

## RESEARCH ARTICLE

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View Journal | View IssueCite this: *Org. Chem. Front.*, 2025, 12, 2346Water mediated synthesis of dialkylphosphine oxides from white phosphorus and *N*-(acyloxy)phthalimides†Guo Tang, \*<sup>a,b</sup> Xuanlin Zhu, ‡<sup>a</sup> Jiali He, ‡<sup>a</sup> Yan Liu<sup>a</sup> and Yufen Zhao <sup>a,c</sup>

The multicomponent synthesis of dialkylphosphine oxides from P<sub>4</sub> without a chlorination step is reported. With the use of *N*-(acyloxy)phthalimides (NHPI esters) as the alkylation reagents, H<sub>2</sub>O as the oxygen source, tris(2,2'-bipyridine)ruthenium dichloride as a photocatalyst, 2,6-lutidine as a base, and *N,N*-dimethylacetamide as a solvent, the reactions are performed under green light irradiation, yielding the desired products in moderate to good yields. Notably, this catalytic system is also capable of synthesizing  $\alpha$ -hydroxy phosphine oxides from P<sub>4</sub> in one-pot. This two step-economic approach, which directly utilizes P<sub>4</sub> as the P-atom source, avoids the traditional chlorination stage and oxidation processes.

Received 31st December 2024,  
Accepted 4th February 2025

DOI: 10.1039/d4qo02447h

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## Introduction

Organophosphorus compounds (OPCs) containing C–P bonds are ubiquitous in both industrial and academic fields, such as materials science, pharmaceutical chemistry, catalysis, and coordination chemistry.<sup>1</sup> In general, the preparation of OPCs predominately relies on white phosphorus (P<sub>4</sub>) as the source of P atoms.<sup>2</sup> However, the conversion of P<sub>4</sub> to OPCs involves (oxy) chlorination process and uses corrosive chlorine gas (Cl<sub>2</sub>) and intermediates such as PCl<sub>3</sub>/PCl<sub>5</sub>/POCl<sub>3</sub>. Another approach toward OPCs is the hydrophosphinylation of alkenes with toxic and explosive PH<sub>3</sub>.<sup>3</sup> To circumvent the use of PCl<sub>3</sub> and PH<sub>3</sub>, significant progress has been made in forming of heteroatom–P bonds, such as S–P bonds<sup>4</sup> or O–P bonds<sup>5</sup> from P<sub>4</sub> (Scheme 1).

Dialkyl phosphine oxides [R<sub>2</sub>P(O)H] containing reactive P(O)–H bonds are versatile building blocks for synthesizing trialkyl phosphine oxides such as  $\alpha$ -hydroxy phosphine oxides. The direct transformation of P<sub>4</sub> into tertiary phosphines (PR<sub>3</sub>) and their oxides containing the same three alkyl/ary groups was investigated by the Cummins, Wolf, and Zhang groups

and others.<sup>6</sup> Due to the complex P–P bond breaking patterns, it is still challenging to construct R<sub>2</sub>P(O)H containing both C–P and P(O)–H bonds directly from P<sub>4</sub> with good selectivity. In 2022, our group made a breakthrough and published the paper entitled “Decarboxylative Selective Phosphorylation of Aliphatic Acids: A Transition-Metal- and Photocatalyst-Free Avenue to Dialkyl and Trialkyl Phosphine Oxides from White Phosphorus”.<sup>7a</sup> Since then, we have reported the selective construction of R<sub>2</sub>P(O)H from P<sub>4</sub> with alkyl bromides and iodides.<sup>7b,c</sup> All these experiments required excess alkylation reagents. Furthermore, strict anhydrous operation and subsequent oxidation workup were needed (Scheme 1B).

One-pot multicomponent reactions enable the efficient and atom-economical synthesis of OPCs directly from P<sub>4</sub>. Significant progress has been made in multicomponent reactions for the synthesis of P–heteroatom bonds (Scheme 1A).<sup>8</sup> In 2020, the first general synthesis of mixed phosphorotrithioates involving P<sub>4</sub>, disulfides, KOH, and alkyl halides was presented.<sup>8a</sup> In 2022, we described a four-component functionalization of P<sub>4</sub> with disulfides, amines and KOH to synthesize phosphoramidodithioates.<sup>8b</sup> In 2023, we published a one-pot direct synthesis of tetra-thiophosphates from P<sub>4</sub>, NaSH, disulfides and alkyl halides with sodium alkyltetra-thiophosphates as the key intermediates.<sup>8c</sup> In 2024, we presented the first copper-catalyzed three-component reaction for generating phosphorothioates from P<sub>4</sub>, disulfides, and alcohols in a single reaction step.<sup>8d</sup>

To construct C–P bonds, many novel phosphorus transfer reagents have been developed in recent years.<sup>9</sup> From the perspective of step-economy, it is highly appealing to directly convert P<sub>4</sub> into OPCs in one-pot process without the use of phosphorus transfer reagents. Compared to the research of P–heteroatom bonds formation, few multicomponent reactions

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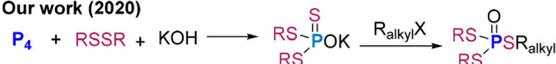
† Electronic supplementary information (ESI) available: Experimental procedures for the synthesis, spectral data and NMR spectra of compounds 3a–3o and 4a–4l. See DOI: <https://doi.org/10.1039/d4qo02447h>

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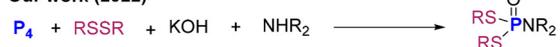


## A) Multi-component reactions for P-Hetero bonds

Our work (2020)



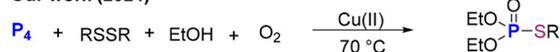
Our work (2022)



Our work (2023)

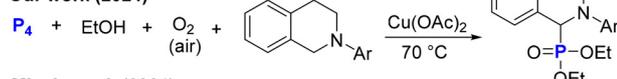


Our work (2024)

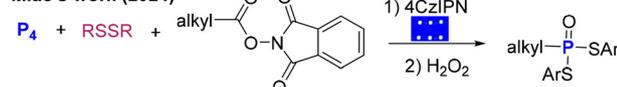


## B) Multi-component reactions for P-Carbon bonds

Our work (2024)

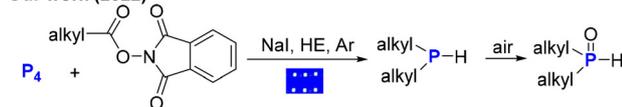


Miao's work (2024)



## C) Synthesis of dialkyl phosphines

Our work (2022)



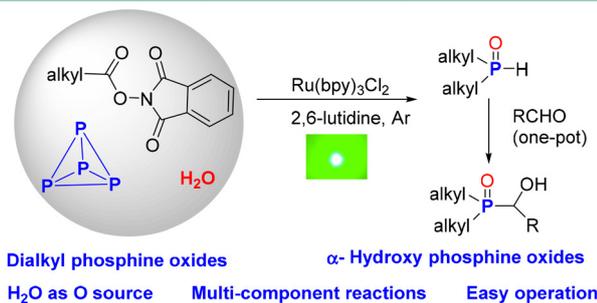
Our work (2023)



Our work (2024)



## D) This work: Multi-component reactions for dialkyl phosphine oxides

Scheme 1 Multi-component reactions with P<sub>4</sub>.

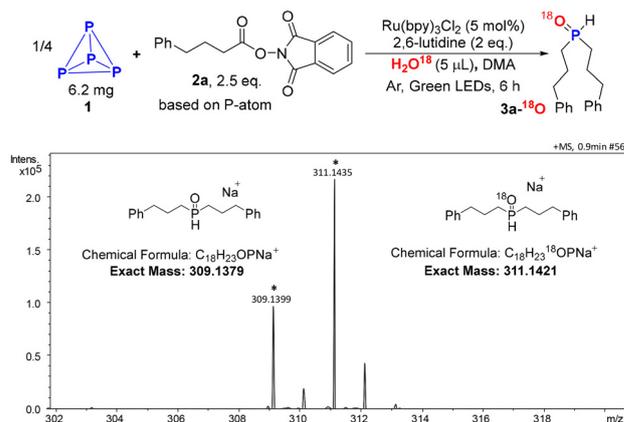
for the construction of C–P bonds have been explored (Scheme 1C). In 2024, the first multicomponent oxidative  $\alpha$ -phosphonylation of amines with P<sub>4</sub> and alcohols was developed.<sup>10a</sup> In the same year, Miao *et al.* reported a visible light-induced three-component reaction integrating P<sub>4</sub>, disulfides, and *N*-(acyloxy)phthalimides (NHPI esters) to produce alkyl-substituted phosphonodithioates.<sup>10b</sup>

As a continuum of our efforts in P<sub>4</sub> chemistry, we envisioned that with the use of H<sub>2</sub>O as the oxygen source, and *N*-(acyloxy)phthalimides as the carbon source, R<sub>2</sub>P(O)H could be directly synthesized from P<sub>4</sub> under an argon atmosphere without additional oxidation processes (Scheme 1D).

## Results and discussion

To verify our conjecture, we commenced our project by employing H<sub>2</sub><sup>18</sup>O, and 4-phenylbutanic acid NHPI ester (**2a**) as model substrates to direct functionalization of P<sub>4</sub> (**1**). When a solution of P<sub>4</sub> (6.20 mg), H<sub>2</sub><sup>18</sup>O (5  $\mu$ L), **2a** (0.5 mmol, 2.5 equivalents per P-atom), 2,6-lutidine in *N,N*-dimethylacetamide (DMA) was irradiated by green LEDs at room temperature for 6 h in the presence of Ru(bpy)<sub>3</sub>Cl<sub>2</sub> under argon atmosphere, **3a**-<sup>18</sup>O was obtained as the main product (Scheme 2).

Inspired by this result, we began optimizing the conditions for the synthesis of R<sub>2</sub>P(O)H **3a** using H<sub>2</sub>O as the oxygen source. After extensive screening of bases, solvents, and catalysts, the desired product **3a** was obtained in 90% yield (entry 1 in Table 1). No R<sub>2</sub>P(O)H and R<sub>2</sub>PH were obtained in the

Scheme 2 Water-<sup>18</sup>O as the O-resource.Table 1 Synthesis of (PhCH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>)<sub>2</sub>P(O)H (**3a**)<sup>a</sup>

Entry	Deviation from standard conditions	Yield (%)
1 <sup>b</sup>	Standard conditions	94 (90)
2	Without H <sub>2</sub> O	Trace
3	H <sub>2</sub> O (0.5 eq.)	13
4	DIPEA, DABCO instead of 2,6-lutidine	60–70
5	NMP instead of DMA	89
6	DMF, DMSO, MeCN instead of DMA	32–49
7	Other organo-PC	0–70
8	Air instead of Ar	0
9	No light or PC	0

<sup>a</sup> Reaction conditions: P<sub>4</sub> (6.20 mg, 0.20 mmol of P atom, a 0.74 M solution of P<sub>4</sub> in toluene, 0.27 mL), **2a** (2.5 equivalents per P-atom), Ru(bpy)<sub>3</sub>Cl<sub>2</sub> (0.01 mmol, 5 mol%), H<sub>2</sub>O (1.1 eq., 4  $\mu$ L), 2,6-lutidine (0.40 mmol, 2 eq.) in DMA (1 mL) irradiated by green LEDs (2  $\times$  3 W, 565–575 nm) at room temperature for 6 h under argon atmosphere. Yield of **3a** was determined by the <sup>31</sup>P{<sup>1</sup>H} NMR analysis of the crude reaction mixture using (C<sub>6</sub>H<sub>5</sub>O)<sub>3</sub>P(O) as an internal standard. <sup>b</sup> Isolated yield in parenthesis. PC = photocatalyst.

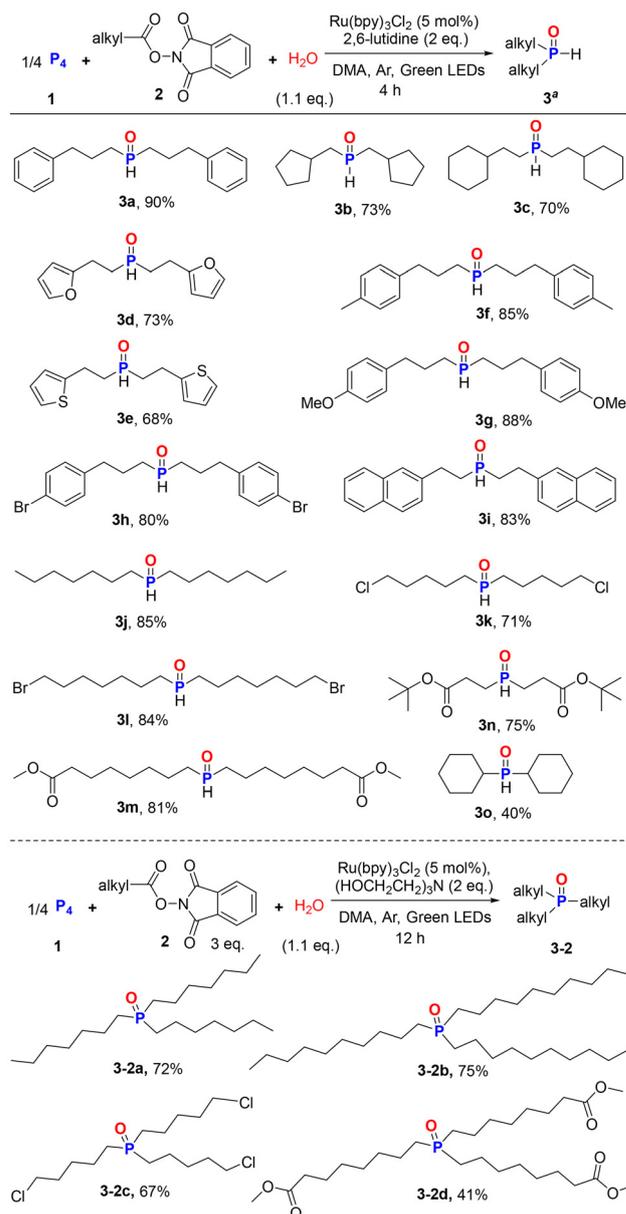


absence of H<sub>2</sub>O (entry 2). Increasing the amount of H<sub>2</sub>O to 0.5 equivalents gave **3a** but in only 13% yield (entry 3). Using H<sub>2</sub><sup>18</sup>O and H<sub>2</sub>O as the O-resources verified that water was the key starting material in this transformation. Other bases, such as DIPEA and DABCO, resulted in lower yield of **3a** (entry 4). DMA and NMP were found to be good solvents for this reaction (entry 5), while other polar solvents, such as DMF, DMSO, and MeCN, led to much lower yields (entry 6). When transition-metal-free photocatalysts were tested, most performed poorly (entry 7), although Na<sub>2</sub>-eosin Y gave the product in 70% yield. No C–P bond formation was observed when the reaction was conducted under air conditions (entry 8). The reaction was inhibited in the absence of either photocatalyst or light (entry 9).

After optimizing the reaction conditions for the synthesis of **3a**, we explored the scope of carboxylic acids for the synthesis of structurally diverse R<sub>2</sub>P(O)H **3** (Scheme 3). Under photoirradiation conditions, various alkyl acid NHPI esters can participate in this process, providing the desired products in good yields. 2-Cyclopentylacetic acid and 3-cyclohexylpropanoic acid NHPI esters gave products **3b** and **3c** in 73% and 70% yields, respectively. Linear acid NHPI esters with heterocyclic rings such as O-heterocycle (**3d**), S-heterocycle (**3e**) underwent phosphorylation without difficulty. Aryl-substituted linear acid NHPI esters with electron donating groups (methyl, methoxy, naphthalene) or electron withdrawing group (bromo) on the benzene ring gave **3f–3i** in 80–88% yields. Simple primary saturated fatty acid ester gave **3j** in 85% yield. Furthermore, a range of synthetically useful functional groups, such as C(sp<sup>3</sup>)–Cl, C(sp<sup>3</sup>)–Br, and ester (**3m** and **3n**), were also compatible with the mild conditions, providing respective dialkyl phosphine oxides in 71–85% yields. Due to steric hindrance, cyclohexanecarboxylic acid ester produced dicyclohexylphosphine oxide in a low yield (**3o**, 40%). When linear secondary or tertiary carbon radical precursors were employed, no C–P bonds were formed. In these failed experiments, white phosphorus remained completely ( $\delta = -529$  ppm). It is speculated that the sharp decrease in the yield of secondary and tertiary carbon free radical reactions may be due to the stability and steric hindrance effect of their free radicals.

We then speculated about whether this water mediated multicomponent strategy could be used to produce valuable trialkyl phosphine oxides (Scheme 3). With Ru(bpy)<sub>3</sub>Cl<sub>2</sub> as the best catalyst and trolamine as the best base, the reaction was performed for 12 h to afford the desired product **3-2a** in 72% yield. A series of chain primary fatty acid ester selected as substrates provided the corresponding trialkyl phosphine oxides **3-2b–3-2d** in satisfactory yields.

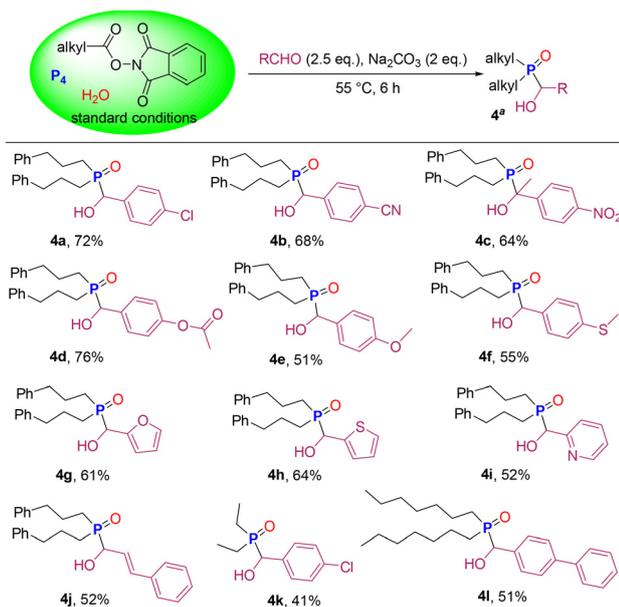
Dialkyl phosphine oxides are often used as starting materials for the preparation of various organophosphorus compounds. Encouraged by the findings described above, we continued to explore the one-pot synthesis of  $\alpha$ -hydroxy phosphine oxides from P<sub>4</sub>, avoiding the purification process of R<sub>2</sub>P(O)H and the use of dangerous Cl<sub>2</sub> and PCl<sub>3</sub>. Pleasingly, we added aldehydes and Na<sub>2</sub>CO<sub>3</sub> to the crude product **3a** solution to produce  $\alpha$ -hydroxy phosphine oxides in good yields



**Scheme 3** Scope of alkyl acid *N*-hydroxyphthalimide esters for the synthesis of dialkyl/trialkyl phosphine oxides.

(Scheme 4). Benzaldehyde with electron withdrawing group (–Cl, –CN, and –NO<sub>2</sub>) and electron donating groups (–OCOCH<sub>3</sub>, –OCH<sub>3</sub>, and –SCH<sub>3</sub>) provided the corresponding products in yields of 51–76% (**4a–4f**). When aromatic heterocyclic formaldehydes such as furan-2-carbaldehyde, thiophene-2-carbaldehyde and pyridine-2-carbaldehyde were employed as substrates, products **4g–4i** were obtained in the range of 52–64%. Cinnamaldehyde, with a double bond, afforded the product **4j** in 52% yield. This method was found to be compatible with primary saturated fatty acid esters. Decarboxylative phosphorylation of propionic acid NHPI ester followed by nucleophilic addition reaction of aldehyde gave product **4k** in 41% yield. Long-chain *n*-octoic acid NHPI ester

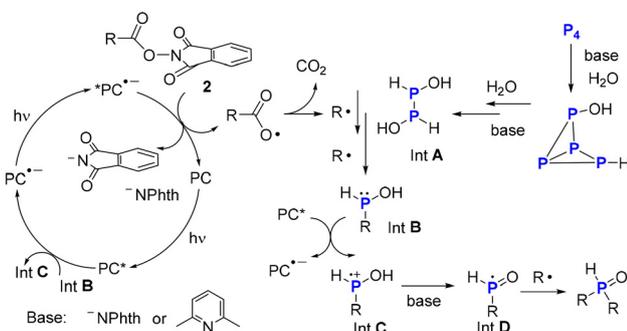




**Scheme 4** One-pot synthesis of  $\alpha$ -hydroxy phosphine oxides from  $P_4$ .

successfully participated in this one-pot synthesis of  $\alpha$ -hydroxy phosphine oxide **4l**.

Furthermore, *in situ* NMR studies on the reaction of NHPI ester **2a** and  $P_4$  were conducted and the corresponding  $^{31}P\{^1H\}$  NMR spectra were shown in Fig. 1–3 (see ESI<sup>†</sup>). When the reaction mixture was irradiated by white light-emitting diodes (LEDs) for 30 min, the signal of the product **3a** and byproduct  $RP(O)(OH)H$  appeared. No  $RPH_2$  or  $R_2PH$  was detected. Based on these results and previous studies,<sup>7</sup> a tentative mechanism for the photoinduced functionalization of  $P_4$  was proposed (Scheme 5). Initially, water breaks the P–P bond of  $P_4$  in a manner like super-basic systems ( $OH^-$  in DMSO) to yield intermediate **A**.<sup>11</sup> Subsequently, the alkyl radical breaks the P–P bond of intermediate **A** to give an unstable intermediate **B**, which then reacts with excited  $PC^*$  and base to form the phosphinoyl radical **D**. Radical **D** can directly combine with another  $R^\cdot$  yields  $R_2P(O)H$  as the main product of the reaction.



**Scheme 5** Proposed reaction mechanism.

## Conclusions

In conclusion, we developed the first water mediated multi-component synthesis of dialkylphosphine oxides and  $\alpha$ -hydroxy phosphine oxides from  $P_4$  without a chlorination step. With the use of  $N$ -(acyloxy)phthalimides as the alkylation reagents,  $H_2O$  as the oxygen source, the reactions are performed under green light irradiation. The desired dialkylphosphine oxides  $[R_2P(O)H]$  can be directly synthesized from  $P_4$  under argon atmosphere without additional oxidation processes in moderate to good yields. Advantageously, the catalytic system can also synthesize  $\alpha$ -hydroxy phosphine oxides from  $P_4$ . These step-economic approaches use  $P_4$  as the P-atom source, avoiding the traditional chlorination stage and oxidation processes.

## Experimental

### Synthesis of dialkyl phosphine oxides from white phosphorus

A Schlenk tube containing  $Ru(bpy)_3Cl_2$  (6.4 mg, 0.01 mmol, 5 mol%) and  $N$ -(acyloxy)phthalimides (0.50 mmol, 2.5 eq.) was evacuated and purged with argon three times.  $P_4$  (6.20 mg, 0.20 mmol of P atom, a 0.74 M solution of  $P_4$  in toluene, 0.27 mL),  $H_2O$  (1.1 eq., 4  $\mu$ L), 2,6-lutidine (0.40 mmol, 2 eq.) and DMA (1 mL) were sequentially added to the system at room temperature. Then the reaction mixture was stirred at room temperature under the irradiation of 6 W green LEDs (565–575 nm) for 6 h. Afterwards, the reaction mixture was quenched with the addition of saturated brine (0.5 mL), and extracted with ethyl acetate ( $3 \times 10$  mL). The combined organic layer was dried over anhydrous  $Na_2SO_4$  and then removed in a vacuum. The residue was purified by flash chromatography using DCM/MeOH (from 40 : 1 to 20 : 1, v/v) as the eluent to afford the desired dialkyl phosphine oxides **3a–3o**.

### One-pot synthesis of $\alpha$ -hydroxy phosphine oxides from white phosphorus

After the above-mentioned reaction mixture was stirred at room temperature under the irradiation of 6 W green LEDs (565–575 nm) for 6 h without any work-up, aldehyde (0.50 mmol, 2.5 eq.) and  $Na_2CO_3$  (42.40 mg, 0.4 mmol, 2.0 eq.) were added in air. The reaction mixture was stirred for 22 hours at  $55\text{ }^\circ\text{C}$  for 6 hours. After completion, saturated brine (0.5 mL) was added; the mixture was diluted by EtOAc (10.0 mL). The mixture was extracted with ethyl acetate ( $3 \times 10$  mL). The combined organic layer was dried over anhydrous  $Na_2SO_4$  and then removed in a vacuum. The residue was purified by flash chromatography using DCM/MeOH (from 40 : 1 to 10 : 1, v/v) as the eluent to afford the desired  $\alpha$ -hydroxy phosphine oxides **4a–4l**.

## Data availability

The data supporting this article have been included as part of the ESI.<sup>†</sup>



## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

We gratefully acknowledge financial support from the National Key Research and Development Program of China (grant no. 2020YFA0608300), the Space Application System of China Manned Space Program (grant no. KJZ-YY-WSM01), and National Natural Science Foundation of China (grant no. 21772163, 21778042, and 41876072).

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