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Recent trends in the direct oxyphosphorylation of C–C multiple bonds

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Due to the wide importance of β -phosphorylated ketones as key building-blocks in the fabrication of various pharmaceutically active organophosphorus compounds, finding new and truly efficient methods for their preparation from simple, low-cost and ubiquitous feedstock materials within a single click is an interesting subject in organic synthesis. Recently, oxyfunctionalization of carbon–carbon multiple bonds has arisen as a straightforward and versatile tool for the synthesis of complex organic molecules from the simple and easily accessible alkenes/alkynes *via* a single operation. In this context, oxyphosphorylation of alkenes/alkynes with P(O)–H compounds has attracted considerable attention as a unique procedure for the construction of β -phosphorylated ketones. In this review, we outline the recent advances and developments in this fast-growing research field with particular emphasis on the mechanistic aspects of reaction.

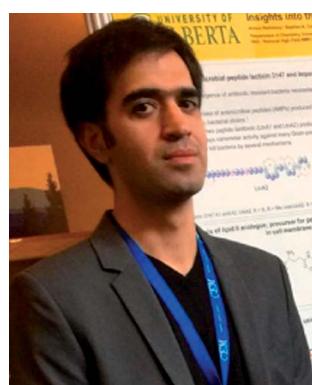
1. Introduction

Organophosphorus compounds play a tremendously important role in the fields of organic chemistry,¹ medicinal chemistry,² and agricultural chemistry.³ Among various organophosphorus compounds, β -phosphorylated ketones (*e.g.*, β -ketophosphonates, β -ketophosphine oxides, β -ketophosphinates) are highly

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valuable building blocks for the synthesis of various value-added chemicals and drug-like molecules.⁴ Moreover, some β -phosphorylated ketones themselves are endowed with outstanding biological properties.⁵ Despite wide importance of these motifs in many fields of chemistry, the methods for their preparation are relatively scarce.⁶ The general protocols for the synthesis of these compounds include the Michaelis–Arbuzov reaction (Scheme 1, route a),⁷ hydration of alkynyl phosphorus compounds (Scheme 1, route b),⁸ acylation of alkyl phosphorus compounds (Scheme 1, route c),⁹ and decarboxylative coupling reaction of alkenyl/alkynyl carboxylic acids with P(O)H compounds (Scheme 1, route d).¹⁰ However, most of these methodologies, if not all, suffer from certain disadvantages, such as requiring harsh reaction conditions, requiring pre-functionalized starting materials, and/or expensive catalysts. Therefore, the development of efficient, practical, and



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environmentally benign methodologies that benefit from simple, inexpensive, and readily available substrates for the construction of this specific class of phosphorus and carbonyl-contain compounds is still highly desirable in synthetic chemistry.

Recently, the double functionalization of carbon–carbon multiple bonds in one-pot processes has emerged as a fruitful tool for the rapid synthesis of complex molecular architectures from safe, low-cost, and prevalent alkenes and alkynes.^{11,12} In this family of reactions, oxyfunctionalization of alkenes/alkynes, which installs a unique functional group and an oxo group in adjacent carbons simultaneously, offers a powerful strategy for the selective synthesis of β -substituted ketones.¹³ In this context, direct oxyphosphorylation of C–C multiple bonds with P(O)–H compounds has gained a considerable amount of attention in the last couple of years in the preparation of numerous synthetically and biologically important β -phosphorylated ketones (Fig. 1). This protocol features readily accessible starting materials, high atom-, step-, and pot-economy, and high bond-forming efficiency, thus providing a straightforward and highly efficient access to the title compounds. Despite significant advances and developments in this field over the past decade, no comprehensive review can be found in the literature covering this novel and rapidly growing research field. In continuation of our recent published reviews on organophosphorus chemistry^{10,14} and modern organic synthesis,¹⁵ herein, we will highlight the most important developments on the synthesis of β -substituted ketones through the direct oxyphosphorylation of alkenes/alkynes with P(O)–H compounds from 2011 to August 2020. It is worthy of note that a particular emphasis is placed on mechanistic aspects of reactions which may allow possible new insights toward designing more efficient and improved catalytic systems.



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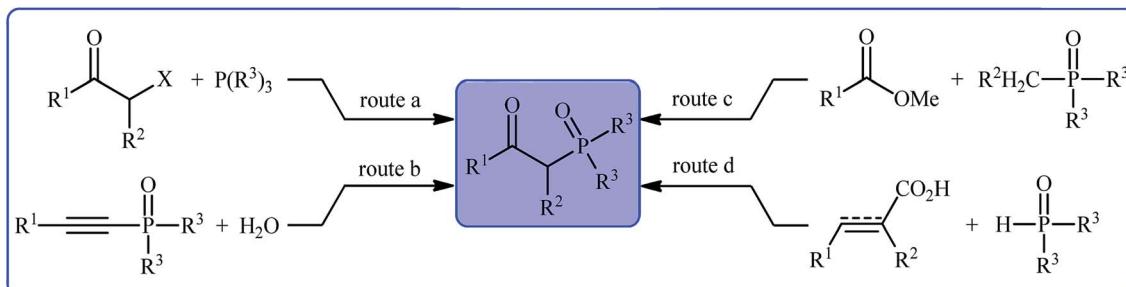
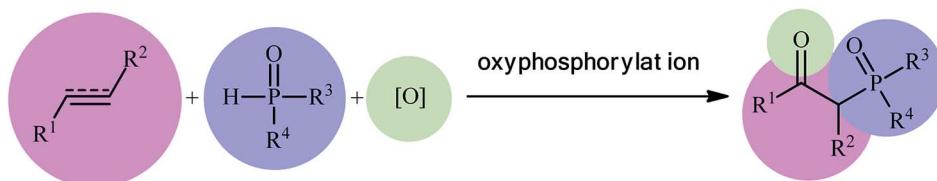
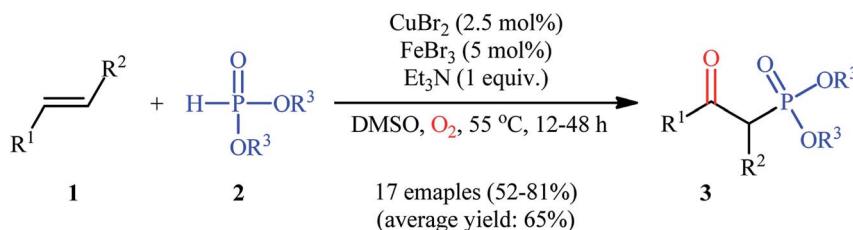
Scheme 1 Main synthetic routes to β -phosphorylated ketones.

Fig. 1 Direct oxyphosphorylation of C–C multiple bonds with H–P(O) compounds.

2. Oxyphosphorylation of alkenes

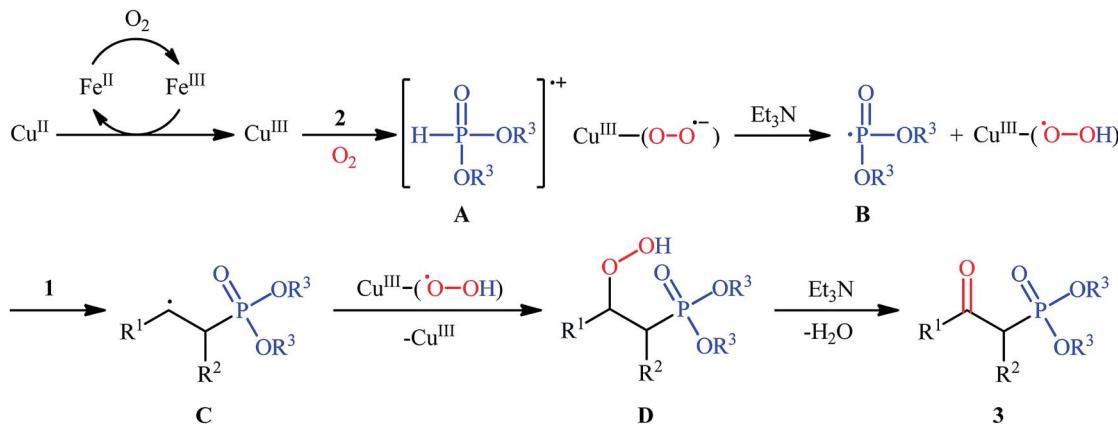
The first mention of the synthesis of β -ketophosphonates through the direct oxyphosphorylation of alkenes with H-phosphonates and dioxygen can be found in a 2011 paper by Ji and Wei.¹⁶ To evaluate the catalytic activity of different transition metal complexes, styrene and diisopropyl phosphonate were chosen as the model substrates. They showed that in the presence of a combination of inexpensive and easily accessible CuBr_2 and FeBr_3 as a catalytic system in DMSO, a variety of electron-rich and electron-deficient styrenes **1** and various H-phosphonates **2** reacted efficiently to give the corresponding β -ketophosphonates **3** in moderate to high yields (Scheme 2). Interestingly, an enamine such as 9-vinylcarbazole was also compatible with this protocol, thus affording the corresponding product in moderate yield. However, 1-octene did not work well in the reaction and therefore no other aliphatic alkenes were examined in this procedure. Of note, the authors demonstrated

the scalability of the reaction since diethyl (2-oxo-2-phenylethyl) phosphonate, a key intermediate for the synthesis of various natural products such as (–)-diospongin B, could be obtained in 26 g scale in good yield of 68%. To make the mechanistic course of this conversion clearer, the authors performed ^{18}O isotope labeling and radical trapping experiments. From these results, they speculated that the reaction starts with the formation of dialkyl phosphonate cation radical **A** and $\text{Cu}^{\text{III}}-(\text{OO}^{\cdot-})$ (superoxide) species through the single electron transfer (SET) from H-phosphonate **2** to *in situ* generated copper(III) species under O_2 . Next, the abstraction of the H^+ by Et_3N from the cation radical **A** leads to phosphonyl radical **B** and $\text{Cu}^{\text{III}}-(\text{OOH})$ (hydroperoxide) species. Subsequently, the reaction of phosphorus-centered radical **B** with alkene **1** gives the alkyl radical **C**, which after interaction with $\text{Cu}^{\text{III}}-(\text{OOH})$ converts to the hydroperoxide intermediate **D** and regenerates the starting Cu^{III} catalyst. Finally, dehydration this intermediate **D** with the aim of Et_3N affords the desired product **3** (Scheme 3).



$\text{R}^1 = \text{Ph, 4-Me-C}_6\text{H}_4, 4\text{-OMe-C}_6\text{H}_4, 4\text{-F-C}_6\text{H}_4, 4\text{-Cl-C}_6\text{H}_4, 4\text{-Br-C}_6\text{H}_4, 4\text{-OCOMe-C}_6\text{H}_4,$
 $4\text{-CN-C}_6\text{H}_4, 3\text{-Me-C}_6\text{H}_4, 2\text{-Me-C}_6\text{H}_4, 2\text{-naphthyl, 2-pyridyl}$
 $\text{R}^2 = \text{H, Me}$
 $\text{R}^1 + \text{R}^2 =$
 $\text{R}^3 = \text{Et, }^i\text{Pr, }^n\text{Bu}$

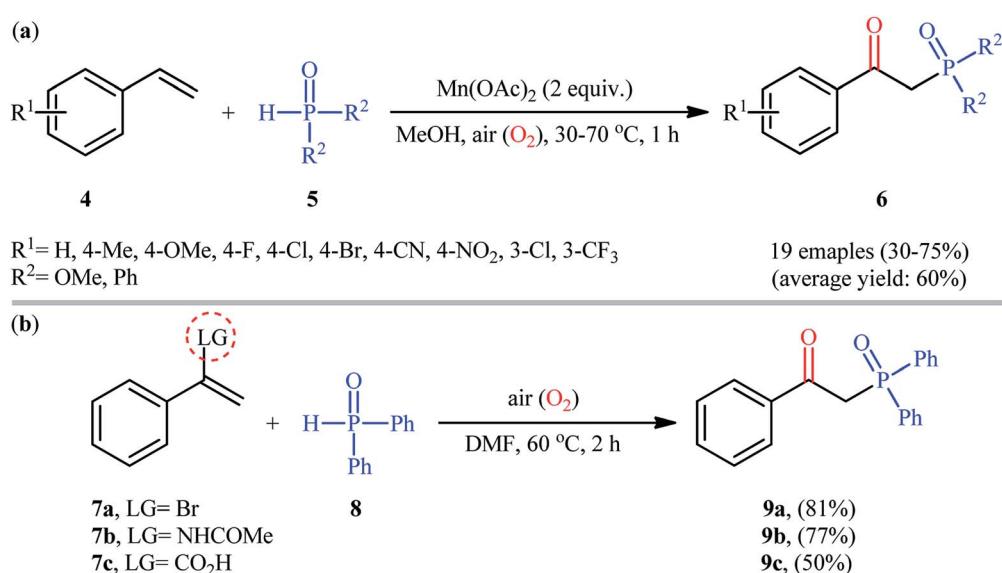
Scheme 2 Ji's synthesis of β -ketophosphonates **3**.

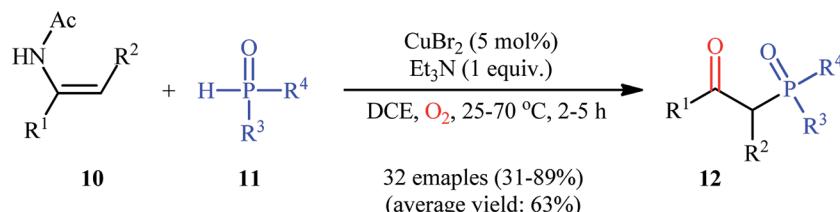
Scheme 3 Possible mechanism of Cu-catalyzed oxyphosphorylation of alkenes 1 with H-phosphonates 2 and O₂.

Afterwards, Zeng and Zou along with their co-workers developed a Mn(OAc)₂-mediated protocol for the oxyphosphorylation of styrene derivatives **4** with P(O)-H compounds **5** (H-phosphonates and secondary phosphine oxides) under aerobic conditions without the use of any base and additive (Scheme 4a).¹⁷ This chemistry showed good functional group tolerance and afforded the target β -ketophosphoryl products **6** in moderate to good yields within 1 h. In this investigation, the authors found some limitation in their methodology, when they attempted to react β -substituted styrenes and non-conjugated terminal alkenes. Unfortunately, in these cases only the hydrophosphorylation products were observed with no formation of the expected β -phosphorylated ketone products. Noteworthy, the reaction of α -substituted styrenes with P(O)-H compounds under the standard conditions also provided the hydroxyphosphorylation products due to the presence of α -substituents. Concurrently, the same

strategy was applied by Zhang's group to the synthesis of a diverse range of β -hydroxyphosphine oxides through the reaction of corresponding 1,1-disubstituted alkenes with H-phosphine oxides and O₂ under catalyst- and additive-free conditions.¹⁸ Following these works, the group of Lei disclosed that if easily removable leaving groups were introduced to α -position of styrenes, β -ketophosphoryl products could be obtained exclusively.¹⁹ Thus, a small series of β -ketophosphonates **9** were synthesized in good yields by simple heating of α -substituted styrenes **7** with diphenylphosphine oxide **8** in DMF under air atmosphere and catalyst-free conditions (Scheme 4b).

Drawing inspiration from these works, Du-Wei and colleagues were able to demonstrate that a library of β -ketophosphoryl products **12** could be obtained in fair to good yields from the reaction of various terminal and internal alkenes **10** with P(O)-H compounds **11** under an oxygen atmosphere using

Scheme 4 (a) Mn-mediated oxyphosphorylation of styrenes **4** with P(O)-H compounds **5**; (b) catalyst-free oxyphosphorylation of 1,1-disubstituted alkenes **7** with diphenylphosphine oxide **8**.



$R^1 = \text{Ph, 4-Me-C}_6\text{H}_4, 4\text{-OMe-C}_6\text{H}_4, 4\text{-F-C}_6\text{H}_4, 4\text{-Cl-C}_6\text{H}_4, 4\text{-CF}_3\text{-C}_6\text{H}_4, 4\text{-CO}_2\text{Me-C}_6\text{H}_4,$
 $3\text{-Me-C}_6\text{H}_4, 2\text{-Me-C}_6\text{H}_4, 2\text{-OMe-C}_6\text{H}_4, 2,4\text{-Cl}_2\text{-C}_6\text{H}_3, 2\text{-naphthyl, 3-pyridyl, 2-thiazolyl}$
 $R^2 = \text{H, Me}$
 $R^1 + R^2 = \text{Ar, Ar', Ar''}$
 $R^3 = \text{Ph, 4-Me-C}_6\text{H}_4, 4\text{-OMe-C}_6\text{H}_4, 4\text{-F-C}_6\text{H}_4, \text{OMe, OEt, O}^{\prime\prime}\text{Bu}$
 $R^4 = \text{Ph, 4-Me-C}_6\text{H}_4, 4\text{-OMe-C}_6\text{H}_4, 4\text{-F-C}_6\text{H}_4, \text{OMe, OEt, O}^{\prime\prime}\text{Bu}$

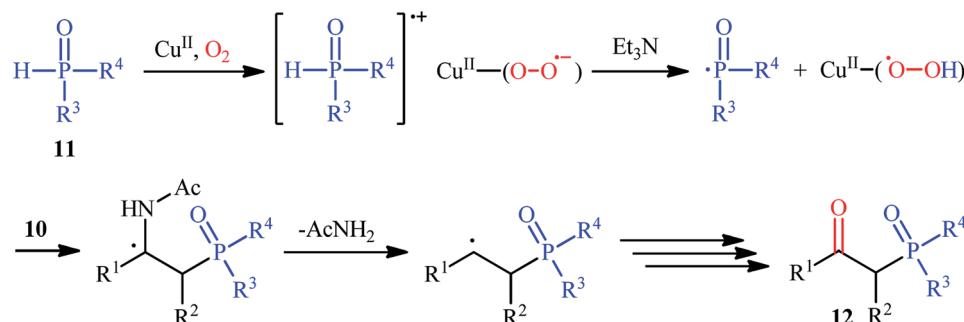
Scheme 5 Cu-catalyzed oxyphosphorylation of enamides 10 with P(O)-H compounds 11 and O₂.

CuBr₂ as a catalyst and Et₃N as a base in DCE (Scheme 5).²⁰ Some of the most important information of this C-P bond forming reaction are listed below: (i) all the three kinds of hydrogen phosphoryl compounds (H-phosphonates, H-phosphine oxides, and H-phosphinates) were compatible in this reaction; (ii) the relative reaction rates of P(O)-H compounds followed the order: H-phosphine oxides > H-phosphonates ≈ H-phosphinates; (iii) terminal alkenes afforded higher yields of the expected products compared to the internal alkenes; (iv) the protocol for oxyphosphorylation of aliphatic alkenes was considerably less efficient; and (v) the electronic nature of the substituents on phenyl ring periphery of aromatic alkenes had a little impact on the rate of reaction. The authors suggested reaction mechanism for this transformation is displayed in Scheme 6 and is closely similar to the one proposed by Ji and Wei.

In 2017, Chen and co-workers achieved CuSO₄·5H₂O-catalyzed aerobic oxyphosphorylation of styrenes 13 with various hydrogen phosphoryl compounds 14 under additive-free conditions.²¹ The reactions were carried out in MeCN at 60 °C, tolerated a wide number of electron-donating and electron-withdrawing functional groups and generally provided the desired β-ketophosphoryl products 15 in modest to excellent yields (Scheme 7a). However, aliphatic alkenes did not work well under standard conditions. It is worthwhile to mention that the optimized reaction condition was also successfully applied in the decarboxylative oxyphosphorylation of alkenyl

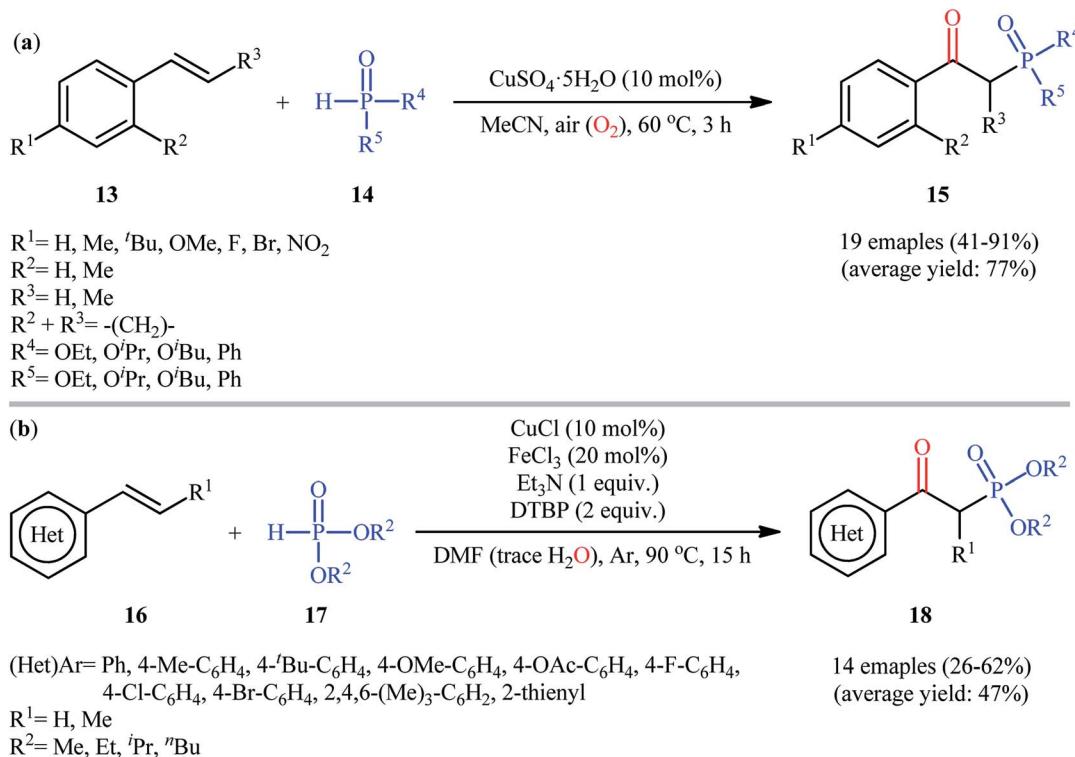
acids with the same set of P(O)-H compounds. To gain insights into the mechanistic course of this reaction, the authors performed the reaction under an inert (nitrogen) atmosphere and no desired product was observed, indicating that the presence of oxygen is indispensable. Moreover, an isotope labeling study in the presence of H₂¹⁸O was conducted, which revealed that the carbonyl oxygen in the products came from oxygen in air. A similar oxyphosphorylation of (hetero)aromatic alkenes with H-phosphine oxides and O₂ was also achieved only using CuCN as the catalyst under additive-free conditions.²² Concurrently, another metal-catalyzed oxyphosphorylation of alkenes was introduced by Gu and Cai.²³ In this methodology, styrene derivatives 16 reacted with H-phosphonates 17 in presence of 10 mol% of CuCl as a catalyst, 20 mol% of FeCl₃ as a co-catalyst, 1 equiv. of Et₃N as a base, and 2 equiv. of DTBP (di-*tert*-butyl peroxide) as the oxidant in DMF under Ar atmosphere at 90 °C to achieve β-ketophosphonates 18 in 26–62% yields (Scheme 7b). In contrast with previous reports which suggested that carbonyl oxygen in the products should originally be from oxygen (air), the authors proposed that carbonyl oxygen in their products originated from water molecules which are present in DMF solvent. Interestingly, when the same reaction was run at 110 °C, vinylphosphonates were obtained as the sole products. They ascribed this observation by the boiling point of water.

Afterwards, Yamamoto–Makino's group extended the scope of alkenes to vinyl ethers.²⁴ Thus, several alkyl/aryl 2-(diethoxyphosphoryl)acetates 21 were synthesized in moderate yields,



Scheme 6 Possible reaction pathway for the formation of β-ketophosphoryl compounds 12.



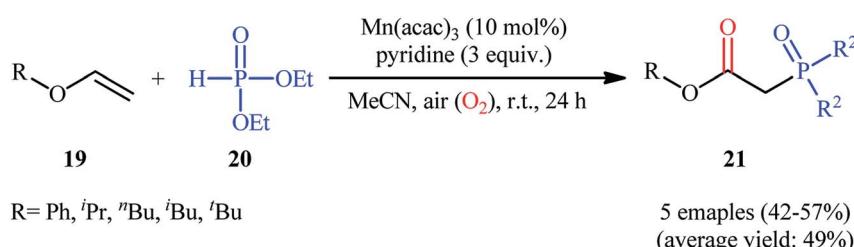


Scheme 7 (a) Synthesis of β -phosphorylated ketones **15** from styrenes **13** and hydrogen phosphoryl compounds **14** under ambient air; (b) Cai's synthesis of β -ketophosphonates **18**.

ranging from 42% to 57%, *via* the $\text{Mn}(\text{acac})_3$ -catalyzed oxyphosphorylation of corresponding vinyl ethers **19** with dimethyl phosphonate **20** and O_2 in MeCN at room temperature (Scheme 8). Noteworthy, $\text{Mn}(\text{acac})_2$ was also found to promote this transformation; albeit, at lower efficiency. However, other manganese salts such as MnBr_2 , $\text{Mn}(\text{OAc})_2$, $\text{Mn}(\text{OAc})_3 \cdot 2\text{H}_2\text{O}$, and $[\text{Mn}^{\text{III}}(\text{Pc})]\text{Cl}$ proved to be completely ineffective. The outcome of this reaction was strongly depended to the base used, and among the various bases (*e.g.*, Et_3N , $^\text{t}\text{Pr}_2\text{Net}$, Cy_2Net , DMAP, DTBMP, TMEDA, DBU, pyridine, 2,2'-bipyridyl, 4-methoxypyridine, 4-(*tert*-butyl)pyridine, 4-(trifluoromethyl)pyridine, 2,6-dimethylpyridine, 2,4,6-trimethylpyridine); pyridine was found to be most effective for this conversion.

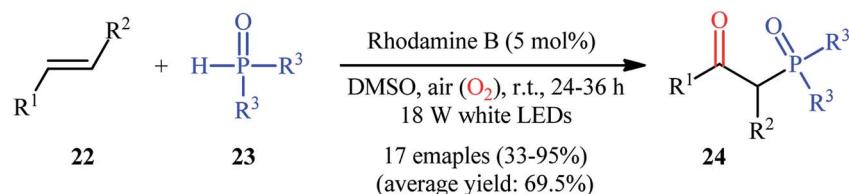
Recently, Zhu *et al.*²⁵ reported a metal-free, visible light photoredox-catalyzed aerobic oxyphosphorylation of alkenes **22** with H-phosphine oxides **23** at room temperature. No additive or base was used and only 5 mol% of inexpensive rhodamine B catalyst was enough for preparation of a variety of β -

ketophosphine oxides **24** in 33–95% yields (Scheme 9). The reaction could also be easily scaled up to the gram-scale as exemplified by the synthesis of 1-(4-chlorophenyl)-2-(diphenylphosphoryl)ethanone on a 1.62 g scale (91%). Remarkably, H-phosphinates were also responded to this reaction. However, H-phosphonates were inert under standard conditions. The radical trapping experiments with 2,2,6,6-tetramethyl-1-piperidinyloxy (TEMPO) pointed toward a radical pathway. On the basis of these results and literature reports, the authors suggested that the reaction proceeds through the following key steps (Scheme 10): (i) photoexcitation of the ground state photocatalyst (PC) by visible light to form the excited state photocatalyst (PC^*); (ii) single-electron transfer from PC^* to phosphinous acid A (the tautomeric form of diarylphosphine oxide **23**) to produce the radical cation intermediate **B** and $\text{PC}^{\cdot-}$ radical anion; (iii) oxidation of $\text{PC}^{\cdot-}$ by O_2 to afford $\text{O}_2^{\cdot-}$ and regenerate the PC; (iv) deprotonation of intermediate **B** to yield the P-centered radical **C**; (v) reaction of



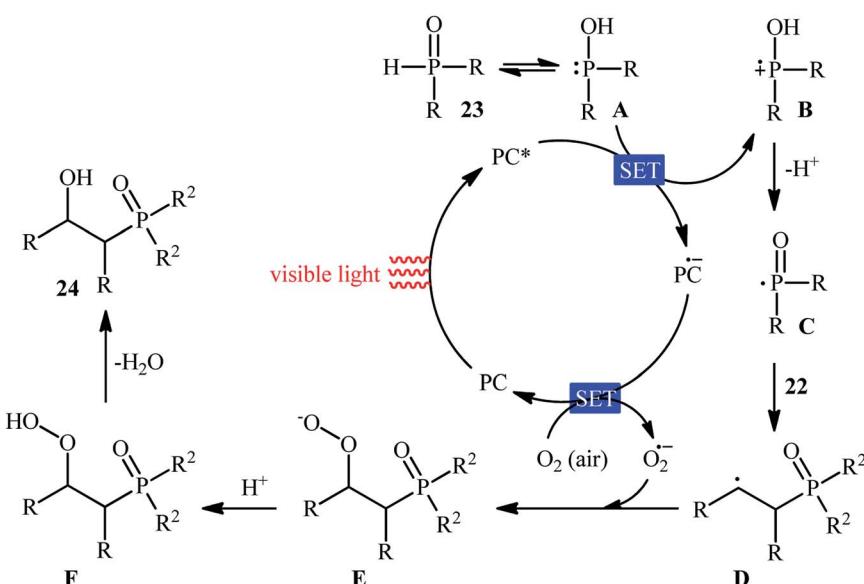
Scheme 8 Mn-catalyzed direct oxyphosphorylation of vinyl ethers **19** with dimethyl phosphonate **20** developed by Yamamoto and Makino.





$R^1 = \text{Ph, 4-Me-C}_6\text{H}_4, 4'\text{-Bu-C}_6\text{H}_4, 4\text{-OMe-C}_6\text{H}_4, 4\text{-Cl-C}_6\text{H}_4, 4\text{-CF}_3\text{-C}_6\text{H}_4, 4\text{-CN-C}_6\text{H}_4,$
 $4\text{-NO}_2\text{-C}_6\text{H}_4, 3\text{-Cl-C}_6\text{H}_4, 2\text{-Cl-C}_6\text{H}_4, 2,3\text{-(OMe)}_2\text{-C}_6\text{H}_3, 2\text{-naphthyl, 2-thienyl}$
 $R^2 = \text{H, Me}$
 $R^1 + R^2 = \text{C}_6\text{H}_4$
 $R^3 = \text{Ph, 4-OMe-C}_6\text{H}_4, 4\text{-F-C}_6\text{H}_4$

Scheme 9 Visible-light-induced direct oxyphosphorylation of alkenes 22 with H-phosphine oxides 23.

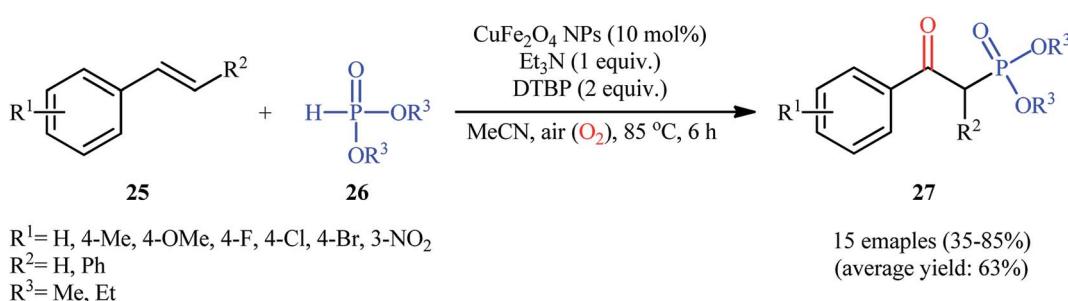


Scheme 10 Proposed mechanism for the reaction in Scheme 9.

intermediate C with alkene 22 to give radical D; (vi) the combination of intermediate D with O_2^- to generate peroxide anion E; (vii) protonation of intermediate E to form intermediate F; and (viii) dehydration of intermediate F to give the desired product 24.

Very recently in 2020, the nanocatalyzed version of this reaction was reported by Moghaddam and colleagues, who

disclosed that the treatment of various styrenes 25 bearing either electron-donating or electron-withdrawing groups at the phenyl ring periphery with H-phosphonates 26 in the presence of 10 mol% of copper ferrite nanoparticles (CuFe_2O_4 NPs), 1 equiv. of Et_3N , and 2 equiv. of DTBP in MeCN under ambient air afforded the corresponding β -ketophosphonates 27 with yield ranging from 35% to 85% (Scheme 11).²⁶ To evaluate the



Scheme 11 Synthesis of β -ketophosphonate derivatives 27 from styrenes 25 and H-phosphonates 26 catalyzed by copper ferrite nanoparticles.



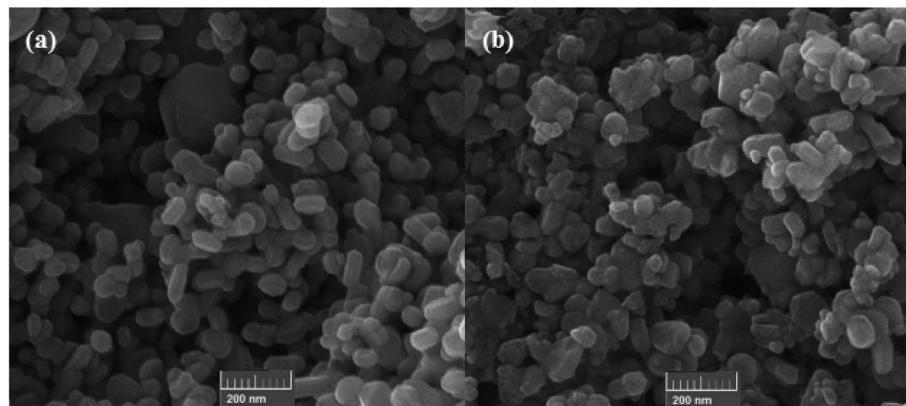


Fig. 2 The FE-SEM images of CuFe_2O_4 NPs (a) before and (b) after six times reuse.

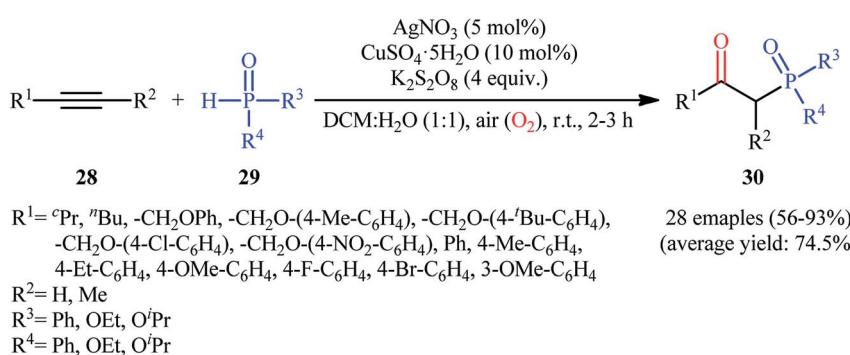
recyclability and reusability of the catalyst, the oxyphosphorylation of styrene with diethyl phosphonate was performed under the optimized reaction conditions. It was found that the catalyst could be easily recovered by means of a magnetic rod and reused six consecutive cycles without significant decrease in its catalytic activity. Noteworthy, the FE-SEM (field emission scanning electron microscopy) images of the catalysts before and after six cycles of the reaction indicated that the morphology, size, and dispersity of the nanoparticles were not approximately transformed during these runs (Fig. 2).

3. Oxyphosphorylation of alkynes

One of the earliest reports on the synthesis of β -phosphorylated ketones *via* the direct oxyphosphorylation of the corresponding alkynes appeared in 2015, when various available aryl/aliphatic alkynes **28** were converted to β -ketophosphoryl compounds **30** through the silver/copper-catalyzed oxyphosphorylation with $\text{P}(\text{O})-\text{H}$ compounds **29** and O_2 in binary solvent $\text{MeCN}/\text{H}_2\text{O}$ with ratio 1 : 1 at room temperature.²⁷ As shown in Scheme 12, this reaction tolerated all the three kinds of hydrogen phosphoryl compounds and both terminal and internal alkynes, and gave the final products in moderate to excellent yields within 2–3 h. The reactivity order for the $\text{P}(\text{O})-\text{H}$ compounds under these reaction conditions was $\text{H-phosphonates} > \text{H-phosphinates} \equiv \text{H-phosphine oxides}$. The results also indicated that terminal

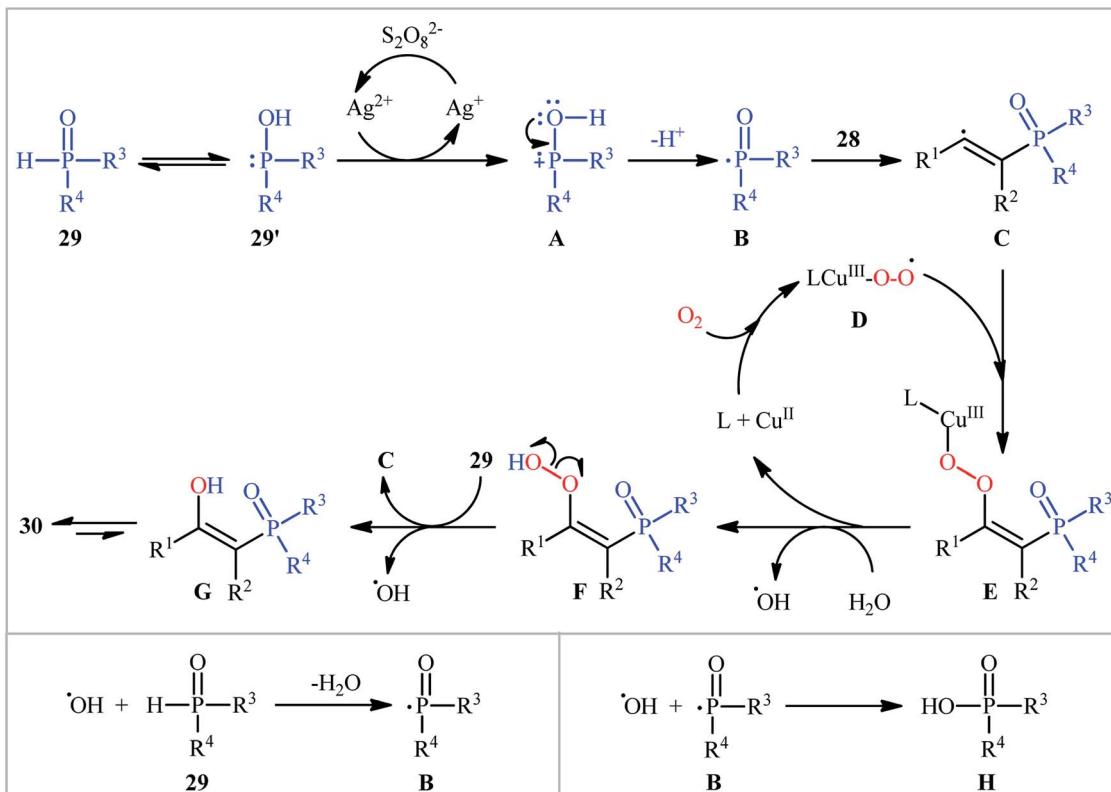
alkynes afforded better yields compared to internal alkynes. Beside good yields, broad substrate scope, mild reaction condition, and low reaction times were the advantages, mentioned for this atom economy synthetic strategy. The plausible mechanism for this appealing reaction is outlined based on a series of control experiments (Scheme 13). At the beginning of the reaction, peroxodisulfate oxidize $\text{Ag}(\text{i})$ cation to $\text{Ag}(\text{ii})$ cation by a single-electron transfer process and the generated $\text{Ag}(\text{ii})$ ion oxidize $\text{P}(\text{O})-\text{H}$ compound **29** to form cation radical **A**. Next, deprotonation of this intermediate gives the phosphoryl radical **B**, which then attacks to the triple bond of alkyne **28** to produce alkenyl radical intermediate **C**. Subsequently, reaction of this radical with *in situ* generated active-oxygen copper complex **D** gives complex **E**, which in the presence of water quickly converts to hydroperoxide species **F**. Then, *via* initial homolysis of the $\text{O}-\text{O}$ bond, the hydroperoxide **F** reacts with another molecule of **2** to form a hydroxyl radical, an enol **G** as well as a phosphoryl radical **B**. Finally, tautomerization of **D** affords the observed products **30**.

In the same year, He's research team presented an efficient copper/iron-catalyzed oxyphosphorylation of terminal alkynes **31** with H-phosphonates **32** and O_2 into β -ketophosphonates **33**.²⁸ The results of this investigation indicated that the optimum condition for this transformation was the combination of $\text{Cu}(\text{acac})_2$ (10 mol%), FeCl_3 (20 mol%), and Et_3N (1

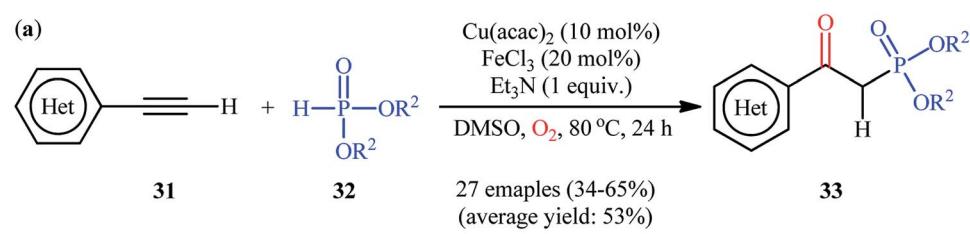


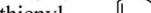
Scheme 12 Ag/Cu-catalyzed direct oxyphosphorylation of alkynes **28** with $\text{P}(\text{O})-\text{H}$ compounds **29** and O_2 .



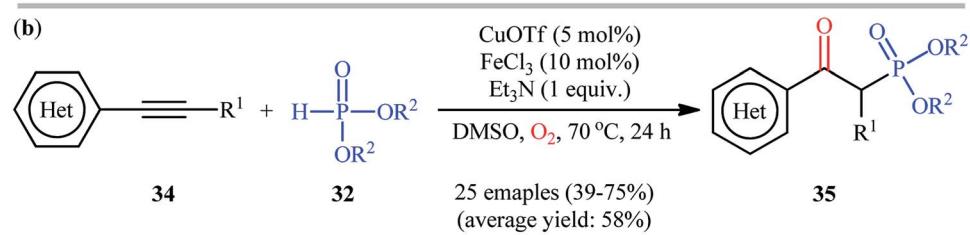


Scheme 13 Mechanistic proposal from the formation of β -ketophosphoryl compounds **30**



(Het)Ar= Ph, 4-Me-C₆H₄, 4-tBu-C₆H₄, 4-Ph-C₆H₄, 4-OMe-C₆H₄, 4-OBn-C₆H₄, 4-F-C₆H₄,
 4-Cl-C₆H₄, 4-Br-C₆H₄, 4-COME-C₆H₄, 4-CO₂Me-C₆H₄, 4-CN-C₆H₄,
 4-OCOCH(NHBoc)Bn-C₆H₄, 3-Me-C₆H₄, 3-Br-C₆H₄, 2-Me-C₆H₄,
 2-Br-C₆H₄, 2-naphthyl, 3-benzo[*b*]thiophenyl, 2-thienyl, 3-thienyl, 

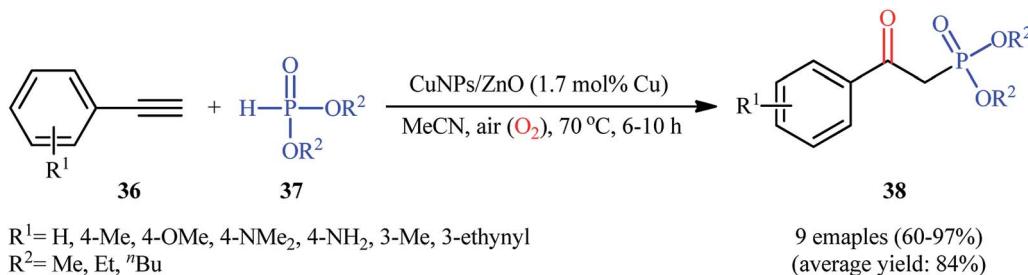
R¹= Me, Et, ⁱPr, ⁿBu, Bn



(Het)Ar= Ph, 4-Me-C₆H₄, 4-Et-C₆H₄, 4-ⁱPr-C₆H₄, 4-^tBu-C₆H₄, 4-Ph-C₆H₄, 4-OMe-C₆H₄, 4-F-C₆H₄, 4-Cl-C₆H₄, 4-Br-C₆H₄, 4-CN-C₆H₄, 4-CF₃-C₆H₄, 4-O^tC₅H₁₁-C₆H₄, 3-Me-C₆H₄, 3-F-C₆H₄, 3-Cl-C₆H₄, 1-naphthyl, 9-phenanthrenyl, 2-thienyl, 2-pyridyl

$\text{R}^1 = \text{H, Me}$
 $\text{R}^2 = \text{Me, Et, } i\text{Pr, } n\text{Bu, Bn}$

Scheme 14 (a) Cu/Fe-catalyzed oxidative coupling of aromatic terminal alkynes **31** and with H-phosphonates **32**; (b) Song's synthesis of β -phosphorylated ketones **35**.

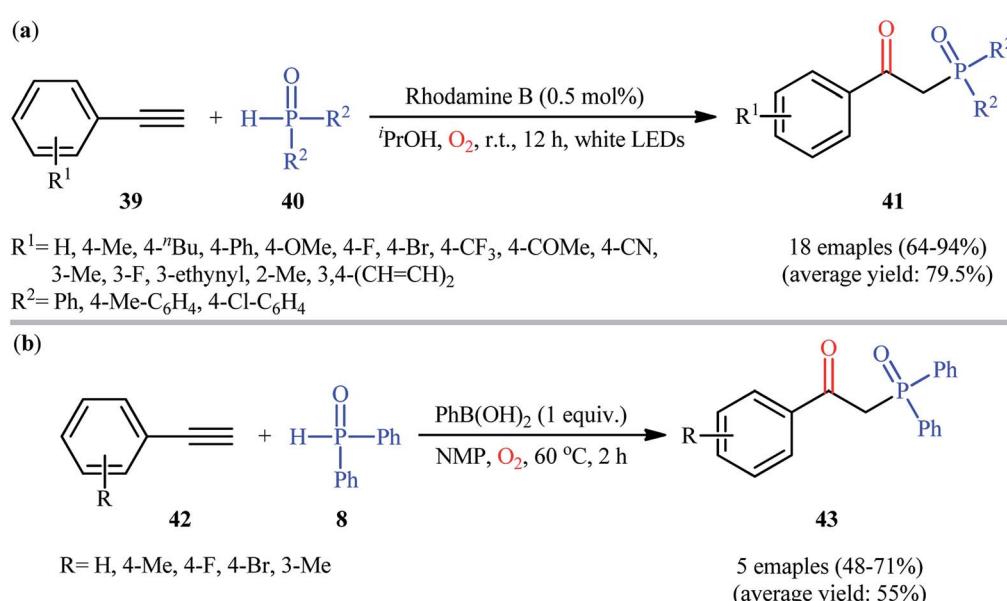
Scheme 15 CuNPs/ZnO-catalyzed oxyphosphorylation of aromatic terminal alkynes **36** with dialkyl H-phosphonates **37**.

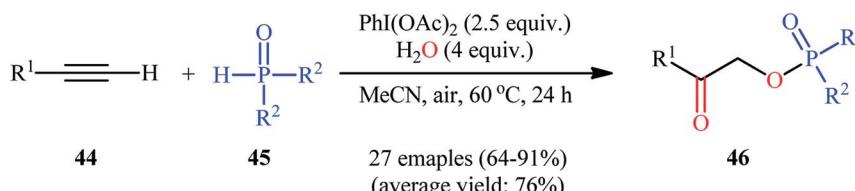
equiv.) as a catalytic system using DMSO as the solvent, at 80 $^\circ\text{C}$. Under optimized conditions, the reaction tolerated a diverse set of aromatic and heteroaromatic terminal alkynes and various symmetrical dialkyl H-phosphonates and gave the desired products in moderate to good yields (Scheme 14a). However, both internal and aliphatic terminal alkynes failed to enter into this reaction. Moreover, strongly electron-poor heteroaromatic alkynes, such as 2- and 3-ethynylpyridines were inert under standard conditions. Shortly thereafter, similar work was reported by Song *et al.*²⁹ Using CuOTf as catalyst and FeCl_3 as cocatalyst, they realized the oxyphosphorylation of various terminal and internal alkynes **34** with the same set of H-phosphonates **32** into the corresponding β -phosphorylated ketones **35** (Scheme 14b). Notably, this method allowed for the oxyphosphorylation of 2-ethynylpyridine, which was inert under the He's reaction conditions. This chemistry was also applicable to the coupling of H-phosphine oxides and H-phosphinates with terminal alkynes. However, the reaction did not work well for aliphatic alkynes.

Later, Moglie and Radivoy along with their co-workers introduced the use of copper nanoparticles (CuNPs)

supported on ZnO as an efficient catalyst for oxyphosphorylation of terminal alkynes with H-phosphonates under aerobic conditions.³⁰ Thus, a number of β -aryl- β -ketophosphonates **38** were successfully synthesized by treatment of the corresponding aromatic terminal alkynes **36** with dialkyl H-phosphonates **37** in the presence of a catalytic amount of CuNPs@ZnO and in the absence of any additive or ligand (Scheme 15). Interestingly, when aliphatic alkynes were subjected to the same reaction conditions, the anti-Markovnikov vinyl phosphonates were formed as the main reaction products, thus suggesting that the presence of a conjugated $\text{C}=\text{C}$ in the starting alkyne is mandatory for the formation of the β -ketophosphonate products. The strategy was also successfully applied to the fabrication of β -ketophosphonates from both aromatic and aliphatic terminal alkenes. It should be mentioned that the catalyst demonstrated not to be compatible with the presence of hydroxyl or carboxylic acid groups in the starting alkyne or alkyne.

In 2016, Bu, Lu, and Cai demonstrated an innovative visible-light-induced organocatalytic oxyphosphorylation of alkynes with H-phosphine oxides under mild conditions.³¹ Thus, in the

Scheme 16 (a) Visible-light-mediated oxidative coupling of terminal alkynes **39** and diaryl phosphine oxides **40**; (b) PhB(OH)_2 -mediated oxyphosphorylation of aromatic terminal alkynes **42** with diphenylphosphine oxide **8**.



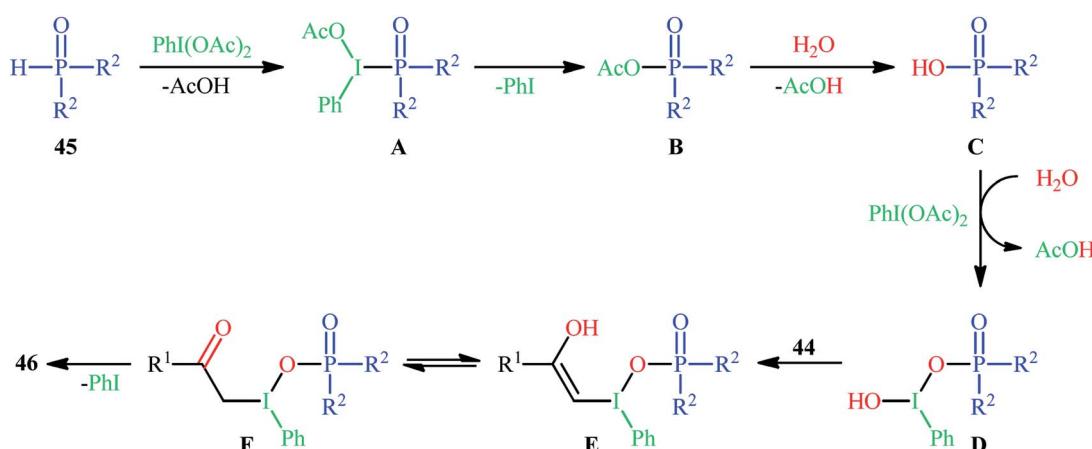
$R^1 = {}^n\text{Bu}, {}^t\text{Bu}, -\text{CH}_2\text{CH}_2\text{CH}_2\text{CN}, -\text{CH}_2\text{-cPent}, -\text{CH}_2\text{Bn}, \text{Ph}, 4\text{-Me-C}_6\text{H}_4, 4\text{-Et-C}_6\text{H}_4, 4\text{-OMe-C}_6\text{H}_4, 4\text{-F-C}_6\text{H}_4, 4\text{-Cl-C}_6\text{H}_4, 4\text{-Br-C}_6\text{H}_4, 4\text{-CF}_3\text{-C}_6\text{H}_4, 4\text{-NO}_2\text{-C}_6\text{H}_4, 4\text{-CO}_2\text{Me-C}_6\text{H}_4, 4\text{-CH}_2\text{CN-C}_6\text{H}_4, 1\text{-naphthyl}, 3\text{-thienyl}, 3\text{-Me-C}_6\text{H}_4, 3\text{-Cl-C}_6\text{H}_4, 2\text{-Me-C}_6\text{H}_4, 2\text{-Cl-C}_6\text{H}_4,$
 $R^2 = \text{Ph}, 4\text{-Me-C}_6\text{H}_4, 4\text{-OMe-C}_6\text{H}_4, 4\text{-F-C}_6\text{H}_4, 4\text{-Cl-C}_6\text{H}_4$

Scheme 17 Hypervalent iodine-mediated direct synthesis of diaryl (2-oxo-2-alkyl/arylethyl)phosphinates **46** from terminal alkynes **44** and diaryl H-phosphine oxides **45**.

presence of only 0.5 mol% of rhodamine B as a photocatalyst in ${}^i\text{PrOH}$ under an oxygen atmosphere and irradiation with a white LED lamp at room temperature, oxidative coupling of aromatic terminal alkynes **39** and diaryl phosphine oxides **40** furnished the corresponding β -ketophosphine oxide derivatives **41** in good to excellent yields (Scheme 16a). However, the system was unfruitful with aliphatic and nitro-substituted aromatic alkynes, and due to the formation of complex mixtures, the products of internal alkynes could not be separated. It is worthy of note that other photocatalysts such as rose bengal, eosin B, eosin Y, and $\text{Ru}(\text{bpy})_3^{2+}$ were tested in this reaction and showed lower catalytic efficiency than that of rhodamine B. Of note, no product was observed in the absence of in the absence of a catalyst. The authors proposed mechanism for this transformation is analogous to the one depicted for alkenes in Scheme 10. Almost at the same time, Zhong and co-workers disclosed that the similar reaction could also successfully carried out under conventional heating conditions ($55\text{ }^\circ\text{C}$) using 5 mol% of low-cost CuCN as a catalyst.³² Concurrently, in the same paper describing catalyst- and additive-free synthesis of β -ketophosphonates through the oxyphosphorylation of styrene derivatives bearing a leaving group at the α -position and H-phosphine oxides,¹⁹ the Lei laboratory described the synthesis of five β -ketophosphine oxides **43** in yield ranging from 48% to 71% *via* the $\text{PhB}(\text{OH})_2$ -mediated oxidative coupling of aromatic

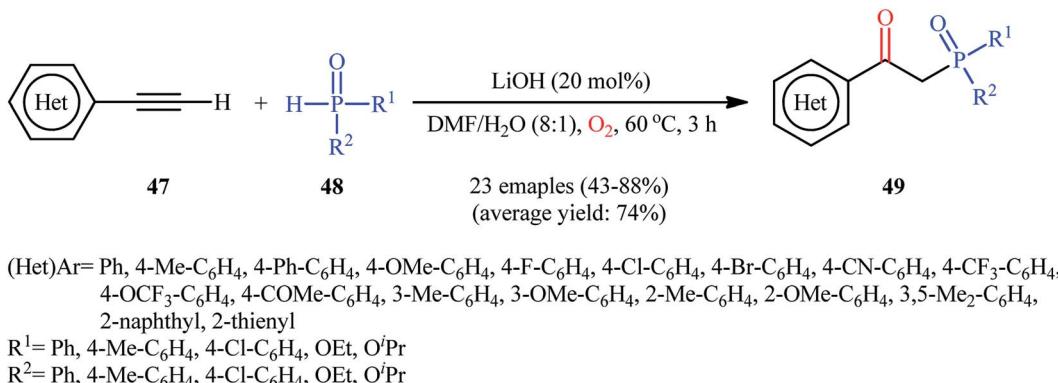
terminal alkynes **42** with diphenylphosphine oxide **8** in NMP (Scheme 16b).

Along this line, Zhang, and Cheng reported a unique and efficient protocol for the metal-free direct synthesis of diaryl (2-oxo-2-alkyl/arylethyl)phosphinates from terminal alkynes, H-phosphine oxides, and water.³³ Careful screening of a number of common oxidants such as H_2O_2 , TBHP, $\text{PhI}(\text{OAc})_2$, $\text{K}_2\text{S}_2\text{O}_8$, and *m*-CPBA; led to $\text{PhI}(\text{OAc})_2$ as the most suitable oxidant for this transformation and among the various organic solvents like THF, DCE, EtOH, DMF, MeCN, 1,4-dioxane, and toluene; MeCN proved to be the most efficient solvent. Both aromatic and aliphatic alkynes **44** and a variety of electron-rich and electron-deficient symmetrical diaryl H-phosphine oxides **45** easily took part in the reaction under optimized reaction conditions and provided the desired diaryl (2-oxo-2-alkyl/arylethyl) phosphinates **46** in good to excellent yields within 24 h (Scheme 17). This method was also applicable for gram-scale synthesis without any difficulty. The mechanistic course of this reaction is shown in Scheme 18. Initially, diaryl H-phosphine oxide **45** reacts with the phenyliodine diacetate to give hypervalent iodine intermediate **A** that, after intermolecular collapse converts to the phosphoric-carboxylic mixed anhydride intermediate **B**. Thereafter, nucleophilic attack of the water molecule on the anhydride **B** affords diarylphosphinic acid **C** that undergoes reaction with phenyliodine diacetate in



Scheme 18 Mechanism that accounts for the formation of diaryl (2-oxo-2-alkyl/arylethyl)phosphinates **46**.





Scheme 19 Base-promoted direct oxyphosphorylation of (hetero)aromatic terminal alkynes **47** with P(O)-H compounds **48**.

the presence of water to form the unstable intermediate **D**. Next, addition of hypervalent iodine reagent **D** to alkyne **44** leads to intermediate **E** which undergoes isomerization to give hypervalent iodine intermediate **F**. Finally, the reductive elimination of this intermediate **F** affords the target product **46** with the releases of indobenzene.

Recently, Shi and Zeng along with their colleagues disclosed the synthesis of β -phosphorylated ketones **49** via base-catalyzed oxyphosphorylation of (hetero)aromatic terminal alkynes **47** with P(O)-H compounds **48** (H-phosphonates, H-phosphine oxides, H-phosphinates) employing 20 mol% of LiOH as the base in an 8 : 1 mixture of DMF and H₂O at 60 °C (Scheme 19).³⁴ A wide panel of important functional groups (e.g., F, Cl, Br, CF₃, OCF₃, OMe, COMe, CN) at different positions of phenyl rings of phenylacetylenes were well tolerated by this reaction, thus indicating its broad applicability. Unfortunately, no comment was made by the authors regarding the applicability of internal and aliphatic terminal alkynes in this protocol.

4. Conclusion

Organophosphorus compounds have aroused a significant interest among synthetic chemists over the past few decades owing to their wide applications in medicinal chemistry, organic synthesis, ligand chemistry, and material science. Among them, β -phosphorylated ketones have attracted significant attention as promising building blocks for the synthesis of various important molecules due to their diverse reaction patterns. In addition to their numerous synthetic utility, some β -phosphorylated ketones themselves are endowed with outstanding biological properties. Therefore, over the years, plenty efforts have been made to identify new and efficient synthetic methods for their preparation from simple and low-cost starting materials. Among the various approaches, oxyphosphorylation of alkenes/alkynes with hydrogen phosphoryl compounds represents an extremely attractive and powerful route for the synthesis of the title compounds which offers several benefits such as high step- and atom-economy as well as the readily available starting materials. As illustrated, both internal and terminals alkenes/alkynes and all the three kinds of P(O)-H compounds (H-phosphonates, H-phosphine oxides,

and H-phosphinates) were compatible in this synthetic strategy. Moreover, most of the reactions covered in this review could be easily scaled up to provide multigram quantities of the target β -phosphorylated ketone products without any difficulty. However, despite remarkable developments over past decade on this attractive research arena, many challenges still remain to be overcome: (i) the reported examples on the use of H-phosphinates and internal alkynes as substrates are scarce and therefore, of course, expanding of the substrate scope to these compounds are necessary; (ii) the majority of metal-catalyzed reactions were conducted in the presence of copper catalysts and under relatively harsh conditions. Thus, the development of new metal-based catalytic systems that allow this difunctionalization reaction under milder conditions would be desirable; and (iii) the synthetic utility of these transformations in the preparation of biologically important compounds should be studied. We conclude this review by hoping that it will inspire researchers to make further progress in this appealing research area.

Conflicts of interest

There are no conflicts to declare.

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