Copper-catalyzed tandem phosphination–decarboxylation–oxidation of alkynyl acids with H-phosphine oxides: a facile synthesis of β-ketophosphine oxides†

Pengbo Zhang, Liangliang Zhang, Yuzhen Gao, Jian Xu, Hua Fang, Guo Tang and Yufen Zhao

The general method for the tandem phosphination–decarboxylation–oxidation of alkynyl acids under aerobic conditions has been developed. In the presence of CuSO₄·5H₂O and TBHP, the reactions provide a novel access to β-ketophosphine oxides in good to excellent yields. This transformation allows the direct formation of a P–C bond and the construction of a keto group in one reaction.

Organophosphorus compounds have broad applications in the fields of organic synthesis, materials science, medicinal chemistry, and ligand chemistry. Thus, development of a new efficient method for C–P bond construction has attracted increasing attention. β-Ketophosphine oxides can facilitate carbon–carbon bond formation and then the diphenylphosphinyl group can be easily removed to give olefins, cyclopropanes, and branched ketones. β-Ketophosphine oxides can also be used for liquid–liquid extraction of metal ions because of their prominent metal-complexing abilities. The traditional methods to prepare β-ketophosphine oxides are based on the acylation of allyl phosphine oxides with carboxylic acid derivatives which employ stoichiometric amounts of the hazardous organometallic reagents. In recent years, our group and other researchers reported many practical approaches to β-ketophosphonates, but these methods are not ideal choices for the synthesis of β-ketophosphine oxides.

In 1966, Nilsson reported the pioneering work of decarboxylative coupling. Since 2002, a series of transition-metal-catalyzed decarboxylative C–C and C–heteroatom bond formation reactions have been extensively developed. Compared with the traditional cross-coupling reactions and C–H activation, decarboxylative coupling reactions using carboxylic acid derivatives have several advantages. Instead of metal waste from organometallic coupling reagents, less toxic carbon dioxide is released as a byproduct after the complete conversion, which reduces the cost of the process for the treatment of waste. It is noteworthy that as a practical alternative, the use of arylpropiolic acids as terminal arylacetylene surrogates is safer and more attractive because arylpropiolic acids are usually solids without an unpleasant smell and are convenient to synthesize, store, and transport. On the basis of this viewpoint, Wu’s group fulfilled the decarboxylative coupling of arylpropiolic acids with P(O)H to construct a Csp–P bond with the assistance of a copper catalyst system. Recently, our group developed an efficient synthesis of E-alkenylphosphine oxides via copper-catalyzed decarboxylative cross-coupling of alkynyl acids with H-phosphine oxides. To the best of our knowledge, the example of β-ketophosphine oxide formation via decarboxylative coupling of alkynyl acids is yet to be reported (Scheme 1).

Reactions involving organophosphorus radicals have a long history, and are useful reactive species in organic synthetic chemistry. Owing to our continuous interest in the P–C bond formation and the reaction of organophosphorus radicals, we present herein our recent progress in constructing valuable bonds involving organophosphorus radicals.

**Scheme 1** C–P bond formation via decarboxylative coupling.
At the outset of our investigation, phenylpropionic acid (1a) and HOPfPh2 (2a) were chosen as the model substrates to survey the reaction conditions. Gratifyingly, when a mixture of 1a (0.2 mmol), 2a (0.8 mmol), CuSO4·5H2O (0.02 mmol) and NH3·H2O (25%, 0.25 mL) in MeCN was heated to 60°C for 2 h, the desired product 3a was obtained in 43% yield (Table 1, entry 1). Subsequently, various Cu(i) and Cu(u) salts were further checked and the results showed that Cu(u) salts were more effective to give the desired product (entries 1–9). A brief survey of bases such as Cs2CO3, K2CO3, NaOAc, (iPr)2NEt, NEt3, pyridine, and NH3·H2O (25%) led to the observation that NH3·H2O (25%) gave 3a in the highest yield (entries 9–16). In our previous synthesis of α-hydroxy phosphonates from H-phosphonates and alcohols, we found that the combined use of Cu(u) and TBHP (tert-butylhydroperoxide) could promote the reaction efficiently. Gratifyingly, the yield increased tremendously when TBHP was employed as an oxidant. However, the other oxidants like K2S2O8, BQ (p-benzoquinone), DTBP (di-tert-butyl peroxide), and H2O2 did not improve the yield (entries 17–22). The solvent systems employed also notably affected the related reaction efficiencies. Conducting the reaction in EtOH, DMF, DMSO and 1,4-dioxane gave the product 3a in very low yield (entries 23–26), while the reaction conducted in MeCN gave a high yield (entry 18). Moreover, the yield was reduced to 30% when using O2 instead of TBHP (entry 22). No desired product was afforded without copper salts (entry 27). The yield of product 3a decreased when

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a Reaction conditions: 1a (0.2 mmol), 2a (0.8 mmol), catalyst (10 mol%), base, oxidant, solvent (1.5 mL) in an open flask at 60°C for 2 h. b Unless otherwise specified, NH3·H2O (25%) was 0.25 mL, other bases were 0.4 mmol. c Yields were determined by 1H NMR. d At 70°C. e At room temperature. f 2a (0.4 mmol).
the temperature was raised to 70 °C or decreased to room temperature (entry 28). However, the attempt to decrease the amount of 2a failed (entry 29). After optimization of the reaction conditions, we established a highly efficient route to the tandem decarboxylation–phosphinination–oxidation of alkynyl acids (entry 18).

With this preliminary result in hand, the generality of the method was explored under the optimized conditions [alkynyl acid (0.2 mmol), P(0-H) (0.8 mmol), CuSO4·5H2O (10 mmol%), TBHP (2 equiv.), NH3·H2O (25%, 0.25 mL) in MeCN at 60 °C in air for 2 h], and the results are summarized in Table 2. In general, a variety of functional groups on the phenyl ring of arylpropionic acids were compatible using this procedure, affording the desired products in good to excellent yields. The methyl substituted arylpropionic acids, such as meta-methyl, para-methyl, and 3,5-dimethyl groups on the aryl ring, reacted with 2a efficiently and gave the desired products 3e–3e in high yields. The ortho-substituted arylpropionic acids exhibited a particularly distinct steric hindrance effect (3b–3e), and the corresponding β-ketophosphine oxide 3b was obtained in a slightly lower yield (75%). Halogen atoms such as fluoro, chloro, bromo, and iodo on the aromatic ring were unaffected under the present reaction conditions to afford the corresponding products 3f–3j in good yields, which could allow for further synthetic transformations. Arylpropionic acids bearing electron-drawing CF3, COOEt, CN groups reacted smoothly to give the corresponding products in good yields (3k–3n). Treatment of diphenyl oxide with methoxy-substituted arylpropionic acid led to the formation of product 3o in 50% yield. Replacing the methoxy group with the trifluoromethoxy group resulted in a higher yield (3p, 87%). More bulky substrates such as 3-(naphthalen-1-yl)propionic acid also smoothly reacted with diphenyl oxide and gave product 3q in 56% yield. In addition, 3-(thiophen-2-yl)propionic acid could also provide the expected product 3s in 46% yield.

The substrate scope was further investigated by reacting phenylpropionic acid (1a) with different organophosphorous reagents. Apart from 2a, di-p-tolyphosphonoxide (2b) and bis(4-chlorophenyl)phosphin oxide (2c) were all suitable substrates, generating the corresponding products 3t and 3u in 86% and 65% yields, respectively. However, diarylphosphon oxide involving a para-bromo substituent 2d produced the desired product 3v in only 43% yield. The butyl(p-toly)(phenyl)phosphone oxide (2e) also efficiently reacted with 1a and led to the corresponding product 3w in 89% yield. Treatment of ethyl phenylphosphate (2f) with 1a afforded the desired product 3x in a lower yield of 42%. When diethyl phosphonate (2g) was used, β-ketophosphonate 3y was obtained in only 22% yield. Alkynyl acid was also examined. Unfortunately, only a trace amount of the desired product was detected by 31P NMR analysis.

With the synthetic β-ketophosphine oxides in hand, we next prepared α-benzyl β-ketodiphenylphosphine oxide 4 from benzy1 bromide and 3a in good yield, which was converted into 1,3-diphenylpropan-1-one 5 in 98% yield via a dephosphinoylation process (Scheme 2).

In an effort to improve our understanding of the reaction profile, a series of isotope labeling studies were conducted. When H2 18O or D2O was added to the reaction mixture under the optimal conditions, no isotope-labeled product was detected in 1H NMR and ESI-MS spectra (Scheme 3a and b). In the absence of air, the transformation could still proceed smoothly to provide 3a in a good yield of 78% (Scheme 3c). It was suggested that the oxygen of the newly formed carbonyl group of 3a mainly originated from TBHP. When 2a was treated with 1.2 equiv. of (phenylethynyl)copper under the optimized reaction conditions, only a trace amount of the desired product was observed, illustrating that (phenylethynyl)copper might not be an intermediate in this process (d). A radical scavenger such as TEMPO could completely restrain the reaction, thus suggesting that the radical processes might be involved. Based on these experimental results and previous mechanistic studies, a plausible mechanism is proposed as shown in Scheme 4.

First, TBHP generates the tert-butoxy radical and the hydroxyl radical in the presence of Cu(II). Then, the tert-butoxy radical triggers the formation of phosphorus radical A from H(O)PPh2. Reaction of the phenylpropionic acid with Cu(II) generates a salt of cupric carboxylate B. Subsequently, addition of phosphorus radical A at the α-position of the triple bond of B gives intermediate C, synthesizing the desired product.
which is ultimately trapped by the hydroxyl radical to form intermediate D. Then D releases one molecular \( \text{CO}_2 \) to produce alkenyl copper intermediate E. Finally, the protonolysis of the intermediate leads to the formation of F, which isomerizes and affords the desired product.

In conclusion, we have successfully developed the first facile method for the preparation of \( \beta \)-ketophosphine oxides via decarboxylation-phosphinolation-oxidation of various alkenyl acids with H-phosphine oxides. Importantly, this transformation method would provide a new pathway for formation of \( \text{Csp}^3-\text{P} \) and \( \text{C} \equiv \text{O} \) bonds in one step. This method is highly efficient and provides a rapid access to a broad spectrum of \( \beta \)-ketophosphine oxides in good to excellent yields. Moreover, the use of an inexpensive \( \text{CuSO}_4 \cdot 5\text{H}_2\text{O} \) catalyst, using readily available alkenyl acids only producing \( \text{CO}_2 \), means that this facile protocol will be attractive for academia and industry.

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Notes and references


