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Photocatalytic stannylation of white phosphorus†

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Organophosphorus compounds (OPCs) are highly important chemicals, finding numerous applications in both academia and industry. Herein we describe a simple photocatalytic method for the stannylation of white phosphorus (P4) using a cheap, commerciallyavailable distannane, (Bu₃Sn)₂, and anthraquinone as a simple photocatalyst. Subsequent 'one pot' transformation of the resulting stannylated monophosphine intermediate (Bu₃Sn)₃P provides direct, convenient and versatile access to valuable OPCs such as acylated phosphines and tetraalkylphosphonium salts.

White phosphorus (P_4) – the most chemically important allotrope of this ubiquitous and abundant element - acts as the common precursor from which all commercially valuable and academically important organophosphorus compounds (OPCs) are prepared. The current methods used for the industrial synthesis of these myriad useful P1 products include the oxidation of P4 with toxic Cl2 gas to generate PCl3 which can subsequently be transformed into a variety of OPCs by reaction with nucleophiles (Scheme 1a). As an alternative route, initial acid- or base-mediated disproportionation of P4 can be used to generate highly toxic PH3 gas which is then employed for the hydrophosphination of unsaturated organic substrates.¹

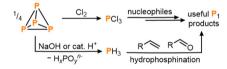
Given the drawbacks of these methods, a highly prominent aim has long been to find ways of bypassing these multi-step procedures. In particular, there is a longstanding desire to develop more step-efficient direct - and, ideally, catalytic methods to functionalize P4 and generate OPCs in a single reaction.

As a result, for several decades comprehensive efforts have been made to better understand the fundamental reactivity of P₄.² However, it is only very recently that it has finally become

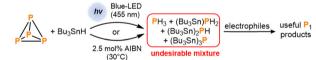
possible to successfully transform P4 directly into a variety of useful P₁ products.³ Moreover, and despite these extensive investigations, the number of successful examples remains extremely low, and those that do exist still suffer from substantial limitations.4 As such, there remains a clear need to expand the range of strategies available for direct, productive P₄ activation, with new catalytic methods being particularly desirable.4a

In one of our own contributions to this area, we recently reported a simple 'one pot' method in which the classical radical reagent trin-butyltin hydride (Bu₃SnH) is used for initial hydrostannylation of

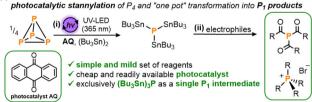
(a) State-of-the-art: indirect industrial routes to P1 products



(b) Hydrostannylation of P_4 and "one pot" transformation into P_1 products



(c) This work photocatalytic stannylation of P₄ and "one pot" transformation into P₄ products



Scheme 1 (a) Current state-of-the-art industrial methods for the synthesis of valuable P₁ products. (b) Recently reported hydrostannylation of white phosphorus (P₄) using Bu₃SnH followed by reaction with electrophiles to generate useful P_1 products in a 'one-pot' fashion.⁵ (c) This work: (i) photocatalytic stannylation of P4 using the photocatalyst anthraquinone (AQ) and hexabutyldistannane (Bu₃Sn)₂; and (ii) subsequent functionalization of the intermediate (Bu₃Sn)₃P with electrophiles into products such as triacylphosphines and tetraalkylphosphonium salts

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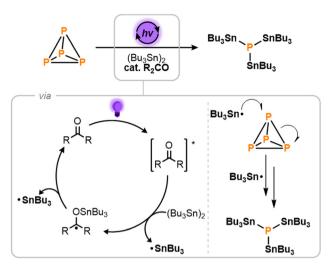
P₄ (Scheme 1b).⁵ This reductive P₄ activation is mediated either by light or by a chemical radical initiator such as AIBN (azobis(isobutyronitrile)) which can initiate a radical chain reaction that breaks down the P4 tetrahedron, yielding a mixture of hydrostannylated phosphines $(Bu_3Sn)_xPH_{3-x}$ (x = 0-3). Key to this mechanism is the attack of stannyl radicals (Bu₃Sn[•]) on the P-P bonds of P₄. The resulting (Bu₃Sn)_xPH_{3-x} mixture can then be converted into a number of important and useful OPCs by reaction with electrophiles.5

Unfortunately, one significant disadvantage of this hydrostannylation strategy is the complexity of the (Bu₃Sn)_xPH_{3-x} mixture, which complicates 'downstream' reaction development by requiring functionalization of two different types of bond (P-Sn and P-H), both of which are distributed over four distinct molecules. Moreover, the presence of gaseous PH₃ as a component of this mixture has been suggested to have a limiting effect on overall yields as it can easily be lost during subsequent manipulations, 4a and it is also problematic from a safety perspective.

These drawbacks would be overcome if the initial P₄ reduction step could instead furnish a single species with just one functionalizable motif, but with reactivity otherwise similar to (Bu₃Sn)_xPH_{3-x}. To achieve this, we describe herein a simple photocatalytic strategy for the atom-precise stannylation of P₄ using the cheap, commercially-available distannane (Bu₃Sn)₂ and simple benzophenone derivatives as photocatalysts (Scheme 1c). This new procedure generates exclusively the stannylated monophosphine (Bu₃Sn)₃P and subsequent, simplified 'one pot' transformations with electrophiles afford valuable OPCs including acylated phosphines and alkylated phosphonium salts.

Based on the analysis above, we sought to develop a new method by which P4 could be selectively transformed into (Bu₃Sn)₃P as the sole product.⁶ It is worth noting that the closely related product (Ph₃Sn)₃P has previously been prepared from P4 using Ph3SnCl as the stannylating reagent, but this required use of a relatively elaborate Ti(III) reagent as a halogen atom abstractor.^{3f} Instead, we imagined that an ideal reagent for such a reaction would be the distannane (Bu₃Sn)₂, which is cheap to purchase and could in principle provide the target phosphine with perfect atom economy. Indeed, Sn-Sn homolysis of (Bu₃Sn)₂ is known to furnish Bu₃Sn[•] radicals, which previous work has shown are capable of adding to P₄. ^{3f,5} However, achieving this homolysis directly requires extreme temperatures or very high energy UV light irradiation that is known to lead to unselective reactivity, and is also unlikely to be compatible with P4.8-10 Fortunately, it has been reported that simple ketones can be used as photocatalysts to access Bu₃Sn[•] radicals by Sn-Sn bond cleavage under much lower energy irradiation.11

The light-driven photocatalytic stannylation of P₄ was therefore targeted, based on the mechanistic proposal outlined in Scheme 2.9 It was anticipated that photoirradiation of the ketone R₂CO would first provide an excited state, [R₂CO]*, 12 capable of reacting with (Bu₃Sn)₂ to generate a stannylated ketyl radical and a free Bu₃Sn• radical.¹¹ The former could then



Scheme 2 Proposed mechanism for the light-driven, photocatalytic stannylation of P₄ in the presence of hexabutyldistannane, (Bu₃Sn)₂, and a ketone photocatalyst, R2CO

thermally release a second Bu₃Sn[•] radical to close the catalytic cycle. Once formed, these Bu₃Sn[•] radicals would then add to the P-P bonds of P4, ultimately breaking it down to generate (Bu₃Sn)₃P as the only P₁ product.¹³

To begin, benzophenone (BP) was chosen as a proof-ofprinciple photocatalyst due to both its simplicity and the fact that its photoreactivity towards hexaalkyldistannanes has been studied previously. 11d Gratifyingly, after an initial optimization the photocatalytic stannylation of P4 could successfully be achieved, with use of 25 mol% BP (all stoichiometries, in both equiv. and mol%, are defined per P atom) and a 3.3-fold excess (5 equiv.) of (Bu₃Sn)₂ providing 50% conversion to the target stannylated phosphine (Bu₃Sn)₃P after stirring under near UV LEDs overnight (Scheme 3; see also ESI,† S3). Control experiments confirmed that all reaction components (P4, (Bu₃Sn)₂, BP, irradiation) were necessary for the reaction to proceed productively (see ESI,† S3, Table S1).

These initial results provided a clear proof-of-principle for the proposed mechanistic strategy. Notably, the observed conversion indicates the activation of at least three Sn-Sn bonds per available equivalent of BP, 14 making this a rare example of a system where P₄ activation has been achieved catalytically, using an otherwise inert substrate. 5,9a,9b,9e,15 Nevertheless, in order to improve the reaction outcome further, a broader range of benzophenone derivates was subsequently screened, with several found to provide markedly improved performance (see ESI,† S5). Particularly impressive results were achieved using



Scheme 3 Initial conditions for the direct, photocatalytic stannylation of P₄ into (Bu₃Sn)₃P optimized using benzophenone (**BP**) as photocatalyst. Stoichiometries in equiv. and mol% are defined per P atom.

UV-LED (365 nm) benzene/acetone, 22 h AQ (12.5 mol%) (Bu₃Sn)₂ (3 equiv.)

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Scheme 4 Optimized conditions for the direct, photocatalytic stannylation of P_4 into $(Bu_3Sn)_3P$ using anthraquinone (AQ) as photocatalyst. Stoichiometries in equiv. and mol% are defined per P atom.

anthraquinone (AQ) and following brief further optimization (see ESI,† S5 and S7) 79% conversion to $(Bu_3Sn)_3P$ could be achieved using significantly reduced loadings of both AQ (12.5 mol%) and $(Bu_3Sn)_2$ (3 equiv.) over the same timeframe (Scheme 4; see also ESI,† S7). Based on the catalytic cycle proposed in Scheme 2, this would correspond to a turnover number (TON) of 10.0 for AQ. Further reductions in catalyst loading to 6.3 mol% or 2.5 mol% were found to lead to even higher TONs (16.8 and 28.2, respectively), albeit at the cost of lower overall conversions (see ESI,† S7, Table S11).

With the stannylation of P4 optimized, attention was then shifted to its subsequent, 'one pot' transformation into other useful P₁ products. Having previously developed procedures for the analogous transformation of the phosphine mixture (Bu₃Sn)_xPH_{3-x} which includes (Bu₃Sn)₃P as a minor component, it was anticipated that addition of electrophiles to photocatalytically-generated (Bu₃Sn)₃P should be similarly productive, ^{4,5} especially since neither the AO photocatalyst nor the (Bu₂Sn)₂ starting material is expected to show appreciable reactivity towards such substrates. And, indeed, in situ addition of a variety of acid chlorides yielded the corresponding triacylphosphines $(R(O)C)_3P$ (R = Ph, Cy, Ad, tBu, iPr, nBu, iPrMe) with good conversions of up to 75% (Scheme 5a(i)). 5,16 Notably, and in comparison to our previously-reported hydrostannylation system, no exclusion of light and no additional base were required for this step, highlighting both the robustness and simplicity of $(Bu_3Sn)_3P$ as a "P³⁻" synthon, relative to $(Bu_3Sn)_xPH_{3-x}$.

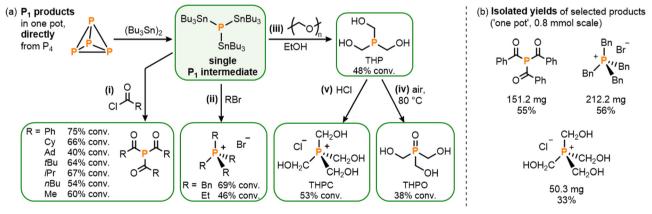
Similarly, reaction of $(Bu_3Sn)_3P$ with alkyl bromides RBr (R = Bn, Et) under moderate heating successfully provided

'one pot' access to the corresponding phosphonium salts, $[R_4P]Br$, including tetrabenzylphosphonium bromide, $[Bn_4P]Br$, which is a known precursor for useful Wittig chemistry (Scheme 5a(ii)). Again, no auxiliary base was required for these reactions, in contrast to the analogous procedures *via* $(Bu_3Sn)_xPH_{3-x}$ where the absence of base leads to a 50% reduction in yield. 5

Finally, another industrially important class of P₁ products was targeted. Hydroxymethyl-substituted phosphine derivatives are used as flame-retardant materials (among a number of other applications),¹⁸ and could be accessed by reacting the stannylated monophosphine (Bu₃Sn)₃P with paraformaldehyde in EtOH to furnish tris(hydroxymethyl)phosphine, (HOCH₂)₃P (THP; Scheme 5a(iii)).^{18a} Subsequent exposure to air then yielded the corresponding phosphine oxide, (HOCH₂)₃PO (THPO; Scheme 5a(iv)),^{18b} while the phosphonium salt tetrakis(hydroxymethyl)phosphonium chloride, [(HOCH₂)₄P]Cl (THPC),^{18c,18d} could be accessed by quenching the *in situ* generated THP with HCl, all in one pot (Scheme 5a(v)).

To demonstrate the viability of these reactions on a preparative scale the triacylphosphine $(Ph(O)C)_3P$ and the phosphonium salts $[Bn_4P]Br$ and THPC were selected as representative examples for isolation (Scheme 5b; see ESI† S9). At 0.8 mmol scale $(PhC(O))_3P$ could be isolated in 55% yield, ¹⁹ which compares well with our previously-reported hydrostannylation method (51%). $[Bn_4P]Br$ could also be isolated in good 56% yield, and THPC in a more modest yield of 33%. ¹⁹

For this last reaction, efforts were also made to recover the Sn-containing compounds present at the end of the reaction. We have previously shown that for the analogous synthesis of THPC *via* (Bu₃Sn)_xPH_{3-x} recovery of the Bu₃SnCl byproduct allows for convenient regeneration and recycling of the Bu₃SnH starting material, thus minimizing the formation of organotin-containing waste. Bu₃SnCl can also be used to regenerate (Bu₃Sn)₂ through a net one-electron reduction,⁸ meaning similar recycling should be feasible for this newer system, provided Bu₃SnCl can again be cleanly recovered. Satisfyingly, Bu₃SnCl could indeed be recovered during THPC workup through simple washing with diethyl



Scheme 5 (a) One-pot synthesis directly from P_4 , via photocatalytically generated P_1 intermediate $(Bu_3Sn)_3P$, of (i) triacylphosphines $(R(O)C)_3P$ (4 equiv. RC(O)Cl, R = tBu, Ph, Me, nBu, Cy, iPr, Ad), (ii) phosphonium salts $[R_4P]Br$ (5 equiv. RBr, R = Bn, Et, 60-80 °C), (iii) tris(hydroxymethyl)phosphine, THP (EtOH, 3 equiv. paraformaldehyde), (iv) tris(hydroxymethyl)phosphonium chloride, THPC (as for (iii) using 12.5 equiv. paraformaldehyde, then 10 equiv. HCl); and (b) Isolated yields for reactions on preparative scale (0.8 mmol). Stoichiometries in equiv. are defined per P atom.

ether, being isolated as part of an otherwise clean mixture with unreacted (Bu₃Sn)₂ in an excellent overall yield of 92% (1.3:1 molar ratio, see ESI† S9).

In conclusion, we have developed a simple, new method for the direct transformation of P4 into a variety of commercially and academically interesting OPCs. The reaction proceeds through a photocatalytic stannylation of white phosphorus, which generates (Bu₃Sn)₃P with perfect atom economy as a single, convenient P₁ intermediate using an inexpensive, commercially available distannane and a simple photocatalyst. This method can be used to prepare a variety of different products through inclusion of a range of different electrophilic substrates, and we have demonstrated that the Sn-containing byproducts of the reaction can in principle be recovered and recycled. These results expand the currently very limited range of strategies that are available for the direct functionalization of P₄, and suggest the intriguing possibility that P₄ activation might also be achievable by reaction with other weak E-E bonds under similar photocatalytic conditions.

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Conflicts of interest

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There are no conflicts to declare.

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