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Direct N-formylation of nitroarenes with CO₂[†]

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Herein we describe a straightforward *N*-formylation of nitroarenes with CO_2 to access *N*-aryl formamides exclusively in the presence of iron and hydrosilane as additives. This protocol showcases a good tolerance of a wide range of nitroarenes and nitroheteroarenes.

Carbon dioxide (CO_2) constitutes an ideal C1 source in organic synthesis owing to its abundance and stability.¹ In particular, CO₂ serves as the best-suited substitute of more reactive C1 building blocks, such as carbon monoxide (CO), methyl iodide, and phosgene.² Despite its high activation barrier and thermodynamic stability,² CO₂ has been widely exploited for decades to access a diverse array of functionalized compounds via carboxylation,³ carbonylation,⁴ or C-H bond construction.⁵ In this context, the incorporation of amines and CO₂ towards synthesis of value-added nitrogen-containing compounds,⁶ especially *N*-formylation of amines with CO_{2} ,⁷⁻¹³ is among the most important chemical transformations due to the versatility of nitrogen-based compounds in academia and industry. Taking the advantage of commercially available hydrosilanes as activating and hydrogenating agents along with various catalysts, such as organic superbases,⁷ carbenes,⁸ ionic liquids and organic salts,⁹ inorganic salts,¹⁰ polar aprotic solvents,¹¹ diazophospholene,¹² and $B(C_6F_5)_3$,¹³ a variety of formamides can be accessed based on amines and CO₂ (Scheme 1a). Additionally, the Ding¹⁴ and Bernskoetter¹⁵ groups described the seminal Ru- and Fe-catalyzed N-formylation of alkylamines based on the CO_2/H_2 system, respectively (Scheme 1b). On the other hand, nitroarenes prove to be reliable aminating agents

for direct synthesis of amines¹⁶ and amides¹⁷ without preforming the more reactive anilines via hydrogenation. Recently, Beller,^{17a} Driver,^{17b} Hu,^{17c} Mankad,^{17f} Wu,^{17d,e} as well as our group^{17g} disclosed the aminocarbonylation reactions to access aryl amides, a class of important compounds in chemical, pharmaceutical, and agrochemical industries, based on nitroarenes and CO or its surrogates. To our knowledge, the merger of CO₂ and nitroarenes for straightforward amide synthesis remains unexplored. We envisioned that the use of suitable and compatible reductants and activating agents could induce the integration of both CO2 and nitroarenes for amide formation. Herein, we unveil the direct N-formylation of nitroarenes with CO_2 in the presence of iron powder and hydrosilane additives (Scheme 1c). This alternative and complementary method would open a conceptually novel avenue for more step-economic and expedient access to formamides without the need of conventional anilines.



Scheme 1 Developments of *N*-formylation based on CO₂.

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We commenced the study on N-formylation of CO₂ using 4-nitroanisole (1a) as the model substrate (Table 1). In the presence of 3 equivalents of Zn powder as reductant and 3 equivalents of phenylsilane as additive, 1a reacted with CO_2 under balloon pressure in dimethylformamide (DMF) solvent at 110 °C to give N-4-methoxyphenyl formamide 2a exclusively in 49% yield (entry 1). In the absence of Zn or hydrosilane, no formulation took place, indicating that both metal reductant and hydrosilane are essential for reaction (entries 2 and 3). When triethoxysilane was used in place of phenylsilane, formylation underwent at 135 °C to offer 2a in 56% yield (entry 4). Further, when Fe powder was employed instead of Zn, 2a was also obtained in 51% yield (entry 5). Noteworthily, the yield could be enhanced to 63% when 10 mol% of KI was added as additive in the Fe-mediated reaction (entry 6). Subsequently, 1,1,3,3-tetramethyldisiloxane (TMDSO) was found to be the superior additive, giving 2a in 75% yield (entries 7-9). Intriguingly, the use of Fe nano powder allowed the reduction of loadings of both Fe and TMDSO to 2 equivalents in association with the yield enhancement to 83% yield (entries 10-12). Such promoting effect would be attributed to the more efficient reduction process provided by the more soluble and finely powdered Fe. In the presence of other polar aprotic solvents such as dimethylacetamide (DMA) and N-methylpyrrolidine (NMP), 2a was formed in similar yields without other side products (entries 13 and 14). Furthermore, no formamide was formed when CO₂ was omitted (entry 15). These control experiments suggested that the carbonyl group of 2a originates from CO₂ rather than solvent.

The optimized protocol (Table 1, entry 11) proved to be general in *N*-formylation of nitroarenes (Scheme 2). Electron-rich (**1a–1f**), electron-neutral (**1g**), and electron-deficient (**1h–1m**) nitroarenes

Table 1Optimization of N-formylation of nitroarenes with CO_2^a					
MeO 1a MeO 1a Me					
Entry	Metal (equiv.)	Hydrosilane (equiv.)	Additive (mol%)	Solvent	Yield ⁱ (%)
1 ^c 2 ^c 3 ^c 4 5	Zn (3) None Zn (3) Zn (3) Fe (3) Fe (3)	PhSiH ₃ (3) PhSiH ₃ (3) None (EtO) ₃ SiH (3) (EtO) ₃ SiH (3) (EtO) ₂ SiH (3)	None None None None None KL (10)	DMF DMF DMF DMF DMF DMF	49 0 56 51 63
7 8 9 10 ^d	Fe (3) Fe (3) Fe (3) Fe (3)	Me(EtO) ₂ SiH (3) PMHS (3) TMDSO (3) TMDSO (2)	KI (10) KI (10) KI (10) KI (10)	DMF DMF DMF DMF	25 32 75 83
11^{d} 12^{d} 13^{d} 14^{d} $15^{d,e}$	Fe (2) Fe (1) Fe (2) Fe (2) Fe (2)	TMDSO (2) TMDSO (2) TMDSO (2) TMDSO (2) TMDSO (2)	KI (10) KI (10) KI (10) KI (10) KI (10)	DMF DMF DMA NMP DMF	83 35 83 75

^{*a*} General procedure: **1a** (0.5 mmol), CO₂ (balloon), metal powder (1–1.5 mmol), hydrosilane (1–1.5 mmol), additive (10 mol%), solvent (1.5 mL), 135 °C, 28 h. ^{*b*} Isolated yield. ^{*c*} Reaction conducted at 110 °C. ^{*d*} Fe nano powder used. ^{*e*} Argon atmosphere without CO₂.

all reacted with CO₂ to form the corresponding N-aryl formamides in moderate to high yields. Additionally, both mono- (1a-2i, 1l) and di-substituted (1j, 1k, 1m-1o) nitroarenes could be incorporated into the formamide products 2a-2o. The sterically encumbered 1-nitro-2,4-dimethylbenzene (10) was also suitable reaction partner. Notably, a wide range of nitro-substituted heterocycles could be utilized for formylation, including benzodioxole (1p), pyrrole (1q), pyridine (1r), N-alkylated indole (1s, 1t), benzofuran (1u), benzothiophene (1v), and benzoxazole (1w), delivering the corresponding N-heteroaryl formamides 2p-2w in 43-85% yields. Noteworthily, the employment of 3 equivalents of Fe metal (Table 1, entry 10) could improve remarkably the formylation of electron-rich nitro(hetero)arenes 1c, 1o and 1s, affording the corresponding formamides 2c, 2o and 2s in 80%, 60% and 63% vields, respectively. On the contrary, the reactions of electrondeficient nitroarenes with the additional equivalent of Fe led to substantial diminishment of yields. Gratifyingly, by adding FeCl₂ (2 equiv.) and slightly increasing the loadings of TMDSO and KI, electron-deficient nitroarenes such as 1j-1m could react more efficiently to afford the formamides in higher yields (48-68%). This protocol was also amenable to multigram synthesis of 2a in synthetically useful yield.

In the course of reaction, nitroarene can be transformed to nitrosoarene, *N*-aryl hydroxylamine, azoxyarene, azoarene, 1,2-diaryl hydrazine, and aniline under reductive conditions.¹⁸ Thus, the reactions of these nitrogen-based species under otherwise



Scheme 2 Substrate scope of nitroarenes in *N*-formylation.^a ^a Reaction conditions: nitroarene (0.5 mmol), CO₂ (balloon), nano Fe (1 mmol), TMDSO (1 mmol), KI (0.05 mmol), DMF (1.5 ml), 135 °C, 28 h. Isolated yields were shown. ^b 3 equiv. of Fe. ^c Modified conditions: nitroarene (1 equiv.), CO₂ (balloon), nano Fe (2 equiv.), FeCl₂ (2 equiv.), TMDSO (3 mmol), KI (30 mol%), DMF (3 ml), 135 °C, 28 h.



identical conditions were examined to probe their roles in formylation (Scheme 3a). Nitrosobenzene (3a), N-phenyl hydroxylamine (3b), azoxybenzene (3c) and azobenzene (3d) all reacted to give formamide 2g in 49-64% yields. Particularly, diphenyl hydrazine (3e) reacted most efficiently to afford 2g in 70% yield. Presumably, nitrobenzene is sequentially reduced to 3a, 3b, 3c, 3d, and finally 3e,¹⁸ which acts as the ultimate intermediate to participate in the formylation reaction. Aniline 3f also reacted to give 2g in 44% yield, suggesting that it is likely the minor intermediate to induce the formylation. Furthermore, when Fe(0) was omitted, the addition of Fe(II) salt, which is a viable oxidized Fe species, could enable the formylation of nitroarene 1a in the presence of excess TMDSO to give formamide 2a in 45% yield. We surmised that hydrosilane can also act as a reductant and a hydrogen source for the reduction of nitroarene *via* the action of Fe-hydride species,^{16a,b} apart from its conventional application as CO₂-activating agent.^{6h}

Based on the experimental results, we proposed a plausible mechanism of the formylation protocol (Scheme 4). Nitroarene 1 is sequentially reduced to 1,2-diaryl hydrazine via the intermediacy of nitrosoarene, N-aryl hydroxylamine, azoxyarene, and/or azoarene¹⁸ (pathway i). 1,2-Diaryl hydrazine then undergoes facile homolysis under heating to give aminyl radicals,^{17g} which are then reduced to amide ion ArNH⁻ (pathways ii and iii). Meanwhile, nitroarene can be reduced to amide dianion ArN²⁻ followed by ArNH⁻, whereas Fe(0) is oxidized to Fe₃O₄¹⁹ (pathway iv and v). A small amount of aniline can also be generated (pathway vi). In the reduction processes, Fe(0) is likely the major reducing agent, while TMDSO likely acts as both mild reducing agent and hydrogenating agent. Owing to the strong nucleophilicity of ArNH⁻ and ArN²⁻, they likely activate both CO₂ and TMDSO via the transition state TS-1, in which the carbamate ion formed further activates TMDSO to trigger the hydride attack to another CO_2 (pathway vii). Such transition state is regarded as the lowest energy pathway for CO₂ activation.^{6h} Via TS-1, formoxysilane is formed with the concomitant regeneration of amide species (pathway viii). We proposed that iodide ion behaves as a





strong nucleophile in polar aprotic solvent DMF, thereby facilitating the deformylation of formoxysilane to form a more reactive formyl iodide in conjunction with silanolate ion (pathway ix). Finally, ArNH⁻ reacts with both formyl iodide and formoxysilane to deliver the *N*-aryl formamide 2 (pathways x and xi). In the same vein, ArNH²⁻ undergoes nucleophilic substitutions with formyl iodide and formoxysilane to give amidate ion, which furnishes formamide 2 upon acidic workup (pathways xii and xiii). On the other hand, the *N*-formylation based on aniline may proceed but tends to be a minor pathway due to the attenuated nucleophilicity of aniline. The detailed reaction mechanism is subjected to a dedicated study in the future.

Under the standard conditions, electron-rich nitro(hetero) arenes tended to react more efficiently to form higher product yields (\sim 50-85%) than electron-deficient and sensitive groupbearing nitroarenes (\sim 40–60%, Scheme 2). We rationalized that the nitrogen-based intermediates derived from the latter are more unstable, especially in the reductive reaction conditions and in the presence of basic by-products (e.g. iron oxides, silanolate ions), and likely undergo over-reduction and decomposition at varying degrees in the course of reactions. Additionally, an additional equivalent of Fe(0) was found advantageous in the reactions of electron-rich nitro(hetero)arenes but detrimental for electrondeficient ones. We speculated that the contribution of Fe(0)and TMDSO-based reduction steps would vary with respect to the electronic effect of nitroarenes. The reduction of electron-rich nitro(hetero)arenes would mainly rely more on highly reducing Fe(0) to deliver more amide ions. Conversely, the excess Fe(0)would over-reduce the electron-deficient nitroarenes to less

reactive anilines or other side products, but additional milder reduction given by Fe(II)/TMDSO system would in turn be more productive for formation of amide ions.

In conclusion, we have developed an alternative strategy to access formamides *via* direct *N*-formylation of nitroarenes with CO₂. By using commercially available Fe powder and 1,1,3,3-tetramethyldisiloxane as additives, a variety of *N*-aryl and *N*-heteroaryl formamides are synthesized. Mechanistic study suggests that both Fe metal and hydrosilane can reduce nitroarenes to anionic nitrogen-based intermediates, which then activate CO₂ and hydrosilane for subsequent formylation.

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Conflicts of interest

There are no conflicts to declare.

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