Check for updates

Volume 54
Number 1
4 January 2018
Pages 1-112

ChemComm

Chemical Communications

rsc.li/chemcomm

ISSN 1359-7345

ROYAL SOCIETY OF CHEMISTRY

COMMUNICATION
S. J. Connon, M. O. Senge et al.
Conformational control of nonplanar free base porphyrins:
towards bifunctional catalysts of tunable basicity

ROYAL SOCIETY OF CHEMISTRY

rsc.li/chemcomm

ChemComm

COMMUNICATION

Direct Carbonylative Amidation of Benzylic Alcohols with Alkylamines via Palladium Catalyzed C-O Bond Activation

Zhiping Yin,[*a] Heng Li,[a] Xiaowen Qin,[a] Mengyang Liu,[a] Chang-Sheng Wang,[b] Xiao-Feng Wu[*c]

A palladium-catalyzed direct carbonylative amidation that enables the efficient coupling of benzylic alcohols with alkylamines under mild conditions has been developed. Key to the success is the use of pentafluoropyridine as a activating group, which facilitates selective alcohol activation and suppresses undesired side reactions.

Amide bond formation stands as a cornerstone-and a persistent challenge-in synthetic chemistry, particularly in the context of modern atom- and step-economical synthesis.[1] This is especially relevant for medicinally privileged structures such as the benzylamide motifs found in widely prescribed Z-drugs like zolpidem and alpidem, where efficient and modular access to this scaffold is highly desirable.[2] While classical condensation methods relying on pre-activated carboxylic derivatives remain indispensable in both academic and industrial settings, their dependence on stoichiometric coupling agents and the concomitant generation of chemical waste imposes clear limitations in terms of sustainability and operational simplicity. Following the pioneering work on transition-metal-catalyzed carbonylative coupling by R. F. Heck, catalytic carbonylation has emerged as a powerful synthetic platform.[3] Among its most notable advances is the three-component coupling of aryl (pseudo)halides, carbon monoxide, and amines, which offers an efficient and direct route to amide bonds.[4] Despite substantial progress, these methods universally require pre-functionalized electrophilic partners, which introduces additional synthetic steps and constrains the modularity of the approach.

In contrast, alcohols are traditionally employed as nucleophiles or solvents. Realizing the direct carbonylative amidation of alcohols with amines via catalytic CO insertion would represent a paradigm shift-transitioning from the conventional logic of pre-functionalization and coupling to a strategy of direct assembly from simple and abundant feedstocks. This transformation demands both activation of the robust C-O bond and a formal polarity inversion at the alcohol's α-carbon to generate an electrophilic acyl equivalent.[5] Achieving such an inversion with high selectivity under catalytic control is essential to suppress competing pathways, particularly the challenge of differentiating two inherently nucleophilic species within a single catalytic manifold. Our group previously demonstrated that nitroarenes could serve as a nitrogen source in the conversion of alcohols to amides, albeit through a distinct redox-neutral C-N coupling pathway limited to aromatic amines.[6] Despite their ready availability, the direct use of alcohols and aliphatic amines in carbonylative couplings has been hindered by the difficulty of the reduction of alkyl nitro compounds. While transition-metal catalyzed carbonylations of benzyl alcohols to phenylacetic acid derivatives have been known for decades (typically proceeding through halide intermediates),[7] Wu's pioneering work showed that dimethyl carbonate could act as a benign activator, avoiding halides entirely.[8] However, the strategic merger of alcohol C-O activation with aliphatic amines for direct amide synthesis remains an open frontier-one that offers considerable potential for more sustainable and efficient synthetic platforms.

Building on our group's sustained interest in nitrogen-heterocycle synthesis,[9] we herein report a catalytic system that directly addresses this challenge: the deoxygenative carbonylation of benzylic alcohols with aliphatic amines. Using a tailored palladium catalyst, we achieve selective activation of the robust benzylic C-O bond, controlled CO insertion, and efficient interception by the amine nucleophile. This strategy effectively reverses the inherent polarity of the alcohol, enabling its use as a latent electrophile in a direct, one-pot coupling to amides. The method operates under mild conditions and provides a modular and step-economical route to a range of functionally diverse benzylic amides.

Our investigation began with benzyl alcohol (**1a**), pentafluoropyridine, and piperidine (**2a**) as model substrates in acetonitrile under a carbon monoxide atmosphere (5 bar), with Pd(OAc)₂/XantPhos as the catalytic system. Recognizing the crucial role of a base in neutralizing the HF byproduct generated

a. School of Pharmacy, Jiangsu University, Zhenjiang 212013, China, E-mail: zhiping_yin@ujs.edu.cn

b. College of Biotechnology and Pharmaceutical Engineering, Nanjing Tech University, 30 Puzhu Road S., Nanjing 211816, PR China

c. Dalian National Laboratory for Clean Energy, Dalian Institute of Chemical Physics, Chinese Academy of Science, 116023 Dalian, Liaoning, China, E-mail: xwu2020@dicp.ac.cn

during the formation of the key intermediate 4-(benzyloxy)-2,3,5,6-tetrafluoropyridine (**1a'**) from **1a** and pentafluoropyridine, we systematically evaluated a variety of bases (Table 1, entries 1–5). The results indicated that KOH afforded the optimal yield (37%), whereas other tested bases (e.g., NaH, t-BuOK, Et₃N, etc.) provided only trace amounts of product or no reaction. Consequently, KOH was established as the most effective base for this transformation and served as the benchmark condition for subsequent optimization. A preliminary investigation into the loading of KOH revealed that decreasing its amount to 1.5 equivalents led to a decreased yield of 28% (Table 1, entry 6). We then optimized the stoichiometry of the three coupling partners while keeping other conditions fixed. A molar ratio of **1a** : pentafluoropyridine : **2a** = 1.0 : 1.5 : 2.5 provided the best result (56% yield, entry 7), whereas a reversed ratio (2.0 : 1.5 : 1.0) significantly reduced the yield to 21% (entry 8), highlighting the importance of a balanced stoichiometry. Subsequently, a systematic screening of ligands within the palladium catalytic system was conducted. Under otherwise fixed conditions, the influence of various phosphine ligands, including XantPhos and DPEPhos was examined, the properties of ligand were proven crucial (Table 1, entries 9-14). Among them, DCyPFc (1,1'-Bis(dicyclohexylphosphino)ferrocene) delivered the best catalytic performance, affording the product in 86% yield (Table 1, entry 13). Notably, the stability of ligand was not influenced with the presence of pentafluoropyridine which was confirmed by ³¹P NMR. With DCyPFc identified as the optimal ligand, we further evaluated the activity of different palladium catalyst precursors. PdCl₂ demonstrated the highest catalytic efficiency, elevating the yield to 91% (Table 1, entry 17). The target product **3a** was ultimately obtained in an isolated yield of 78%. Finally, solvent screening confirmed acetonitrile as optimal; alternative solvents such as toluene, DMF, and DCE were less effective (see SI, Section 3.5). These optimized conditions (PdCl₂/DCyPFc, KOH, CH₃CN, 5 bar CO) were used for all subsequent substrate-scope investigations.

Table 1. Optimization of reaction conditions[a]
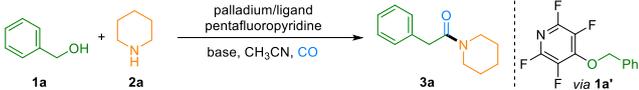
Entry	Catalyst/Ligand	Bite angle	Base	Yield % [b]
1	Pd(OAc)₂/XantPhos	108.11	KOH	37
2	Pd(OAc)₂/XantPhos	108.11	NaOH	26
3	Pd(OAc)₂/XantPhos	108.11	NaH	Trace
4	Pd(OAc)₂/XantPhos	108.11	t-BuOK	8
5	Pd(OAc)₂/XantPhos	108.11	Et₃N	Trace
6[c]	Pd(OAc)₂/XantPhos	108.11	KOH	28
7[d]	Pd(OAc)₂/XantPhos	108.11	KOH	56
8[e]	Pd(OAc)₂/XantPhos	108.11	KOH	21
9[d]	Pd(OAc)₂/DPEPhos	103.93	KOH	63
10[d]	Pd(OAc)₂/N-Xantphos	114.2	KOH	41
11[d]	Pd(OAc)₂/PPh₃	/	KOH	N.R.
12[d]	Pd(OAc)₂/TFP	/	KOH	Trace
13[d]	Pd(OAc)₂/DCyPFc	99.18	KOH	86
14[d]	Pd(OAc)₂/DPPF	99.18	KOH	79
15[d]	Pd(PPh₃)₄/DCyPFc	99.18	KOH	86
16[d]	Pd(PhCN)₂Cl₂/DCyPFc	99.18	KOH	71
17[d]	PdCl₂/DCyPFc	99.18	KOH	91 (78[f])

[a] Reaction conditions: Step 1: 1a (0.1 mmol, 1.0 equivalent, 10 µL), pentafluoropyridine (0.2 mmol, 2.0 equivalent, 22 µL), base 2.5 equivalent, CH₃CN (1 mL), N₂, r.t. and 5h, Step 2: 2a (0.2 mmol, 2.0 equivalent, 20 µL), palladium catalysts (5 mol%), ligand (10 mol%), CO(5 bar), 110 °C and 20 h. N.R. = no reaction. [b] GC yields of mixture **3a** were determined by using dodecane as the internal standard. [c] KOH (8.4 mg, 0.15 mmol, 1.5 equivalent). [d] 1.0 equivalent = 0.1 mmol, 1a : pentafluoropyridine : **2a** = 1.0 : 1.5 : 2.5 eq. [e] 1.0 equivalent = 0.1 mmol, **1a** : pentafluoropyridine : **2a** = 2.0 : 1.5 : 1.0 eq. [f] Isolated yield is given in parenthesis. TFP = Tri-2-furanylphosphine.

Under the optimized conditions, the substrate scope of this carbonylative amidation was systematically examined. As illustrated in Table 2, the reaction exhibited broad compatibility with a wide range of benzylic alcohols. Electron-deficient substrates bearing halogen (F, Cl), trifluoromethyl, or cyano groups reacted effectively, affording the corresponding amides in 20–86% yield, though the trifluoromethyl- and cyano-substituted derivatives exhibited moderate reactivity (Table 2, **3f**, **3g**). The decreased yield was due to the decreased stability of the corresponding palladium intermediate. Electron-rich benzylic alcohols with methyl, *tert*-butyl, methoxy, or phenyl substituents were also well tolerated, delivering products in 34–81% yield (Table 2, **3h**–**3k**). The method further extended to other aromatic systems such as naphthalene and thiophene, which provided moderate yields (33-48%, **3l**, **3m**). It must be noted that trace or no target product were observed for (*E*)-3-phenylprop-2-en-1-ol and pyridin-3-ylmethanol (**3n**), likely due to the inherent strength of their C-O bonds and the potential coordination of the pyridine moiety under the reaction conditions (see SI 2.3 for failed examples, including aliphatic alcohols).

We then evaluated the generality of this transformation with respect to various alkylamines. A wide range of both primary and secondary alkylamines proved to be competent coupling partners, delivering the corresponding amides in 44-96% yield (Table 3, **4a**–**4q**). Cyclic secondary amines of varying ring sizes performed effectively: five- and seven-membered ring analogues afforded the desired amides in 94% and 46% yield, respectively. Similarly, substituted piperidines such as 4-methylpiperidine and morpholine were well tolerated, yielding products in 44-63% (Table 3, **4c**, **4d**). Acyclic secondary amines also participated smoothly. Diethylamine, dipropylamine, and even the more sterically hindered diisopropylamine all provided the corresponding

amides in >60% yield (Table 3, **4e-4g**). Among primary alkylamines, linear analogues including propylamine, butylamine, and isobutylamine reacted with high efficiency (93–94% yield, Table 3, **4h-4j**). Notably, more hindered primary amines such as isopropylamine and tert-butylamine remained viable substrates, furnishing products in 79–89% yield, further highlighting the practical utility of this method (Table 3, **4k, 4l**). The protocol also exhibited excellent compatibility with strained and alicyclic amines, with cyclopropylamine and cyclohexylamine delivering amides in up to 96% yield (Table 3, **4m**, **4n**). Furthermore, various benzylamines and aromatic amines proved to be valuable substrates, providing the corresponding products in 62–70% yield (Table 3, **4o-4q**).
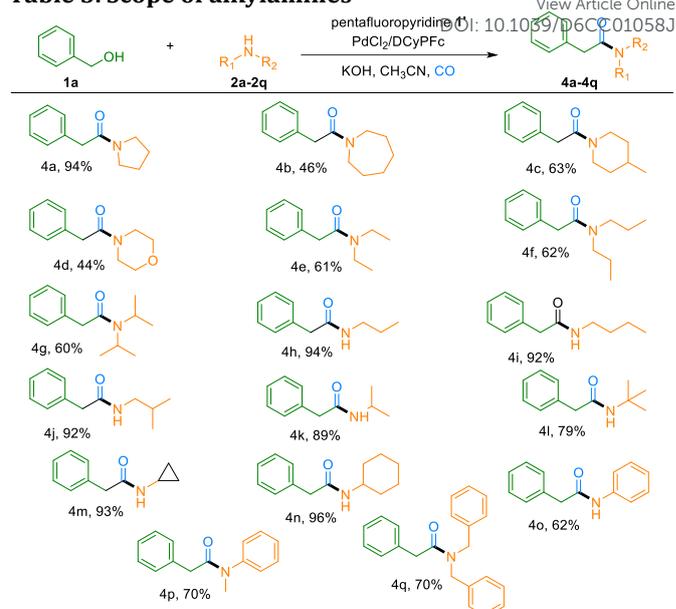
Table 2. Scope of benzylic alcohols [a]
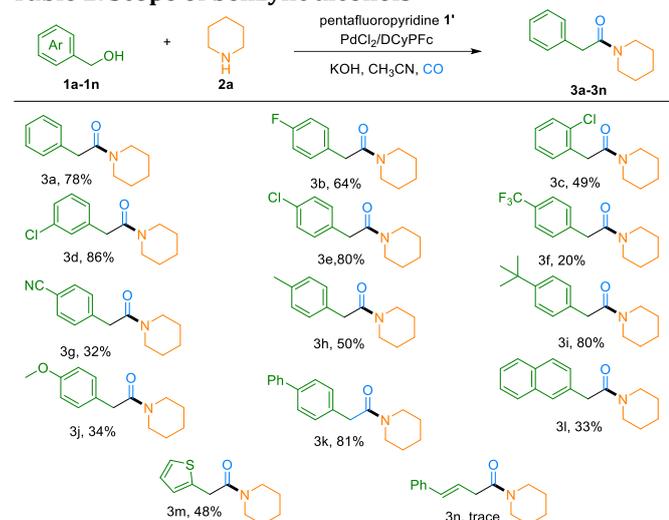
[a] Reaction conditions: Step 1: **1a-1n** (0.2 mmol, 1.0 equivalent), pentafluoropyridine (0.3mmol, 1.5 equivalent, 33 μL), KOH (0.5 mmol, 2.5 equivalent, 28 mg), CH$_3$CN (2 mL), N$_2$, r.t. and 5h, Step 2: **2a** (0.5 mmol, 2.5 equivalent, 49μL,), PdCl$_2$ (5 mol%, 1.8 mg), DCyPFc (10 mol%, 11.6 mg), CO (5 bar), 110 °C and 20 h.

To assess practical applicability, we performed a gram-scale synthesis (5 mmol) under 5 bar CO atmosphere . The reaction proceeded smoothly, delivering the desired product in 72% isolated yield (Scheme 1, **I**), demonstrating the scalability of this method. Mechanistic investigations through control experiments provided further insight. Omitting pentafluoropyridine under standard conditions completely inhibited product formation, indicating its essential role in C-O bond activation. This finding was corroborated by the isolation and structural confirmation of intermediate **1a'**, when subjected to the standard reaction conditions, the purified intermediate cleanly furnished the target product in 95% yield (Scheme 1, **II, III**). Significantly, alternative C-O activation strategies failed to generate **3a** (Scheme 1, **IV, V**), highlighting the distinctive efficiency of the pentafluoropyridine-enabled pathway. Together, these results provide important insights into the proposed catalytic cycle.

Table 3. Scope of alkylamines [a]

[a] Reaction conditions: Step 1: **1a** (0.2 mmol, 1.0 equivalent, 21 μL), pentafluoropyridine (0.3mmol, 1.5 equivalent, 33 μL), KOH (0.5 mmol, 2.5 equivalent, 28 mg), CH$_3$CN (2 mL), N$_2$, r.t. and 5h, Step 2: **2a-2q** (0.5 mmol, 2.5 equivalent), PdCl$_2$ (5 mol%, 1.8 mg), DCyPFc (10 mol%, 11.6 mg), CO (5 bar), 110 °C and 20 h.

### Scheme 1. Large-scale synthesis and control experiments.
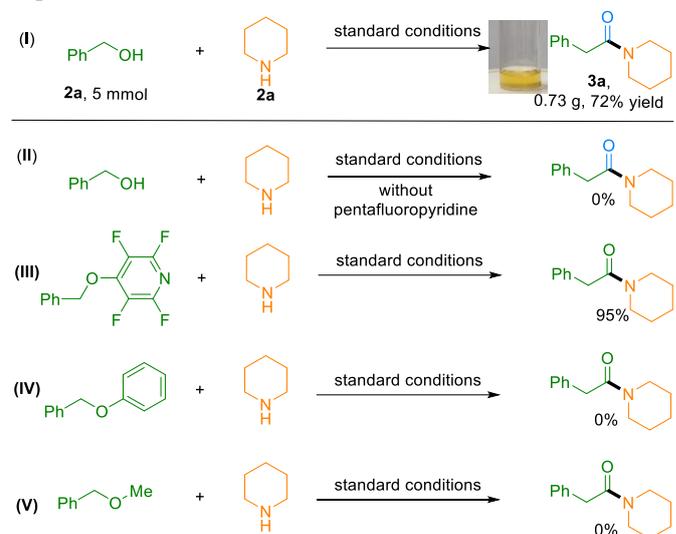
Based on experimental evidence and literature precedents,[6, 10] we propose a plausible catalytic cycle for this direct C-O activation carbonylation (Figure 1). The reaction commences with the in situ formation of activated intermediate **1a'** from benzyl alcohol and pentafluoropyridine, which undergoes oxidative addition with Pd(0) to afford the benzylpalladium(II) species **A**. Following CO coordination and migratory insertion, the acylpalladium complex **C** is generated. Subsequent ligand exchange with piperidine furnishes intermediate **D**, from which reductive elimination delivers the amide product and regenerates the active Pd(0) catalyst for the next catalytic

cycle. This mechanistic proposal aligns with all experimental observations and offers a coherent rationale for this unprecedented deoxygenative carbonylation reactions.

Figure 1 Proposed mechanism.
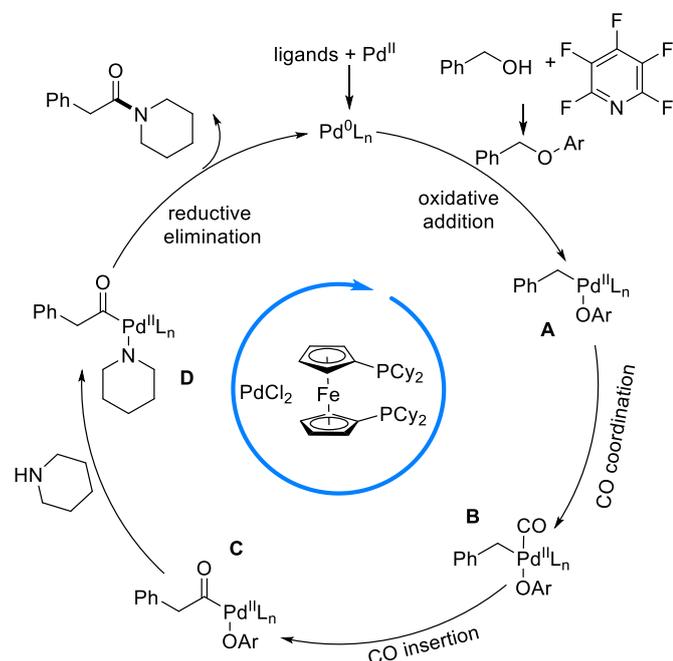
In summary, we have developed a palladium-catalyzed system that achieves the direct carbonylative amidation of benzylic alcohols with alkylamines. This transformation effectively reverses the inherent polarity of the alcohol, activating it as an electrophilic coupling partner, while avoiding the need for prefunctionalized substrates or stoichiometric activators. The protocol is operationally simple, scalable, and compatible with a wide range of functional groups, including those present in pharmaceutically relevant motifs. By providing a concise and sustainable route to benzylic amides from abundant feedstocks, this work not only advances the field of catalytic C–O bond functionalization but also establishes a practical platform for the modular synthesis of amide-containing compounds in organic and medicinal chemistry.

Conflicts of interest

There are no conflicts to declare.

Data availability

The data supporting this article have been included as part of the Supplementary Information.

Notes and references

(1) (a) V. R. Pattabiraman, J. W. Bode, *Nature* 2011, **480**, 471-479. (b) S. Kumari, A. V. Carmona, A. K. Tiwari, P. C. Trippier, *J. Med. Chem.* 2020, **63**, 12290-12358. (c) B. T. Matsuo, P. H. R. Oliveira, E. F. Pissinati, K. B. Vega, I. S. de Jesus, J. T. M. Correia, M. Paixao, *Chem. Commun.* 2022, **58**, 8322-8339.

(2) (a) S. Z. Langer, S. Arbilla, J. Benavides, B. Scatton, *Adv Biochem Psychopharmacol* 1990, **46**, 61-72. (b) D. Aakash, B. R. Kaur, K. Ramanjot, K. Sanjiv, J. U. Kumar, S. Harinder, B. Sandeep, K. Dinesh, D. P. Kishore, *Curr. Top. Med. Chem.* 2017, **17**, 238-250. (c) N. N. Zhang, Y. L. Zhao, D. M. Fan, J. B. Xiao, K. W. Cheng, M. F. Wang, *Food Hydrocoll.* 2020, **108**, 106073.

(3) (a) H. Konishi, K. Manabe, *Tetrahedron Lett.* 2019, **60**, 151147. (b) A. Schoenberg, R. F. Heck, *J. Org. Chem.* 1974, **39**, 3327-3331. (c) A. Schoenberg, I. Bartoletti, R. F. Heck, *J. Org. Chem.* 1974, **39**, 3318-3326.

(4) (a) J. Cao, Z.-J. Zheng, Z. Xu, L.-W. Xu, *Coord. Chem. Rev.* 2017, **336**, 43-53. (b) B. Gabriele, R. Mancuso, G. Salerno, *Eur. J. Org. Chem.* 2012, 6825-6839. (c) Z.-P. Bao, L.-C. Wang, X.-F. Wu, *ACS Catal.* 2025, **15**, 19580-19606.

(5) (a) P. Villo, A. Shatskiy, M. D. Kärkäs, H. Lundberg, *Angew. Chem. Int. Ed.* 2023, **62**, e202211952. (b) E. Bisz, M. Szostak, *ChemSusChem* 2017, **10**, 3964-3981. (c) J. Cornella, C. Zarate, R. Martin, *Chem. Soc. Rev.* 2014, **43**, 8081-8097. (d) T. B. Boit, A. S. Bulger, J. E. Dander, N. K. Garg, *ACS Catal.* 2020, **10**, 12109-12126. (e) S.-Q. Zhang, X. Hong, *Acc. Chem. Res.* 2021, **54**, 2158-2171.

(6) X. W. Qin, H. Li, S. H. Sun, S. Y. Shen, T. F. Xu, X. F. Wu, Z. P. Yin, *Org. Lett.* 2025, **27**, 12524-12529.

(7) (a) J. Duan, J. Jiang, J. Gong, Q. Fan, D. Jiang, *J. Mol. Catal. A: Chem.* 2000, **159**, 89-96. (b) S. Jayasree, A. Seayad, R. V. Chaudhari, *Org. Lett.* 2000, **2**, 203-206. (c) S. Sabater, M. Menche, T. Ghosh, S. Krieg, K. S. L. Rück, R. Paciello, A. Schäfer, P. Comba, A. S. K. Hashmi, T. Schaub, *Organometallics* 2020, **39**, 870-880. (d) Y. Zhu, D. Chen, M. Luo, H. Xia, *Chin. J. Chem.* 2024, **42**, 2765-2772. (e) Y. Zhang, Q. Cao, Y. Xi, X. Wu, J. Qu, Y. Chen, *J. Am. Chem. Soc.* 2024, **146**, 7971-7978.

(8) Y. Li, Z. Wang, X.-F. Wu, *Green Chem.* 2018, **20**, 969-972.

(9) (a) Z. Yin, J. Xu, X.-F. Wu, *ACS Catal.* 2020, **10**, 6510-6531. (b) Z. Yin, W. Shi, X.-F. Wu, *J. Org. Chem.* 2023, **8**, 4975-4994. (c) I. A. Khan, J. Luo, H. B. Shi, Y. Zou, A. Khan, Z. S. Zhu, W. M. Xu, D. Y. Wang, M. Huang, *Food Chem.* 2022, **368**, 130845. (d) N. N. Liang, B. Q. Shi, X. T. Hu, W. T. Li, X. W. Huang, Z. H. Li, X. N. Zhang, X. B. Zou, J. Y. Shi, *Food Chem.* 2024, **460**, 140570. (e) T. An, C. Liu, W. Yuan, X. Qin, Z. Yin, *Chem. Commun.* 2024, **60**, 3389-3392. (f) C. Liu, T. An, W. Yuan, H. Dai, X. Liang, Z. Yin, *Chem. Commun.* 2023, **59**, 12891-12894.

(10) Y. Jiang, Y. Wei, Q.-Y. Zhou, G.-Q. Sun, X.-P. Fu, N. Levin, Y. Zhang, W.-Q. Liu, N. Song, S. Mohammed, B. G. Davis, M. J. Koh, *Nature* 2024, **631**, 319-327.

The data supporting this article have been included as part of the Supplementary Information.