃P might exhibit better catalytic properties than *t*-Bu₃P for the solid-state mechanochemical C–N cross-coupling reactions. Pleasingly, the reaction using Ad₃P proceeded efficiently under mechanochemical conditions to furnish **3a** in good yield (72%; Table 1, entry 2). In contrast, the use of Buchwald ligands, which are commonly employed in solution-based cross-coupling reactions, resulted in poor yields (Table 1, entries 3–6). Notably, the yield was very low when XPhos, which is the optimal ligand under the conditions reported by Su,[31] was used (8%, Table 1, entry 4).[51] No reaction was observed when we used Pd-PEPPSI-iPent, which is the optimal palladium-based catalyst under the conditions reported by Browne[32] (Table 1, entry 7).[51] We also confirmed that the use of 1,5-cod as a LAG additive and supporting ligands for the palladium-based catalyst are essential for this transformation (Table 1, entries 8 and 9).

Table 1. Ligand Effect on the Mechanochemical Solid-State C–N Cross-Coupling Reaction between 1a and 2a in Air.[a]
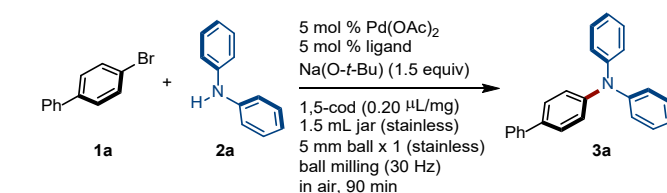
entry	ligand	NMR yield (%)
1	*t*-Bu₃P•HBF₄	14
2	Ad₃P	72
3	SPhos	<5
4	XPhos	8
5	*t*BuXPhos	<5
6	DavePhos	15
7[b]	Pd-PEPPSI-iPent	<5
8[c]	Ad₃P	10
9	none	<5

[a]Conditions: **1a** (0.50 mmol), **2a** (0.50 mmol), Pd(OAc)₂ (0.025 mmol), ligand (0.025 mmol), Na(O-*t*-Bu) (0.75 mmol), and 1,5-cod (0.20 μL/mg) in a stainless-steel ball-milling jar (1.5 mL) with a stainless-steel ball (diameter: 5 mm). Yields were determined by ¹H NMR analysis with CH₂Br₂ as an internal standard. [b]Pd-PEPPSI-iPent (5 mol %) was used instead of Pd(OAc)₂. [c]The reaction was carried out without 1,5-cod.

t-Bu₃P•HBF₄ → $t\text{-Bu}_3\text{P}\cdot\text{HBF}_4$

Ad₃P

SPhos

XPhos

tBuXPhos

DavePhos

$R = $ (isobutyl group, Me / Me)

Pd-PEPPSI-iPent

To further improve the reactivity of the Pd(OAc)₂/Ad₃P system, the effect of the ball-milling conditions on the solid-state C–N cross-coupling reaction was investigated (Table 2). For this purpose, the reaction between **1a** and **2a** was performed in a larger stainless-steel jar (5.0 mL) using one stainless-steel ball (diameter: 10 mm) (Table 2, entry 2). Unfortunately, this resulted in a lower yield (51%) than conducting the reaction in a 1.5 mL stainless-steel milling jar using a 5-mm ball (Table 2, entry 1). The use of an agate or ZrO₂ jar also significantly decreased the yield of **3a** (16% and 5%, respectively; Table 1, entries 3 and 4). In our previous study on the development of a mechanochemical solid-state Suzuki–Miyaura cross-coupling reaction,[28] the product yields have increased significantly with increasing mechanical impact, which suggested that strong mechanical stimulation facilitates uniform dispersion of the reactants, palladium-based catalysts, and bases in the solid-state reaction mixture. With this in mind, we investigated the effect of the number of balls (Table 1, entries 5 and 6). As anticipated, the yield of **3a** improved dramatically when three balls (diameter: 10 mm) were used (>99%; Table 1, entry 6). Under these optimized conditions, the reaction using $t\text{-Bu}_3\text{P}\cdot\text{HBF}_4$ provided **3a** in good yield (67%; Table 1, entry 7), albeit that the use of Ad₃P affords the most satisfactory results.

Table 2. Investigation of the Effect of Varying the Mechanochemical Parameters.[a]

Reaction scheme:
1a (Ph-C₆H₄-Br) + **2a** (H-N(Ph)₂), 5 mol % Pd(OAc)₂, 5 mol % Ad₃P, Na(O-t-Bu) (1.5 equiv), 1,5-cod (0.20 μL/mg), ball milling (30 Hz), in air, 90 min → **3a** (Ph-C₆H₄-N(Ph)₂)

entry	jar/ball material	jar size (mL)	number of balls (mass, g)	ball size (mm)	NMR yield (%)
1	stainless steel (SUS420J2)	1.5	1 (0.5 g × 1)	5	72
2	stainless steel (SUS420J2)	5.0	1 (4.0 g × 1)	10	51
3	Agate	5.0	1 (1.4 g × 1)	10	16
4	ZrO₂	10	2 (3.0 × 2 g)	10	5
5	stainless steel (SUS420J2)	5.0	2 (4.0 × 2 g)	10	82
6	**stainless steel (SUS420J2)**	**5.0**	**3 (4.0 × 3 g)**	**10**	**>99**
7[b]	stainless steel (SUS420J2)	5.0	3 (4.0 × 3 g)	10	67

[a]Conditions: **1a** (0.50 mmol), **2a** (0.50 mmol), Pd(OAc)₂ (0.025 mmol), Ad₃P (0.025 mmol), Na(O-t-Bu) (0.75 mmol), and 1,5-cod (0.20 μL/mg). [b]t-Bu₃P•HBF₄ was used as the ligand instead of Ad₃P.

With the optimized conditions in hand, we explored the scope and limitations of aryl bromides in the present solid-state C–N cross-coupling reaction system in air (Table 3). This reaction is applicable to a wide range of substrates and permits the synthesis of various triarylamines that contain large polycyclic hydrocarbon cores. For example, naphthalene, pyrene, triphenylene, fluorene, acenaphthene, phenanthrene, and anthracene efficiently formed the corresponding triarylamines (**3a–3i**) in moderate to high yields. Unfortunately, the present catalytic system is very sensitive to steric hindrance in the aryl halides, i.e., substrates such as 1-bromonaphthalene derivatives (**1g** and **1h**) and 9-boromoanthracene (**1i**) provided the corresponding coupling products (**3g–3i**) in low yield (12–37%). Triarylamines that contain internal alkyne (**3j**) and stilbene (**3k**) moieties were synthesized in good yield (84% and 80%, respectively) under the optimized conditions. Unfortunately, the reaction between 2-chloronaphthalene (**1b'**) and diphenylamine (**2a**) afforded the product (**3b**) in very low yield (8%).

Next, we explored various amine nucleophiles (Table 3). Both bis(4-methylphenyl)amine (**2b**) and bis(4-methoxyphenyl)amine (**2c**) were reacted with **1c** under the optimized conditions to form the desired products (**3l** and **3m**) in high yield (87% and 93%, respectively). Diarylamines that contain a naphthyl group (**2d** and **2e**) are also compatible under the applied conditions (64% and 60% yield, respectively). Finally, this method was applied to boron-substituted or ketone-containing substrates to form the corresponding functionalized triarylamines (**3p** and **3q**) in 42% and 96% yield, respectively.

Table 3. Substrate Scope.[a]

Conditions: **1** (0.50 mmol), **2** (0.50 mmol), Pd(OAc)₂, 5 mol% Ad₃P, Na(O-t-Bu) (1.5 equiv), 1,5-cod (0.20 µL/mg), 5 mL jar (stainless), 10 mm ball x 3 (stainless), ball milling (30 Hz), in air, 90 min

3a 87% yield
3b 96% yield (8% yield)[b]
3c 80% yield
3d 83% yield
3e 73% yield
3f 93% yield
3g 37% yield
3h 30% yield
3i 12% yield
3j 84% yield
3k 80% yield
3l 87% yield
3m 93% yield
3n 64% yield
3o 60% yield
3p 42% yield
3q 96% yield
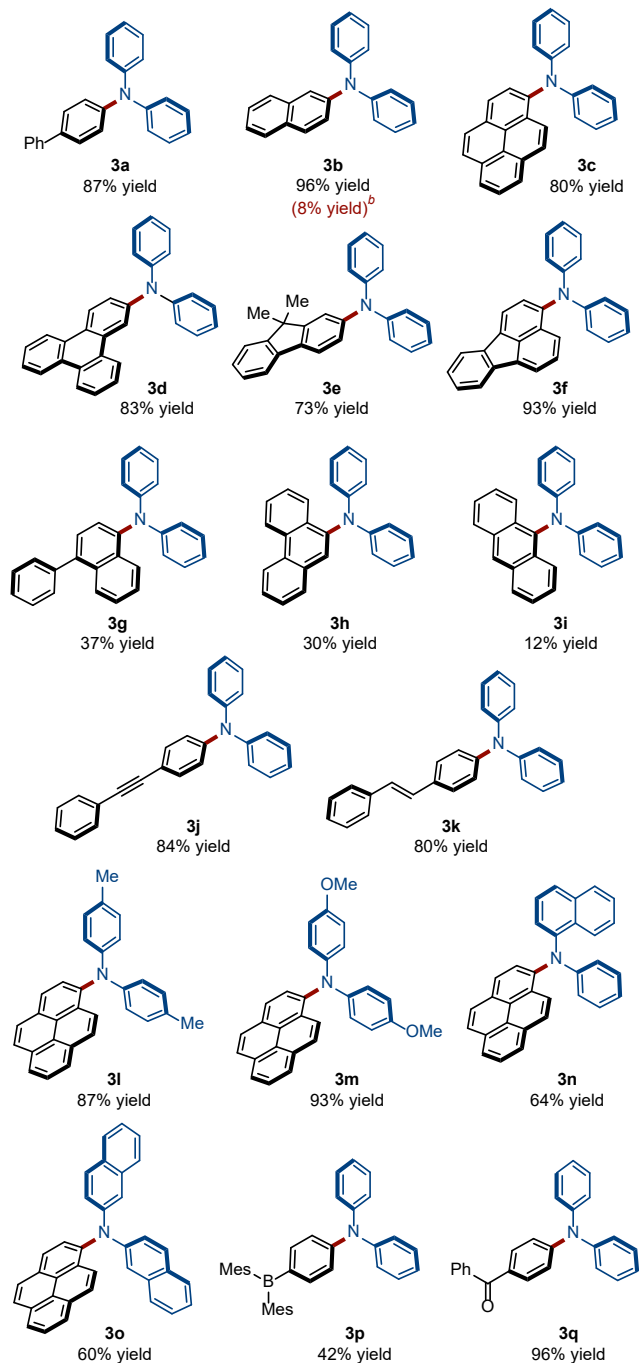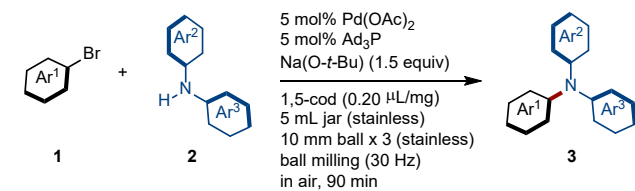
[a]Conditions: **1** (0.50 mmol), **2** (0.50 mmol), Pd(OAc)₂ (0.025 mmol), Ad₃P (0.025 mmol), Na(O-t-Bu) (0.75 mmol), and 1,5-cod (0.20 µL/mg) in a stainless-steel ball-milling jar (5 mL) with three stainless-steel balls (10 mm). [b]2-Chloronaphthalene (**1b'**) was used instead of **1b**.

To demonstrate the practical utility of this protocol, we investigated the gram-scale synthesis of triarylamines under mechanochemical conditions in air (Scheme 2). The reaction of **1b** with **2a** was conducted on a 4.5 mmol scale

under the modified mechanochemical conditions using a stainless-steel ball milling jar (10 mL) with three stainless-steel balls (diameter: 10 mm). Pleasingly, the gram-scale reaction proceeded smoothly, providing **3b** in 80% yield. This result demonstrates the potential utility of the present mechanochemical protocol for the large-scale preparation of valuable synthetic targets in air.

Scheme 2. Gram-scale Synthesis.
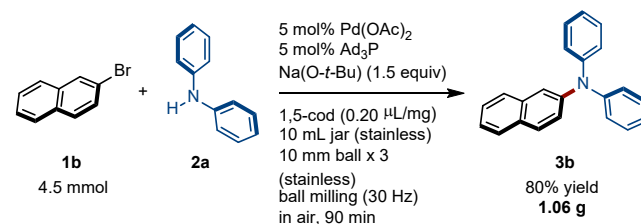
To demonstrate the applicability of the developed mechanochemical conditions to the synthesis of triarylamine-based organic materials, we synthesized hole-transporting materials for organic light-emitting diodes (Scheme 3).[42-44, 52,53] Under the optimized conditions, the reaction of 4,4'-dibromo-1,1'-biphenyl (**1o**) and **2d** afforded the targeted triarylamine (**3r**) in excellent yield (96%).[52,53] This result suggests outstanding potential of this solid-state C–N coupling reaction for the efficient preparation of materials-science-oriented nitrogen-containing polyaromatic compounds in an environmentally friendly manner.

Scheme 3. Mechanochemical Synthesis of a Hole-transporting Material.
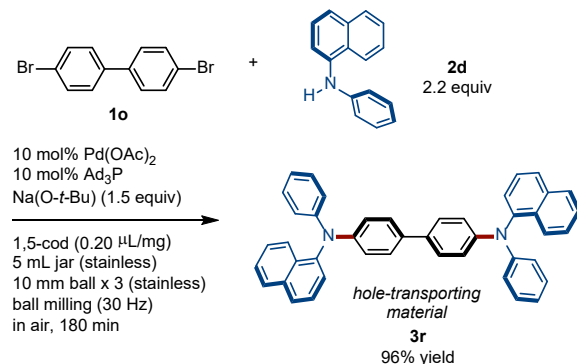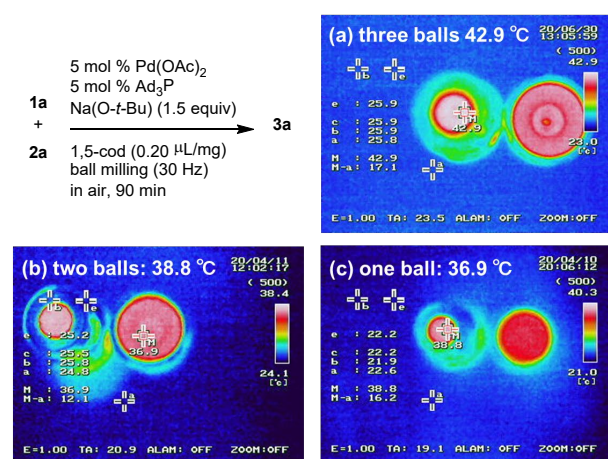

To determine the mechanism by which increasing the number of balls facilitates the solid-state Pd(OAc)₂/Ad₃P-catalyzed C–N coupling reaction, we conducted preliminary mechanistic studies (Figure 1). First, in order to investigate whether the acceleration effect originates the frictional heat during ball milling, the temperature inside the milling jar during the mechanochemical coupling reaction of **1a** and **2a** was measured using thermography immediately after opening the jar (Figure 1A). The thermography images showed that the temperature after the coupling reactions under the optimized conditions using different numbers of 10-mm balls was ~40 °C (<±3 °C) in all cases, which discounts the possibility of a thermally induced acceleration due to heat generated by the ball milling.

Subsequently, we measured the kinetics of reactions with different numbers of balls (Figure S2–S4). As periodic sampling of the reaction runs requires stopping the mill

4

and opening the jar, each data point was obtained from an individual reaction. The reaction using three balls (diameter: 10 mm) did not proceed during the first 30 min but was subsequently rapidly completed within 90 min (Figure S2). Unfortunately, the reactions using one or two balls were not reproducible, impeding a kinetic analysis (Figure S3 and S4). At this point, we think that this irreproducibility is due to the poor mixing efficiency of the solid-state reaction mixture under these conditions. To test this hypothesis, the surface morphology of the milled **2a** was analyzed using scanning electron microscopy (SEM) (Figure 1B). The SEM image of commercially available **2a** shows fine particles with a diameter range of <500 μm [Figure 1B(a)]. The distortion of the particles became more pronounced with increasing number of balls under the applied mechanochemical conditions [Figure 1B(b)–1B(d)]. This result implies that increasing the number of balls increases the surface area of the reactant by crushing it into finer particles, thus facilitating the solid-state coupling reactions and providing good reproducibility.
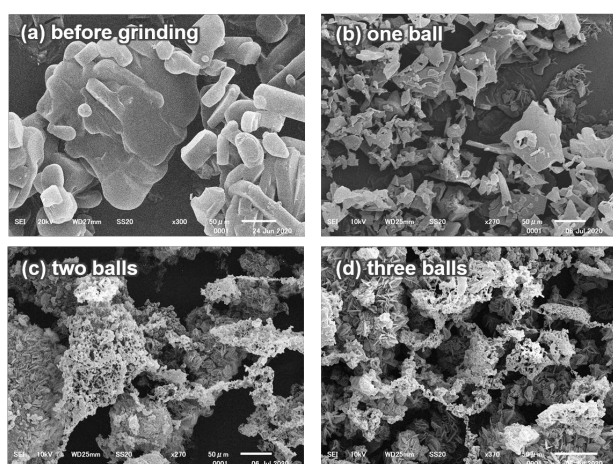

Figure 1. Mechanistic Studies. (A) Thermographic images of the milling jar after the solid-state cross-coupling reactions between **1a** and **2a** under the optimized conditions. (B) SEM images of **2a** (a) before ball milling and after ball milling (30 Hz) using (b) one ball, (c) two balls, (d) or three balls (ball diameter: 10 mm) for 60 min. Scale bars in the SEM images (bottom left): 50 μm.

In summary, we have developed a glove-box-and-Schlenk-line-free protocol for the solid-state C–N cross-coupling reactions catalyzed by the Pd(OAc)$_2$/Ad$_3$P system. The present mechanochemical cross-coupling reactions can be conducted in the solid-state in air and does not require glove-box and Schlenk-line techniques. The present approach would therefore provide a practical and sustainable method to complement conventional solution-based palladium-catalyzed C–N bond-forming reactions for the synthesis of valuable triarylamine-based materials.

ASSOCIATED CONTENT

Supporting Information

Experimental procedures and characterization of new compounds are provided in the Supporting Information. This material is available free of charge via the Internet at http://pubs.acs.org.

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Notes

The authors declare no competing financial interest.

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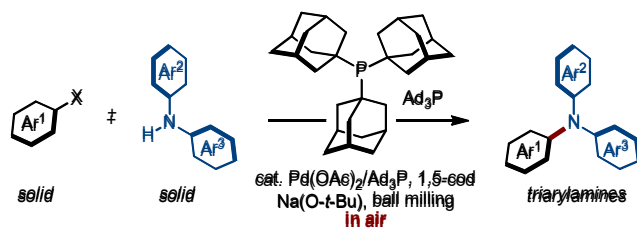
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TOC figure:

solid ‡ *solid*   cat. Pd(OAc)$_2$/Ad$_3$P, 1,5-cod   *triarylamines*
Na(O-*t*-Bu), ball milling
in air

• *glove-box and Schlenk-line-free protocol*   • *general and scalable*   • *mechanistic studies*

Synopsis: The development of operationally simple mechanochemical solid-state C–N cross-coupling reactions in air is reported.