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Synthesis of 1,2,4-azadiphosphole derivatives based on vanadium-catalyzed [2+2+1] cycloaddition reactions of azobenzenes with phosphaalkynes†

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A new synthetic method is described to construct 1,2,4-azadiphosphole derivatives based on vanadium-catalyzed [2+2+1] cycloaddition reactions. Reactions of azobenzenes as nitrogen sources with phosphaalkynes as phosphorous counterparts in the presence of VCl$_2$(thf)$_2$ as a catalyst afford the corresponding 1,2,4-azadiphospholes.

Phosphorus heterocycles have provided important structural motifs to explore materials science and coordination chemistry.[1] To access diverse skeletons, development of new and efficient synthetic methods is quite important. In addition to the conventional synthetic methods typically forming carbon–phosphorous single bonds, the use of phosphaalkynes as substrates is beneficial to construct phosphorous-containing π-systems.[2] Stoichiometric reactivity of phosphaalkynes with transition metal complexes has been studied extensively,[3] but catalytic reactions of phosphaalkynes under transition metal catalysis have been limited to several sporadic examples.[4,5] Recently, our group has reported catalytic [2+2+2] cycloaddition reactions to produce phosphabenzenes and [3+2] cycloaddition reactions to produce 1,3-azaphospholes based on the use of phosphaalkynes as substrates.[5] To investigate further utility of phosphaalkynes to prepare aromatic compounds containing phosphorus atoms, we have focused on the synthesis of 1,2,4-azadiphospholes as next targets.
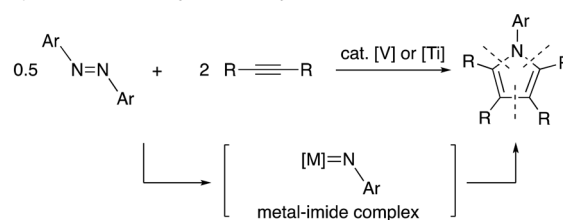
Synthetic examples of 1,2,4-azadiphospholes have been limited to only a few reports.[6,7] To the best of our knowledge, the first synthesis of the 1,2,4-azadiphosphole skeleton was reported in 1991, where a 1,2,4-azadiphosphole derivative was prepared based on a thermal dimerization of an amino-substituted phosphaalkyne.[6a] Later, other groups reported stoichiometric reactions of titanium- and vanadium-imide complexes as nitrogen sources with phosphaalkynes.[6b–d] Although these stoichiometric reactions provided effective

synthetic methods of 1,2,4-azadiphospholes, transition metal-catalyzed synthesis of 1,2,4-azadiphospholes has never been achieved until now.
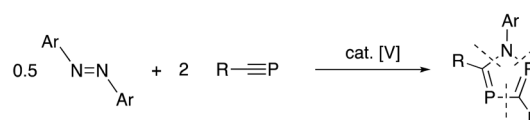
Recently, titanium-catalyzed [2+2+1] cycloaddition reactions have been reported by Tonks and co-workers to prepare pyrroles from reactions of azobenzenes with alkynes (Scheme 1a).[8] In this reaction system, titanium-imide species generated from azobenzenes worked as a key intermediate. More recently, a similar pyrrole synthesis using vanadium-catalyzed reaction system has been reported by Tonks, Mashima, Tsurugi and co-workers.[9]

Based on the research background, we have envisaged metal-catalyzed [2+2+1] cycloaddition reactions of azobenzenes with phosphaalkynes to produce 1,2,4-azadiphospholes (Scheme 1b). As a result, we have found that some vanadium complexes worked as effective catalysts toward the formation of 1,2,4-aza-diphospholes. Herein, we report experimental results in detail.

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Scheme 1 Catalytic [2+2+1] cycloaddition reactions with azobenzenes as nitrogen sources in heterocycle synthesis.

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First, investigation of reaction conditions was carried out with the use of azobenzene (**1a**) and 1-adamantylphosphaethyne (**2a**) as typical substrates. When the reaction of **1a** (1 equiv.) with **2a** (2 equiv.) in the presence of 20 mol% of VCl$_3$(thf)$_3$ was carried out in toluene at 110 °C for 16 hours, the desired 1,2,4-azadiphosphole (**3a**) was obtained in 48% yield (Table 1, entry 1). Replacing VCl$_3$(thf)$_3$ with VBr$_3$(thf)$_3$ and VCl$_3$(py)$_3$ was not effective in the current reaction system (Table 1, entries 2 and 3). Then, we investigated the use of VCl$_2$(thf)$_2$, VCl$_2$(1,4-dioxane)$_2$ and VI$_2$(thf)$_4$. Among these catalysts, VCl$_2$(thf)$_2$ showed the highest reactivity to afford the desired product **3a** in 60% yield (Table 1, entries 4–6). Next, addition of ligands was examined. When bidentate phosphine ligands such as 1,1-bis(diphenylphosphino)methane (dppm) or 1,2-bis(diphenylphosphino)ethane (dppe) was used, only lower yields of the product were obtained (Table 1, entries 7 and 8) while no formation of the desired product was observed when 2,2′-bipyridine was used as a ligand (Table 1, entry 9). Separately, we confirmed no formation of **3a** in the absence of vanadium complexes. Based on these results, we have chosen the use of VCl$_2$(thf)$_2$ as the optimal catalyst.

With the optimized reaction conditions in hand, the use of other substrates was investigated. The use of various functionalized azobenzene derivatives was examined (Scheme 2). When 3-methyl-, 3-ethyl- and 3-isopropyl-substituted azobenzenes were used, the corresponding products (**3b**, **3c** and **3d**) were obtained in 55%, 51% and 46% yield, respectively. Effect of alkyl groups at the 4-position of the benzene rings in azobenzene was also examined. Introduction of methyl, ethyl, isopropyl, *tert*-butyl, *sec*-butyl and *n*-butyl groups afforded the corresponding products (**3e–3j**) in good yields. The reaction system was tolerant toward oxygen functional groups such as 4-methoxyl and 4-methoxycarbonyl groups to give the desired products (**3k** and **3l**) in 52% and 47% yields, respectively. Unfortunately, an
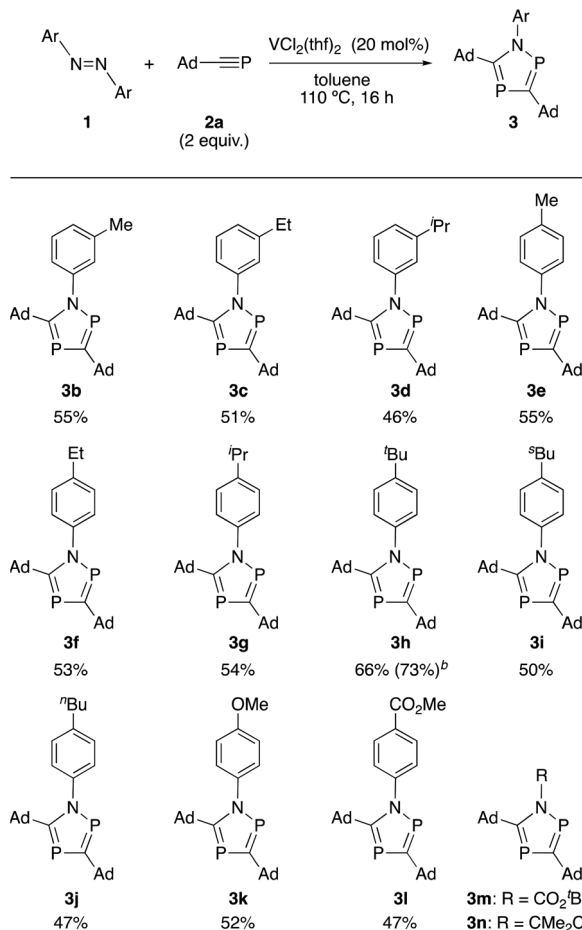
Scheme 2 Reactions of azobenzene with 1-adamantylphosphaethyne **2a**. [a] Reactions of azobenzenes **1** (0.1 mmol) with **2a** (0.2 mmol) in the presence of VCl$_2$(thf)$_2$ (20 mol%) in toluene (2.0 mL) at 110 °C for 16 h. [b] ^{1}H NMR yield with C$_6$Me$_6$ as an internal standard.

azodicarboxylate ester was not applicable in this reaction system resulting in no formation of the corresponding product (**3m**). We also examined 2,2′-azobis(isobutyronitrile) as an aliphatic azo compound, but the corresponding product (**3n**) was not obtained at all.

Next, the use of various phosphaalkynes was investigated (Scheme 3). A phosphaakyne bearing 3,5-dimethyl-1-adamantyl moiety was successfully transformed into the corresponding product (**3o**) in 50% yield. The use of phosphaalkyne bearing a benzene ring was also successful to afford the corresponding product (**3p**) in 41% yield. Introduction of various alkoxyphenyl moieties into the phenyl ring was also investigated and the corresponding products were obtained in moderate yields (**3q–3t**). When the biphenyl substituent was introduced in phosphaalkyne, the corresponding product (**3u**) was obtained in 41% yield.

As mentioned above, recently vanadium-catalyzed [2+2+1] cycloaddition reactions of azobenzenes with alkynes to afford the pyrroles were reported.[9] In this reaction system, bis-(trimethylsilyl)aniline was used as an additive along with VCl$_3$(thf)$_3$ catalyst, and a vanadium-bis(imide) complex was proposed to be a catalytically active species. Bis(trimethylsilyl)aniline

Table 1 Investigation of reaction conditions[a]

Entry	[V] catalyst	Ligand	Yield (%)
1	VCl$_3$(thf)$_3$	—	48
2	VBr$_3$(thf)$_3$	—	42
3	VCl$_3$(py)$_3$	—	0
4	VCl$_2$(thf)$_2$	—	60
5	VCl$_2$(1,4-dioxane)$_2$	—	47
6	VI$_2$(thf)$_4$	—	27
7	VCl$_2$(thf)$_2$	dppm	27
8	VCl$_2$(thf)$_2$	dppe	20
9	VCl$_2$(thf)$_2$	2,2′-Bipyridine	0

[a] Reactions of **1a** (0.10 mmol) with **2a** (0.20 mmol) in the presence of a catalyst (20 mol%) and a ligand (20 mol%) in toluene (2.0 mL) at 110 °C for 16 h.
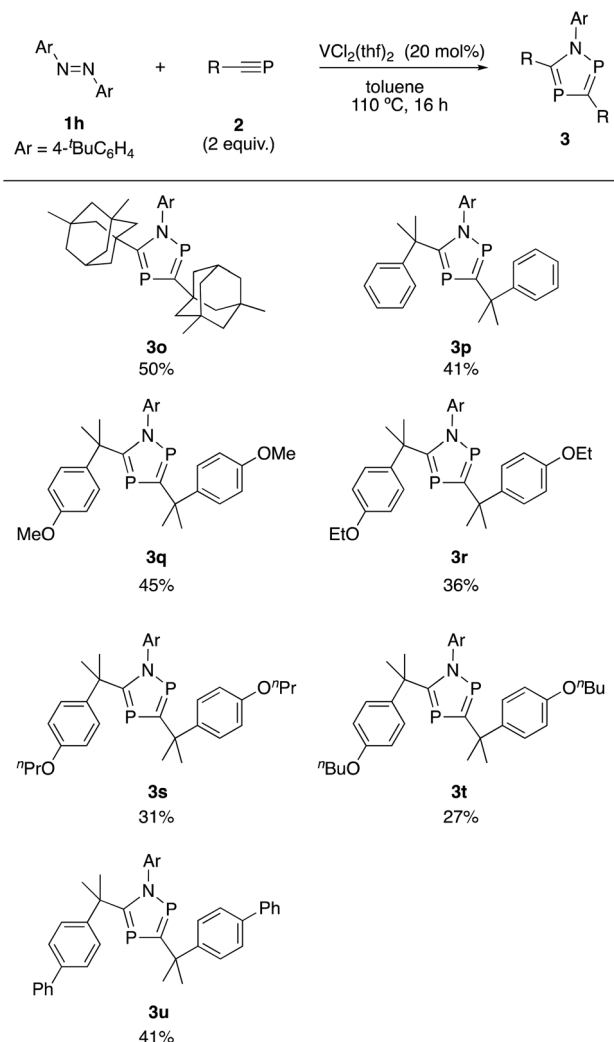
Scheme 3 Reactions of 1,2-di(4-tert-butyl)diazene **1h** with phosphaalkynes **2**. [a]Reactions of **1h** (0.1 mmol) with phosphaalkynes **2** (0.2 mmol) in the presence of VCl$_2$(thf)$_2$ (20 mol%) in toluene (2.0 mL) at 110 °C for 16 h.

supplies an imide moiety on the vanadium center through a redox neutral ligand exchange process and the second imide moiety comes from azobenzene substrates through an oxidative process, where the generated vanadium-bis(imide) complex is supposed to undergo further reactions to give the pyrroles.
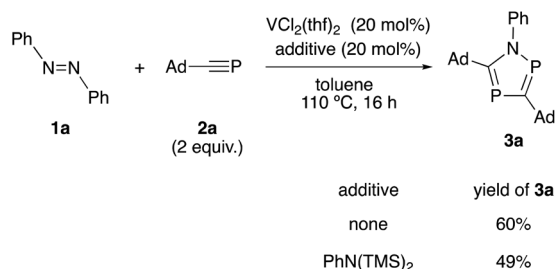
Scheme 4 Effect of bis(trimethylsilyl)aniline as additive.
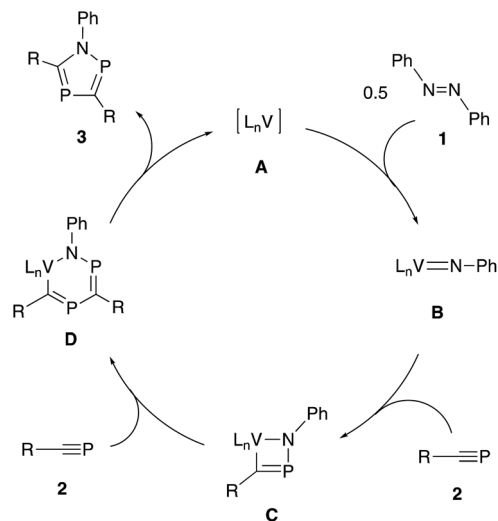
Scheme 5 Plausible reaction pathway.

According to the experimental result, we investigated the effect of bis(trimethylsilyl)aniline as an additive in the current reaction system (Scheme 4). When bis(trimethylsilyl)aniline (20 mol%) was added along with VCl$_2$(thf)$_2$ catalyst in the reaction of **1a** with **2a**, the yield of **3a** did not improve. This result suggests that the introduction of imide moiety from bis(trimethylsilyl)aniline was not effective in this reaction system. Previously, vanadium-mono(imide) complexes were reported to react stoichiometrically with phosphaalkynes to give the corresponding 1,2,4-azadiphospholes.[6b,c] Based on the experimental result, we consider that a vanadium-mono(imide) species may work as a catalytically active species in the current reaction system.

Based on these observations, a plausible reaction pathway is shown in Scheme 5. First, vanadium complex **A** reacts with an azobenzene **1** to give vanadium-imide complex **B**. Then, [2+2] cycloaddition reaction occurs between **B** and a phosphaalkyne **2** to afford a four-membered ring intermediate **C**. Subsequent insertion of the second phosphaalkyne **2** forms a six-membered ring intermediate **D**. Finally, reductive elimination affords the desired 1,2,4-azadiphosphole **3** accompanied by the regeneration of the initial vanadium complex **A**.

In summary, we have succeeded in the development of the first vanadium-catalyzed [2+2+1] cycloaddition reactions of azobenzenes with phosphaalkynes to synthesize a variety of 1,2,4-azadiphospholes. We believe the present study has opened a new aspect of synthetic application for the utilization of phosphaalkynes as building blocks in the construction of phosphorous-containing heterocyclic skeletons.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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