An efficient method for the synthesis of amides via Pd-catalyzed oxidative carbynylation of C(sp³)–H bonds with CO and amines is described. The route efficiently provides substituted phenyl amides from alkanes.

The importance of amides in chemistry and biology is well recognized and, consequently, a variety of methods have been developed for their synthesis, with notable examples including the Schmidt, Schotten–Baumann and Ugi reactions. These methods are based on the reactions of activated acid derivatives (acid chlorides and anhydrides) or acid/base induced rearrangement reactions. Recently, attention has been devoted to developing new routes to amides that do not require acid or base, but to achieve this goal, relatively expensive starting materials such as aldehydes are required. Metal catalyzed routes enable amides to be generated from starting materials other than carboxylic acids and these reactions are summarized in two excellent review articles. Perhaps the most notable achievement in this regard has been the direct catalytic conversion of alcohols and amines into amides.

The synthesis of amides involving transition metal catalyzed C–X (X = H, Br, I etc.) bond functionalization followed by carbynylation with CO has received considerable attention as it has a high atom economy. Seminal research includes the catalytic aminocarbonylation of alk-1-ynes and the application of a homogeneous PdCl₂–PPh₃ catalytic system for direct oxidative aminocarbonylation using CO and oxygen under basic conditions. The catalytic aminocarbonylation of alkenes using a Co/C catalyst has also been reported, and palladium catalyzed carbynylation reactions of aryl bromides have been used to prepare benzamides from aryl bromides at atmospheric pressure. Other notable developments include transition metal-free alkoxy-carbonylation of aryl halides, and the synthesis of amides by the activation of aromatic C–H bonds.

Palladium-phosphine complexes are widely used to catalyze the formation of carbon–carbon, carbon–nitrogen and carbon–oxygen bonds. Bis-phosphine ligands are particularly useful in these reactions and the influence of the ligand bite-angle on C–C and C–X bond forming cross coupling reactions has been reviewed. The wide bite-angle bis-phosphine, Xantphos, has found a number of important uses. Notably, Huang and co-workers reported the synthesis of esters from alkanes using a PdCl₂–Xantphos catalyst in the presence of BuOOBu. Azidocarbonylation reactions may also be catalyzed by a Pd₂dba₂–Xantphos system and intermolecular amidation of aryl halides using Pd(OAc)₂–Xantphos has also been reported. Pd-catalyzed direct oxidative carbynylation of allylic C–H bonds with carbon monoxide has also been reported. We found that PdCl₂ combined with various bis-phosphines including Xantphos, Nixantphos, (±)Binao or (R)-Phanephos catalyze the formation of amides in the absence of acid or base, by the direct functionalization of C(sp³)–H bonds with subsequent CO insertion – the outcome of these studies is described herein.

The viability of the reaction was explored with toluene 1 and aniline 2 as substrates under CO (50 atm) using various PdCl₂-based catalysts due to their excellent performance in carbynylation reactions (Scheme 1, Table 1). High yields of the desired product (compound 3 in Scheme 1) are obtained with the wide bite-angle bis-phosphines, Xantphos, Nixantphos, or (R)-Phanephos and also with (±)-Binao (Table 1, entries 9–12 and 16–19). The yield of 3 is very low in the absence of a ligand co-catalyst (Table 1, entries 1 and 2) or in the presence of other bis-phosphines and mono-phosphate ligands (Table 1, entries 3–8). The influence of the bis-phosphine bite-angle is apparent (Table 1, entries 3–12), with ligands with bite-angles >90° having very low yields.

Scheme 1 Reaction of toluene, aniline and CO to afford N2-diphenylacetamide.
Optimization of reaction conditions for the reaction of toluene and aniline with CO

| Entry | $T$ (°C) | Ligand | Bite angle, $\beta$ | Oxidant | Yield of 3 (%) |
|-------|----------|--------|-------------------|---------|----------------|
| 1     | 100      | Xantphos | —      | DTBP     | 0.4            |
| 2     | 100      | Xantphos | —      | DTBP     | 0              |
| 3     | 100      | Dppbe   | $82^{22a}$ | DTBP     | 0.9            |
| 4     | 100      | Triphos | $84^{23}$  | DTBP     | 2              |
| 5     | 100      | PPh$_3$ | —      | DTBP     | 10             |
| 6     | 100      | Dpppe   | $86^{24}$ | DTBP     | 0              |
| 7     | 100      | Tpp     | —      | DTBP     | 16             |
| 8     | 100      | Dppf    | $99^{25}$ | DTBP     | 3              |
| 9     | 100      | Xantphos| $102^{26}$| DTBP     | 34             |
| 10    | 100      | Xantphos| $102^{2}$ | DTBP     | 36             |
| 11    | 100      | (±)-Binap | $93^{25}$ | DTBP     | 31             |
| 12    | 100      | (R)-Phanephos | $101^{26}$ | DTBP     | 46             |
| 13    | 100      | Xantphos | $102^{2}$ | DTBP     | 21             |
| 14    | 100      | Xantphos | $102^{2}$ | DTBP     | 30             |
| 15    | 100      | Xantphos | $102^{2}$ | DTBP     | 20             |
| 16    | 125      | Xantphos | $102^{2}$ | DTBP     | 66             |
| 17    | 125      | Xantphos | $102^{2}$ | DTBP     | 54             |
| 18    | 125      | (±)-Binap | $93^{25}$ | DTBP     | 62             |
| 19    | 125      | (R)-Phanephos | $101^{26}$ | DTBP     | 68             |
| 20    | 125      | Xantphos | $102^{2}$ | DTBP     | 48             |
| 21    | 125      | Xantphos | $102^{2}$ | DTBP     | 0              |
| 22    | 125      | Xantphos | $102^{2}$ | Ag$_2$O  | 0              |
| 23    | 125      | Xantphos | $102^{2}$ | AgOAc    | 0              |
| 24    | 125      | Xantphos | $102^{2}$ | H$_2$O$_2$| 0.3            |
| 25    | 125      | Xantphos | $102^{2}$ | K$_2$S$_2$O$_4$ | 7   |
| 26    | 125      | Xantphos | $102^{2}$ | DTBP     | 3              |
| 27    | 125      | Xantphos | $102^{2}$ | DTBP     | 0.4            |

Dppbe = 1,2-bis(diphenylphosphino)benzene, Tapp = tris(o-methoxyphenyl)-phosphine, Dppf = 1,1'-bis(diphenylphosphino)ferrocene, Dpppe = 1,2-bis(diphenylphosphino)ethane, DTBP = Bu$_4$O$_2$Bu. Reaction conditions: 1 (15 ml), 2 (1 mmol), PdCl$_2$ (5 mol% based on aniline), ligand (0.06 mmol), oxidant (1.2 mmol), CO (50 atm), 125 °C, 24 h. $a$ Without PdCl$_2$, $b$ CO (30 atm). $c$ CO (40 atm). $d$ With H$_2$O (15 ml), 1 (15 mmol), and 2 (1 mmol). $e$ Pd$_2$(dba)$_3$ (0.05 mmol). $f$ Pd(PPh$_3$)$_2$ (0.05 mmol). $g$ Yields were determined by GC analysis relative to aniline with $n$-decane as an internal standard. $h$ The bite angle given corresponds to that in Dpppe/PdBr$_2$. $i$ The structure of Xantphos is similar to Xanthos and it is therefore assumed that their bite-angles are the same.

generally exhibiting better activity (the exception being dppf). Moreover, in the absence of an oxidant or in the presence of a weak oxidant, i.e. Ag$_2$O, compound 3 is not obtained (Table 1, entries 21 and 22). Aniline reacts more favourably with itself to afford 1,3-diphenylurea when CO$_2$ is employed as the oxidant or H$_2$O as solvent (Table 1, entry 16). Under these conditions a similar yield (68%) is obtained with the (R)-Phanephos ligand (Table 1, entry 19).

Substrate scope of the Pd-catalyzed amide formation with aniline

| Entry | RH | Yield$^a$ (%) |
|-------|----|---------------|
| 1     |   | 0             |
| 2     |   | Branched (46)$^f$ Linear (13)$^f$ |
| 3     |   | Branched (53)$^f$ Linear (12)$^f$ |
| 4     |   | 43 (39)       |
| 5     |   | 60            |
| 6     |   | 66 (62)       |
| 7     |   | 68            |
| 8     |   | 55 (51), 38$^b$ |
| 9     |   | 57 (51), 31$^b$ |
| 10    |   | Trace         |

Reaction conditions: RH (15 ml), 2 (1 mmol), PdCl$_2$ (5 mol% based on aniline), Xantphos (0.06 mmol), DTBP (1.2 mmol), CO (50 atm), 125 °C, 24 h. $a$ Yields were determined by GC analysis relative to aniline with $n$-decane as an internal standard (isolated yield in parentheses). $b$ With (R)-Phanephos (0.06 mmol). $c$ N,2-diphenylpropanamide. $d$ N,3-diphenylpropanamide.

Xanthos or (R)-Phanephos and DTBP as the oxidant (Table 2). Products resulting from carbonylation of C(sp$^3$)-H (aromatic) bonds are not observed (Table 2, entry 1). Cyclohexane reacts in the presence of DTBP to afford the corresponding amide in reasonable yield (Table 2, entries 4 and 5), albeit lower than the yield of the product obtained with toluene (Table 2, entries 6 and 7), presumably due to the higher bond dissociation energy of the C–H bond in cyclohexane compared to toluene. With diphenylmethane the main product is 1,3-diphenylurea, presumably due to steric hindrance. Presumably the yield of the branched product is higher than the linear product using ethylenbenzene as the substrate for the same reason (Table 2, entries 2 and 3). Moreover, electron withdrawing –F and –Cl substituents in the para-position favor the reaction; however, for the carbonylation of toluene, ethylenbenzene and cyclohexane, the ligand (R)-Phanephos is superior to Xanthos, whereas for substrates with electron withdrawing –F and –Cl substituents in the para-position, Xanthos is superior (Table 2, entries 2–11). These combined data confirm that the yield of the desired product is related, at least in part, to the C(sp$^3$)-H bond dissociation energies. The PdCl$_2$-Xanthos catalyst tolerates anilines with electron withdrawing or donating substituents (Scheme 2 and Table 3), although electron withdrawing groups are more favorable for this reaction. The system is, unfortunately, inactive with allylamines.

Scheme 2 Reaction of toluene and substituted anilines with CO.
Table 3 Influence of substituents attached to aniline on the carbonylation reaction

| Entry | R            | Yield of 3′-a (%) |
|-------|--------------|------------------|
| 1     | H            | 62               |
| 2     | 3-NO₂        | 64               |
| 3     | 3-CN         | 57               |
| 4     | 4-CN         | 69               |
| 5     | 3-F          | 54               |
| 6     | 4-F          | 66               |
| 7     | 3-Cl         | 64               |
| 8     | 4-Cl         | 66               |
| 9     | 3,5-CF₃      | 78               |
| 10    | 3-Me         | 51               |
| 11    | 4-Me         | 48               |
| 12    | 3-OMe        | 61               |
| 13    | 4-OMe        | 32               |
| 14    | 2,4,6-Me     | 8                |

Reaction conditions: 1 (15 ml), 2′ (1 mmol), PdCl₂ [5 mol% based on 2′], Xantphos (1.20 mmol) DTBP (1.2 mmol), CO (50 atm), 125 °C, 24 h. *Isolated yield based on the amine.

Scheme 3 The proposed reaction mechanism.

The full mechanistic details of this transformation have not been determined, however, in the presence of the radical scavenger TEMPO the reaction is completely suppressed, indicative of a radical process, which is similar to the one proposed by Huang and co-workers for the formation of esters from alkanes and alcohols using a similar catalytic system. A plausible reaction mechanism is shown in Scheme 3. In the presence of a ligand, sequential oxidation of the Pd(0) bis-phosphine catalyst generated in situ with the anilino and benzyl radicals produced in the presence of DTBP leads to the formation of intermediate (B). Subsequent insertion of CO gives intermediate (C) which can undergo reductive elimination to afford the final product. The concentration of aniline strongly influences the yield of the product, i.e. at high concentrations the yield of 1,3-diphenylurea is increased (see ESI), presumably because the aniline can more easily react with itself and CO to form 1,3-diphenylurea.

ESI-MS was used to analyze the reaction and a peak that may be tentatively assigned to [(Xantphos)Pd(CH₃)₂Ph]+ (Fig. S1, ESI§) was observed. This species could be derived from either B or C (see ESI§ for further details).

In summary, a convenient and efficient method for the synthesis of amides via Pd-catalyzed oxidative carbonylation of C-H bonds with CO has been devised. The method represents a practical and efficient approach for the synthesis of substituted phenyl amides from simple alkanes.

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