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# Air and water stable secondary phosphine oxides as diazaphospholene precatalysts†

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Air-stable secondary phosphine oxides (SPOs) are readily formed from diazaphospholene bromides. In the presence of pinacolborane, these SPOs are transformed into catalytically active diazaphospholene hydrides. A silyl triflate transforms the SPOs into phosphonium triflates. The use of diazaphospholene SPOs as reduction reaction precatalysts was validated by imine reduction, conjugate reduction, pyridine hydroboration, and asymmetric reduction.

Diazaphospholenes have recently emerged as a class of catalyst for reduction reactions. Gudat showed that highly air-sensitive diazaphospholene hydrides were stoichiometric reducing reagents for aldehydes, ketones, and conjugate acceptors.<sup>1</sup> Kinjo showed that ammonia borane could regenerate a diazaphospholene hydride in the reduction of an azo compound, allowing catalysis with diazaphospholenes.<sup>2a</sup> Kinjo then showed that pinacolborane (HB(pin)) and diphenylsilane could be terminal reductants in aldehyde and carbon dioxide reductions, respectively. Importantly, pinacolborane transformed an alkoxydiazaphospholene to a diazaphospholene hydride in Kinjo's work (eqn (1), Fig. 1).<sup>2b,c</sup> Our research group has exploited the alkoxydiazaphospholene intermediate from Kinjo's aldehyde-reduction catalytic cycle as a precatalyst for diazaphospholenes. We used this system to catalyze imine reductions and render Gudat's conjugate reductions catalytic.<sup>3a</sup> Kinjo and our group separately reported that diazaphospholenes catalyze the reduction of pyridines, with the distinct difference that our group used a neutral precatalyst, while Kinjo used a cationic triflate.<sup>4</sup> We subsequently employed the pre-catalyst strategy to form a chiral diazaphospholene for imine reduction.<sup>3b</sup> Cramer disclosed an elegant chiral diazaphospholene precatalyst based on an alkoxydiazaphospholene for conjugate reduction and Claisen reactions.<sup>5</sup> Other reported

Previous diazaphospholene (and arsene) pre-catalyst strategies:



Fig. 1 Comparison of diazaphospholene precatalysts.

systems relevant to diazaphospholene pre-catalysts include Radosevich's trisamidophosphine that forms a diazaphospholene in the presence of HB(pin),<sup>6</sup> and diazaarsolenes bearing alkoxy substituents reported by Melen. Arsenic hydrides have not been isolated from these systems, however these arsenic catalysts do engage in reduction catalysis in analogy to the diazaphospholene systems.<sup>7</sup> Yang and Cheng showed that diazaphospholenes are the most hydridic neutral compounds yet quantified on the Mayr nucleophilicity scale, promising a rich future for diazaphospholenes as reduction catalysts and reagents.<sup>8</sup>

With this intense interest in diazaphospholenes, further developments to increase the accessibility of these catalysts in organic synthesis is warranted. This work shows that the secondary phosphine oxide (SPO) hydrolysis products of diazaphospholene alkoxydes or halides are air and water stable, can be purified by silica gel chromatography, and can serve as precatalysts for diazaphospholene hydrides (eqn (2), Fig. 1) and the corresponding phosphonium triflates.

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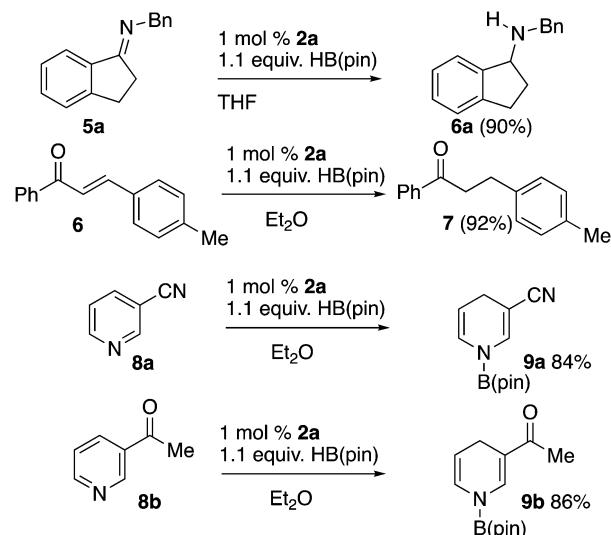
Scheme 1 Synthesis of a diazaphospholene SPO, and its transformation to hydride.

During our initial investigations of diazaphospholenes, we observed that samples of diazaphospholene precatalyst **1a** in standard NMR tubes hydrolyzed to SPO **2a** because of the incursion of water (Scheme 1).<sup>3</sup> SPOs of diazaphospholenes have been rationally synthesized as ligands for metals *via* hydrolysis of amidodiazaphospholene **1b**.<sup>9</sup> We also noted that when HB(pin) was present with **1a**, SPO formation was not observed on the same timeframe. We considered that HB(pin) could either be scavenging the water or could be reducing the SPO. While this work was in progress, Webster and co-workers reported the reduction of dialkyl and diaryl SPOs directly to uncomplexed phosphines using pinacolborane at room temperature.<sup>10</sup>

In a more direct synthesis of **2a** we avoided the preparation of intermediate **1a** or **1b** by combining diazaphospholene bromide **3a** with triethylamine in dichloromethane followed by addition of water to afford SPO **2a**. Compound **2a** is a solid that was stable to column chromatography in air (Scheme 1). NMR spectral data matched that previously reported for **2a**.<sup>9a</sup> Additionally, a crystal of **1a** grown from ethyl acetate/hexanes has the same unit cell as that previously reported for crystals of **2a**.<sup>9a,b</sup> HB(pin) transforms **2a** to hydride **4a** in various solvents, including benzene-*d*<sub>6</sub>, and THF. While 1 equivalent of HB(pin) was sufficient for the transformation, the reaction proceeded significantly faster when greater than 1 equivalent of HB(pin) was used.

We used **2a** in reductions employing HB(pin) that are known to be catalyzed by diazaphospholenes, and have been documented to not undergo an uncatalyzed reaction with HB(pin) to validate the function of **2a** as a precatalyst (Scheme 2). Imine **5a** was reduced to the corresponding amine with HB(pin) in the presence of 1 mol% of **2a**.<sup>3a</sup> Chalcone **6** was reduced to compound **7** with 1 mol% of **2a**.<sup>3a,5a,11</sup> Precatalyst **2a** also effected the reduction of pyridines with HB(pin). Nicotinonitrile **8a** was reduced to dihydropyridine **9a**, while 3-acetylpyridine **8b** was reduced to dihydropyridine **9b** without reduction of the ketone. These results show similar chemoselectivity and selectivity to previous pyridine reductions performed with precatalyst **1a**.<sup>4</sup>

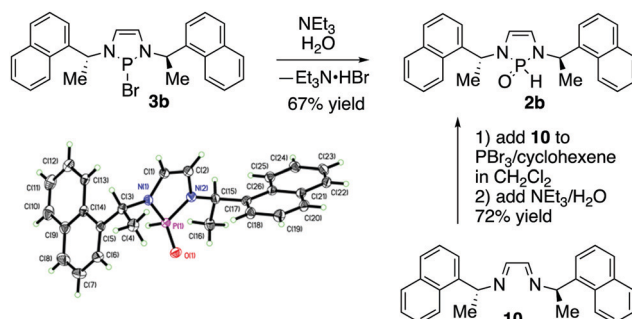
We subsequently explored the chemistry of a chiral SPO precatalyst (Scheme 3). Compound **2b** was prepared from the hydrolysis of the corresponding bromide and was also an air and water stable substance that could be purified by silica gel chromatography.<sup>3b</sup> In a further simplification of the procedure,



Scheme 2 Reduction reactions with SPO precatalyst **2a**.

isolation of **3b** could be completely avoided. Diimine **10** was cyclized with PBr<sub>3</sub> and cyclohexene according to MacDonald and co-workers' procedure for diazaphospholene generation.<sup>12</sup> After one hour, triethylamine and water were added directly to the reaction mixture. After aqueous work-up and chromatography, **2b** was obtained in good yield. This procedure allowed preparation of **2b** in one pot from starting material **10**, without the isolation of any air-sensitive intermediates. Prior NMR spectroscopy of **3b** had shown equivalence of the naphthylethyl sidechains and back-bone protons, implying a *C*<sub>2</sub> symmetric structure, which strongly suggests that the P-Br bond is ionized in solution, resulting in planarization of the phosphorus.<sup>3b,13</sup> NMR spectroscopy of **2b** shows less symmetry: the naphthylethyl side-chains and back-bone protons of the diazaphospholene are now inequivalent, a consequence of the breaking of *C*<sub>2</sub> symmetry because of the distinct H and O phosphorus substituents in the SPO. X-ray crystallography confirmed the structure of **2b**.<sup>14</sup>

Asymmetric reduction reactions were explored with chiral precatalyst **2b** (Scheme 4). Reduction of imine **5b** to amine **6b** (eqn (1), Scheme 4) proceeded with comparable enantioselectivity to that previously observed with chiral diazaphospholenes with an 85:15 e.r. at room temperature, and an increase to 92:8 e.r. at -35 °C.<sup>16</sup> A limitation for the use of **2b** was revealed when



Scheme 4 Asymmetric catalysis with SPO precatalyst **2b**.

sterically hindered imine **5c** gave a low yield of essentially racemic product **6c** under these conditions (eqn (2), Scheme 4). To further understand these distinctive results, we repeated the experiments in NMR tubes and observed the reaction progress by  $^{11}\text{B}$  and  $^{31}\text{P}$  NMR spectroscopy. After the components were mixed, the reaction of eqn (1) showed almost complete consumption of HB(pin) according to the  $^{11}\text{B}$  NMR spectrum after 5 minutes. The  $^{31}\text{P}$  NMR spectrum of this reaction showed a mixture of precatalyst **2b** and hydride **4b**. The reaction of eqn (2) exhibited substantially different behavior. The  $^{11}\text{B}$  NMR spectrum showed essentially no conversion after both 5 minutes and two hours, with mostly unreacted HB(pin) present. At both the 5 minute and two hour mark, the  $^{31}\text{P}$  NMR spectrum of the reaction of eqn (2) showed a relatively complex mixture of phosphorus containing products, including putative **4c**, and  $\text{PH}_3$ . These endocyclic cleavage products have previously been noted in the decomposition of diazaphospholenes, and do not appear to be catalytically active in imine reduction.<sup>3</sup> Accordingly, with a readily reduced substrate such as **5b**, pre-catalyst activation and imine reduction occur, while with a challenging substrate such as **5c**, reduction does not occur before decomposition of the catalyst. The observed formation of **6c** in the reaction of eqn (2) is presumably due to an uncatalyzed reduction of the imine with HB(pin) mediated by the water introduced during the quenching of the reaction.<sup>15</sup>

Our recently reported cationic diazaphospholene catalysts are able to reduce substrate **5c** in a highly enantioselective manner, showing that **2b** has inferior reactivity.<sup>16</sup>

Given this limitation, we sought to determine if **2a** and **2b** could be used to generate phosphonium cations *in situ*, enabling access to more active phosphonium triflate reduction catalysts. Exposure of **2a** or **2b** to TBS triflate in dry chloroform-*d* on an NMR scale immediately resulted in the formation of phosphonium ions **10a** and **10b**, as evidenced by appearance of a signal in the  $^{31}\text{P}$  NMR spectrum at +202.3 ppm for **2a**, and 208.6 ppm for **2b** (Scheme 5). Additionally, the formerly diastereotopic naphthylethyl groups in **2b** became equivalent in the  $^1\text{H}$  NMR spectrum, indicating the generation of  $C_2$  symmetry by planarization of the phosphorus centre. The mixture containing **2a** proved competent in

Scheme 5 Catalysis with phosphonium cations generated from SPO precatalysts **2a** and **2b**.

the reduction of pyridine **8c** to the dihydropyridine **9c**. Pyridines such as **8c** with no electron withdrawing group would not be expected to readily undergo reduction with neutral diazaphospholenes.<sup>5b</sup> Kinjo has effected reductions of such pyridines employing phosphonium cations.<sup>5a</sup> In the chiral system, exposure of **2b** to TBS triflate, followed by addition of imine **5c**, and HB(pin) restored asymmetric induction with **5c**, leading to selective formation of **6c**. This contrasts with the absence of selectivity observed with precatalyst **2b** for the reduction of **5c** in the absence of the silyl triflate. The observed reactivity and selectivity are close to that previously observed in the reduction of compound **5c** catalyzed by isolated phosphonium ion **10b**, and it would be expected that the large scope of substrates reduced by isolated **10b** could also be prepared under these *in situ* conditions. In addition, endocyclic cleavage of **10b** with excess HB(pin) has never been noted, enabling the reduction of more challenging substrates such as **5c** to compete with catalyst decomposition.<sup>16</sup>

In conclusion, we have validated the use of SPOs derived from diazaphospholenes as precatalysts for the catalytically active diazaphospholene hydrides. Given growing interest in diazaphospholenes, the demonstration that secondary phosphine oxides can serve as bench-purifiable and air stable precatalysts will further help to popularize diazaphospholene chemistry. The stability of these entities to chromatography will enable the separation of diazaphospholene precursors from complex product mixtures, potentially allowing access to diazaphospholenes that cannot currently be accessed cleanly. We have also demonstrated generation of phosphonium cations from diazaphospholene SPOs, showing how these highly active asymmetric reduction catalysts can also be accessed from air-stable precursors.

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## Conflicts of interest

The authors declare the following competing financial interest(s): Dalhousie University has filed patents on chiral phosphonium triflates and secondary phosphine oxides for asymmetric imine reduction, from which royalty payments may be derived.

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