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Diazomethane umpolung atop anthracene: an electrophilic methylene transfer reagent†

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Formal addition of diazomethane's terminal nitrogen atom to the 9,10-positions of anthracene yields $\text{H}_2\text{CN}_2\text{A}$ (**1**, $\text{A} = \text{C}_{14}\text{H}_{10}$ or anthracene). The synthesis of this hydrazone is reported from Carpino's hydrazine $\text{H}_2\text{N}_2\text{A}$ through treatment with paraformaldehyde. Compound **1** has been found to be an easy-to-handle solid that does not exhibit dangerous heat or shock sensitivity. Effective umpolung of the diazomethane unit imbues **1** with electrophilicity at the methylene carbon center. Its reactivity with nucleophiles such as $\text{H}_2\text{C}(\text{PPh}_3)$ and N-heterocyclic carbenes is exploited for $\text{C}=\text{C}$ bond formation with elimination of dinitrogen and anthracene. Similarly, **1** is demonstrated to deliver methylene to a nucleophilic singlet d^2 transition metal center, $\text{W}(\text{ODipp})_4$ (**2**), to generate the robust methyldiene complex [**2** $=\text{CH}_2$]. This behavior is contrasted with that of the Wittig reagent $\text{H}_2\text{C}(\text{PPh}_3)$, a more traditional and Brønsted basic methylene source that upon exposure to **2** contrastingly forms the methyldiyne salt $[\text{MePPh}_3][\text{2}\equiv\text{CH}]$.

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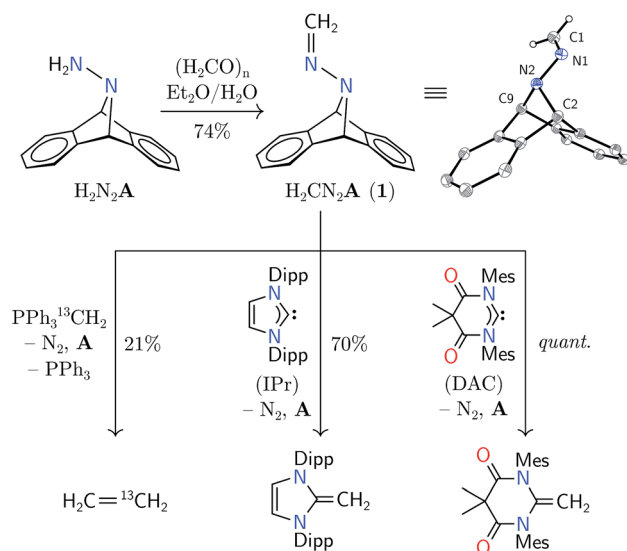
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Diazomethane is infamous for the dangers associated with its use.¹ Despite its synthetic versatility, diazomethane's high toxicity and propensity to explode should give a chemist pause before committing to its use. In an effort to offer an alternate methylene source using an anthracene-based strategy,^{2–8} we report herein the synthesis and some initial reactivity studies of $\text{H}_2\text{CN}_2\text{A}$ (**1**, $\text{A} = \text{C}_{14}\text{H}_{10}$ or anthracene), a molecule conceived as a formal adduct between diazomethane and anthracene. An initial survey of the reactivity patterns of **1** has revealed it not to be a simple substitute for diazomethane, instead characterizing it as a unique electrophilic methylene source. Its electrophilicity differentiates **1** from common metal-free methylene transfer reagents such as diazomethane and methylene triphenylphosphorane.

Synthesis of hydrazone **1** proceeded from Carpino's hydrazine $\text{H}_2\text{N}_2\text{A}$ upon paraformaldehyde treatment in a biphasic diethyl ether–water mixture,^{9,10} providing the target molecule in 74% isolated yield (Scheme 1). An X-ray diffraction study of its structure revealed expected metrical data.¹¹

Hydrazone **1** was found to be an air-stable and crystalline solid, easily manipulable and displaying no propensity for detonation upon heating or shock. The solid was found to be volatile by thermogravimetric analysis (TGA), which showed gradual sample evaporation up to 120 °C without any discrete

mass-loss events that would be expected from its fragmentation into diazomethane and anthracene. Within a sealed capillary, **1** melted without explosion (116–119 °C). After heating the melt to 140 °C, NMR spectroscopic analysis of the resolidified solid showed 74% recovery of **1** with 26% anthracene production. Its behavior in solution was similar, evincing only slow



Scheme 1 Synthesis of methylene hydrazone **1** and initial studies of methylene transfer (Mes = mesityl, Dipp = 2,6-diisopropylphenyl), shown alongside its structure from an X-ray diffraction study. Thermal ellipsoids are shown at the 50% probability level. Selected distances [Å] and angles [°]: N1–C1 1.275(2), N1–N2 1.389(1), N2–C2 1.508(1), N2–C9 1.521(2); N2–N1–C1 118.3(1).

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fragmentation into anthracene at temperatures greater than 120 °C. The volatility of this compound foiled attempts at analysis of its thermal behavior by molecular beam mass spectrometry (MBMS), limiting our ability to comment on the fragments directly produced by its thermal fragmentation.^{2–5}

Having established **1** to pose a low explosion risk, we were encouraged to proceed to test its reactivity as a methylene synthon. Our initial investigations rapidly uncovered contrasting reactivity patterns *vis-à-vis* those characteristic of diazomethane. For example, methylation of carboxylic acids, a hallmark of diazomethane reactivity,¹² did not proceed upon treatment with excess pivalic acid, acetic acid, or trifluoroacetic acid. These experiments were informative, and led us to consider more closely the electronic structure of **1**.

Hydrazones are known to be carbon ambiphiles;¹³ however, **1** did not demonstrate nucleophilicity. Such behavior is not unexpected, as the π_{CN} is known to be polarized away from the carbon center, although less so than an imine π_{CN} or a ketone π_{CO} bond.¹⁴ The polarization of this bond suggests that **1** should be expected to exhibit moderate electrophilicity at its methylene carbon. This would effectively induce umpolung of the diazomethane unit as diazomethane generally reacts as a carbon nucleophile.¹⁵

The predicted reversal of philicity was initially confirmed by successful methylene transfer in the reaction between **1** and H₂CPPH₃. Combination of these two reagents in benzene-d₆ yielded ethylene in 21% yield over 12 h in concert with anthracene, triphenylphosphine, and, presumably, dinitrogen. The reaction was found to produce several unidentified byproducts by NMR spectroscopy, explaining the low yield of ethylene; however, isotopic labelling of the ylide led to H₂C=¹³CH₂ from **1** and H₂¹³CPPH₃, and H₂C=CD₂ from **1** and D₂CPPH₃, confirming ethylene formation through the unification of the electro- and nucleophilic methylene units. Although the yield was low, this mode of reactivity was instructive for our further studies.

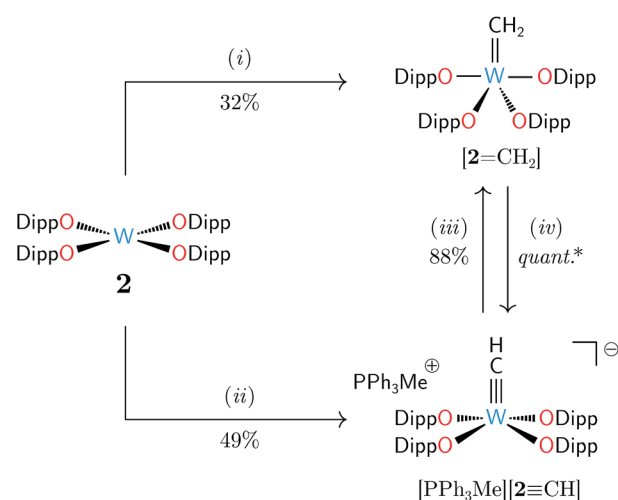
The electrophilicity of **1** lent itself well to the synthesis of N-heterocyclic olefins from N-heterocyclic carbenes (NHCs).¹⁶ In benzene-d₆ solution, **1** reacted with nucleophilic IPr (1,3-bis(2,6-diisopropylphenyl)imidazol-2-ylidene) to yield the corresponding olefin in 70% yield after 13 h at 80 °C.¹⁶ As a nucleophile with increased electrophilicity, the Bielawski *N,N'*-diamidocarbene (“DAC”) was found to react in essentially quantitative yield to form a new C=C bond over 24 h at 22 °C.¹⁷ This mode of reactivity differs markedly from that of diazoalkanes, which have been documented to react with NHCs at their electrophilic N-terminus to produce azines with a new C=N–N=C moiety.¹⁸ Heating **1** with triphenylphosphine or tricyclohexylphosphine has not yielded the analogous yields, suggesting a modest Lewis acidity at the carbon center of **1**.

It is rare for diazomethane to be used in transition metal chemistry for the synthesis of a stable methyldene complex.¹⁹ In fact, the use of diazoalkanes in d-block chemistry is often complicated by their propensity for side reactions other than alkylidene delivery.^{20,21} The reactivity differences between **1** and diazomethane thus encouraged us to attempt the use of **1** in methyldene complex synthesis to see if engagement of the

terminal nitrogen in bonding to anthracene subdues deleterious alternate reaction pathways.

We identified [W(ODipp)₄] (**2**, ODipp = 2,6-diisopropylphenoxide)^{22,23} as a d² transition metal complex well poised to behave as a methylene acceptor.²⁴ Complex **2** is synthetically easy to access, and its square-planar geometry features a nucleophilic lone pair of electrons housed in a metal-centered d_{z²}-like orbital, analogously to related tantalum and molybdenum singlet d² species.^{8,25} Treatment of **2** with excess **1** gave facile formation of the anticipated methyldene complex [2=CH₂] after mild heating in benzene to 55 °C for 35 h (Scheme 2). Characteristically deshielded proton and carbon resonances of the CH₂ unit were found by NMR spectroscopy: ¹H δ 8.95 ppm and ¹³C δ 232.9 ppm with scalar coupling constants of ²J_{WH} = 156.0 Hz, ¹J_{WC} = 185.0 Hz, and ¹J_{CH} = 155.6 Hz. The ¹J_{CH} coupling constant was typical of metal alkylidenes lacking significant agostic character.^{26–29} The success of **1** in this capacity was exciting, as the rarity of terminal, isolable methyldene complexes³⁰ makes new methods for their generation welcome developments.

Crystallization from pentane at –35 °C overnight enabled an X-ray diffraction study of [2=CH₂] (Fig. 1, left) that confirmed the molecular structure. Although the data were not of high quality, the coordination geometry about the tungsten center was unambiguously identified to be intermediate between square pyramidal and trigonal bipyramidal ($\tau = 0.48$),³¹ and the alkylidene bond was identified with a W⋯C interatomic distance of 1.864(4) Å. This bond length is typical of a W=C double bond³² and similar to values reported for other tungsten(vi) methyldenes.^{29,33–35} Compound [2=CH₂] was not found to react productively with ethylene or 1-hexene upon heating to 70 °C in benzene-d₆ for 18 h, confirmed by a lack of isotopic migration from [2=¹³CH₂] to the olefins.³⁶ Under these conditions, [2=CH₂] also did not react with mesitaldehyde or 4,4'-



Scheme 2 Comparative reactivity of W(ODipp)₄ (**2**): (i) H₂CN₂ (**1**, 10 equiv.), benzene, 55 °C, 35 h; (ii) H₂CPPH₃ (2.0 equiv.), THF, 25 °C, 30 min; (iii) lutidinium triflate (1.0 equiv.), THF, 25 °C, 5 min; (iv) PPh₃CH₂ (1.0 equiv.), THF, 25 °C, 30 min. (*) NMR spectroscopic analysis showed (iv) to be quantitative.



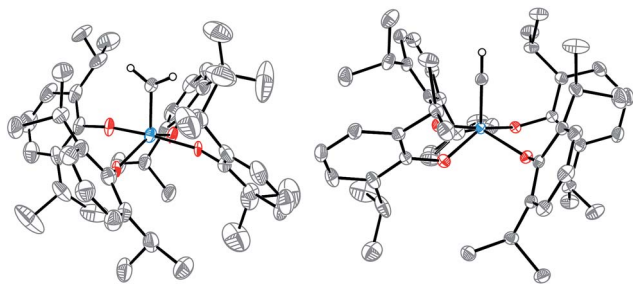


Fig. 1 Molecular structures of (left) $[2=CH_2]$ and (right) $[MePPh_3][2\equiv CH]$ from single-crystal X-ray diffraction studies shown with thermal ellipsoids at the 50% probability level. All hydrogen atoms except for the methyldene and methynide hydrogens are omitted for clarity, as is the $[MePPh_3]$ cation. Interatomic distances for tungsten-carbon multiple bonds: (left) $W1=C1$ 1.864(4) Å, (right) $W1\equiv C1$ 1.749(1) Å.

dimethylbenzophenone to form $[2\equiv O]$ and the corresponding olefins. Despite this, $[2=CH_2]$ is notable as an example of a methyldene complex with aryloxides as the exclusive supporting ligands. As such, it is an interesting structural model for methyldene complexes supported by silica or alumina surfaces implicated in alkane or olefin metathesis.^{37–39}

The reactivity of **1** was particularly satisfying after discovery of the contrasting behavior of H_2CPh_3 , a known reagent for CH_2 delivery to transition metal centers.^{40–42} Treating a solution of **2** with H_2CPh_3 (1 equiv.) in THF at 25 °C rapidly consumed 50% of **2** and formed the methynide salt $[MePPh_3][2\equiv CH]$. Doubling the amount of H_2CPh_3 gave total consumption of **2** and provided $[MePPh_3][2\equiv CH]$ in 49% isolated yield (Scheme 2). Variation of the stoichiometry and temperature of this reaction did not lead to conditions for $[2=CH_2]$ formation, indicating competitive deprotonation of intermediate $[2=CH_2]$ by Brønsted basic H_2CPh_3 . Such acid-base chemistry is postulated to play a critical role in the formation of surface-bound alkylidenes and alkynylidynes for alkane and olefin metathesis,^{37,38} meaning $[2=CH_2]$ serves also as an interesting reactivity model for alkynidyne synthesis mediated through proton transfer. This was corroborated by independent deprotonation of $[2=CH_2]$ with H_2CPh_3 , and highlights the utility of **1** as a weakly Brønsted basic source of methylene. Protonation of $[MePPh_3][2\equiv CH]$ using lutidinium triflate presents a complementary route to $[2=CH_2]$.

An X-ray crystallographic study of $[MePPh_3][2\equiv CH]$ revealed a $W\cdots C$ interatomic distance of 1.749(1) Å and a square pyramidal ($\tau = 0.21$) coordination geometry about tungsten. A search of the CSD revealed this to be the first catalogued example of a structurally characterized metal methynidyne in an all-oxygen ligand environment, and the first catalogued example of a tungsten(vi) methynidyne complex.

As interest in metal methyldene species is rapidly growing both in homogeneous and heterogeneous catalysis,^{37–39,43–45} we hope **1** can be further exploited in their syntheses. Compound **1** has also shown promise in formation of new C=C bonds with H_2CPh_3 and NHCs, and may find use in construction of terminal olefins.

Conflicts of interest

There are no conflicts to declare.

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