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# Direct observation of *o*-benzyne formation in photochemical hexadehydro-Diels–Alder (*hν*-HDDA) reactions†

Xiaonan Ma,<sup>ab</sup> Jan Maier,<sup>id</sup> c Michael Wenzel,<sup>a</sup> Alexandra Friedrich,<sup>id</sup> c Andreas Steffen,<sup>cd</sup> Todd B. Marder,<sup>id</sup> \*c Roland Mitrić,<sup>id</sup> \*a and Tobias Brixner<sup>id</sup> \*a

Reactive *ortho*-benzyne derivatives are believed to be the initial products of liquid-phase [4 + 2]-cycloadditions between a 1,3-diyne and an alkyne *via* what is known as a hexadehydro-Diels–Alder (HDDA) reaction. The UV/VIS spectroscopic observation of *o*-benzyne derivatives and their photochemical dynamics in solution, however, have not been reported previously. Herein, we report direct UV/VIS spectroscopic evidence for the existence of an *o*-benzyne in solution, and establish the dynamics of its formation in a photoinduced reaction. For this purpose, we investigated a bis-diyne compound using femtosecond transient absorption spectroscopy in the ultraviolet/visible region. In the first step, we observe excited-state isomerization on a sub-10 ps time scale. For identification of the *o*-benzyne species formed within 50–70 ps, and the corresponding photochemical hexadehydro-Diels–Alder (*hν*-HDDA) reactions, we employed two intermolecular trapping strategies. In the first case, the *o*-benzyne was trapped by a second bis-diyne, *i.e.*, self-trapping. The self-trapping products were then identified in the transient absorption experiments by comparing their spectral features to those of the isolated products. In the second case, we used perylene for trapping and reconstructed the spectrum of the trapping product by removing the contribution of irrelevant species from the experimentally observed spectra. Taken together, the UV/VIS spectroscopic data provide a consistent picture for *o*-benzyne derivatives in solution as the products of photo-initiated HDDA reactions, and we deduce the time scales for their formation.

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## Introduction

After the discovery of a [4 + 2]-cycloaddition reaction between a conjugated diene and an alkene by Diels and Alder in 1928,<sup>1</sup> it took almost 70 years until Johnson<sup>2–6</sup> and Ueda<sup>7–11</sup> explored the [4 + 2]-cycloaddition between a diyne and an alkyne. This latter reaction is now known as a “hexadehydro-Diels–Alder” (HDDA) reaction so named by the Hoya group in 2012 as a logical extension of the dehydro-Diels–Alder reaction.<sup>12</sup> Instead of forming cyclohexene derivatives, the HDDA reaction<sup>13–16</sup> is believed to produce a highly reactive *o*-benzyne intermediate, and is of much current interest due to its unusual reactivity/mechanism<sup>17–27</sup> and numerous

applications in synthetic chemistry.<sup>28–35</sup> A series of reports within the last three years,<sup>36–43</sup> examining the HDDA reaction from a more mechanistic viewpoint, demonstrate the immense versatility of HDDA-generated *o*-benzynes, including reaction with an alkyne forming a highly reactive benzocyclobutadiene<sup>21</sup> or reaction with electron-deficient thioamides giving stabilized ammonium ylides in a new type of [3 + 2]-cycloaddition.<sup>24</sup> These and other examples emphasize the importance of HDDA-generated *o*-benzynes for the discovery of new reaction pathways.

Although the HDDA reaction is becoming increasingly useful in synthesis, intriguing questions remain. Does *o*-benzyne really form as an intermediate during the HDDA reaction? Can we confirm the existence of *o*-benzyne derivatives as initial products of the HDDA reaction in solution? In reported investigations on the HDDA reaction, *o*-benzyne was inferred indirectly by “back-tracking” its structure from the products resulting from trapping reactions, *i.e.*, a secondary product of the HDDA reaction.<sup>14–16</sup> A trapping reaction of unique interest is the double hydrogen transfer, first reported in 2009 by Tsui and Sterenberg.<sup>44</sup> This transfer of two vicinal hydrogen atoms from solvents was further investigated in more detail by Hoya *et al.*<sup>17</sup> correlating the reactivity of different solvents with the degree of eclipsing of two adjacent C–H bonds among their low-energy conformers. A subsequent promising trapping reaction is that of perylene with

<sup>a</sup>Institut für Physikalische und Theoretische Chemie, Universität Würzburg, Am Hubland, 97074 Würzburg, Germany. E-mail: xiaonanma@tju.edu.cn; brixner@uni-wuerzburg.de; roland.mitric@uni-wuerzburg.de

<sup>b</sup>Institute of Molecular Plus, Tianjin University, No. 92 Weijin Road, Nankai District, 300072 Tianjin, China

<sup>c</sup>Institut für Anorganische Chemie, Institute for Sustainable Chemistry & Catalysis with Boron, Universität Würzburg, Am Hubland, 97074 Würzburg, Germany. E-mail: todd.marder@uni-wuerzburg.de

<sup>d</sup>Institut für Anorganische Chemie, Technische Universität Dortmund, Otto-Hahn-Str. 6, 44227 Dortmund, Germany

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traditional<sup>45,46</sup> and HDDA-generated<sup>47</sup> *o*-benzyne derivatives, generating new polycyclic aromatic hydrocarbons. However, trapping experiments do not provide a direct observation of the proposed *o*-benzyne intermediate, and UV/VIS spectroscopic evidence for *o*-benzyne in solution remains elusive.

The formation, characterization, and application of benzyne has fascinated chemists for more than 90 years due to their Kekulé-type diradical structure, special role in combustion and astrochemistry, and great potential in synthetic chemistry.<sup>48–53</sup> *o*-Benzyne was first proposed as an intermediate in 1927 by Bachmann and Clarke,<sup>54</sup> and IR,<sup>55–61</sup> microwave,<sup>62–66</sup> NMR,<sup>67,68</sup> and photoelectron<sup>69–74</sup> spectra of *o*-benzyne have been reported in the last half century. A recent report,<sup>75</sup> however, explains that the NMR data<sup>68</sup> of an *o*-benzyne generated inside of a hemicarcerand do not belong to an *o*-benzyne inside the molecular cage, but instead, the signals result from a species formed from a reaction of the *o*-benzyne with the molecular cage. This had been one of the very few reports of the spectroscopic identifications of an *o*-benzyne in “solution” at room temperature. There are only few investigations of electronic absorption spectra in the ultraviolet/visible (UV/VIS) range, in the gas phase or in low-temperature matrices.<sup>76–82</sup> To the best of our knowledge, the electronic absorption spectrum, *i.e.*, the UV/VIS spectrum, of *o*-benzyne in solution, has not been reported.

In addition to the thermal HDDA reaction discussed above, the photochemical HDDA (*hν*-HDDA) reaction has also been reported recently, initiated by UV excitation at low temperature (−70 °C). The reaction cascade and corresponding trapping product were found to be identical to those formed thermally.<sup>83</sup> Extending the previous work which employed continuous-wave illumination, we show herein that using time-resolved excitation and transient absorption spectroscopy makes the direct detection of *o*-benzyne possible, and allows us to deduce their formation dynamics. In our study, we employed ultrafast spectroscopy that has proven to be a powerful tool by which to elucidate chemical reaction mechanisms, identifying the product as well as the pathways leading to it.

Thus, we performed ultrafast pump-probe (transient absorption) spectroscopy in the UV/VIS region and accurate quantum chemical calculations to explain the observed transient spectra in order to investigate the excited-state dynamics and photochemical reaction of bis-diyne compound **Me-BD-Me** (Fig. 1, top left), where Me is −CH<sub>3</sub> and the bis-diyne (BD) is −4-C<sub>6</sub>H<sub>4</sub>−(C≡C)<sub>2</sub>−(CH<sub>2</sub>)<sub>3</sub>−(C≡C)<sub>2</sub>−4-C<sub>6</sub>H<sub>4</sub>−. We combine transient absorption measurements with intermolecular trapping strategies (Fig. 1, bottom) in tetrahydrofuran (THF) solution (path 1), using self-trapping (paths 2 and 3), and the reaction with perylene (path 4). Thus, we obtained spectroscopic evidence for the existence of the *o*-benzyne *hν*-HDDA cascade and recorded the highly plausible transient UV/VIS absorption spectrum of an *o*-benzyne species in solution for the first time.

## Results and discussion

### Transient absorption experiment

The transient UV/VIS absorption spectra of **Me-BD-Me** in THF following UV excitation at a central wavelength of  $\lambda = 295$  nm

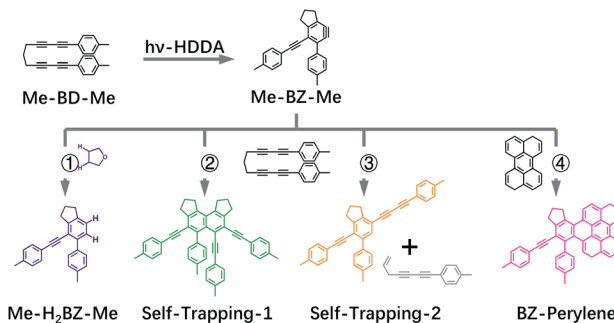


Fig. 1 Compounds and reactivity scheme. Chemical structure of the bis-diyne investigated, 1,11-bis(*p*-tolyl)undeca-1,3,8,10-tetrayne (**Me-BD-Me**, top left) with Me = −CH<sub>3</sub>, BD = −4-C<sub>6</sub>H<sub>4</sub>−(C≡C)<sub>2</sub>−(CH<sub>2</sub>)<sub>3</sub>−(C≡C)<sub>2</sub>−4-C<sub>6</sub>H<sub>4</sub>−. We also show the *hν*-HDDA reaction-generated *o*-benzyne species (**Me-BZ-Me**, top center) and the following trapping reactions with tetrahydrofuran (bottom left, path 1), a second **Me-BD-Me** (self-trapping, bottom center, path 2 and 3), and perylene (bottom right, path 4).

are shown in Fig. 2, which display the optical density difference spectra ( $\Delta OD$ ) in the 320–670 nm regime. The transient absorption spectra of **Me-BD-Me** in CHCl<sub>3</sub> and CH<sub>3</sub>CN are displayed in Fig. S1† with similar manner.

The transient absorption signal is dominated by a cascaded sequence of positive peaks extending from the VIS to the near-UV range with dynamic changes observed from sub-picosecond to several-nanosecond time scales. The positive peaks can be attributed to excited-state absorption of initially populated

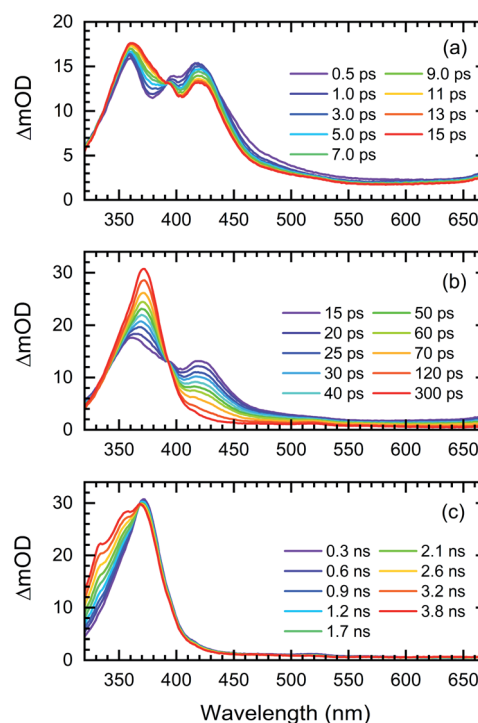


Fig. 2 Transient UV/VIS absorption spectra in 320–670 nm regime. Data are shown for **Me-BD-Me** dissolved in THF upon photoexcitation at 295 nm in the delay ranges of (a) 500 fs to 15 ps, (b) 15 ps to 300 ps, and (c) 300 ps to 3.8 ns.



states as well as the ground-state and excited-state absorption resulting from intermediates or final photoproducts, to be assigned below. No negative ground-state bleach from the initial bis-diyne is observed in the recorded spectral window as the measured static UV/VIS spectrum of **Me-BD-Me** in THF (Fig. S2†) indicates its electronic absorption to be mostly in the deep-UV (<300 nm) range. The transient absorption spectra of **Me-BD-Me** in THF, CHCl<sub>3</sub>, and CH<sub>3</sub>CN are qualitatively similar, and three spectral bands are observed in all three cases. At early pump-probe time delays (Fig. 2a, S1a and d†), the pronounced positive band at 350–450 nm (subsequently referred to as band 1) appears immediately upon excitation and loses amplitude on a several-picosecond time scale (see below for a global fit and quantitative data). Band 1 of **Me-BD-Me** in different solvents shows multi-peak character (Fig. S3†). Taking **Me-BD-Me** in THF as an example (Fig. 2a), two pronounced peaks (2.98 eV/416 nm and 3.44 eV/360 nm) with comparable intensities were observed at 0.5 ps delay for **Me-BD-Me**, while an additional shoulder (3.14 eV/395 nm) of the 2.98 eV/416 nm peak is also observed. Subsequently, the 2.98 eV/416 nm peak rapidly decays together with the shoulder, while the 3.44 eV/360 nm peak undergoes red-shifting as a new band grows at 3.34 eV/371 nm (Fig. 2b). Similar spectral character and behaviors can also be observed for **Me-BD-Me** in both CHCl<sub>3</sub> and CH<sub>3</sub>CN. Considering the fast decay behavior, we tentatively attribute the multi-peak band 1 to the excited-state absorption (ESA) band of the initially populated S<sub>1</sub> state of **Me-BD-Me**. The observed decay of band 1 within hundreds of picoseconds might correspond to the electronic deactivation (S<sub>1</sub> → S<sub>0</sub>) or the following photochemical transformation to bleach the S<sub>1</sub> state. During the peak red-shifting process in the range of 50 to 100 ps delay, the 416 nm peak of the ESA band also decays and disappears with a similar time scale, which confirms that the red-shifting is a result of the decay of ESA and the growing of a new band at 371 nm. We performed TD-DFT calculations<sup>84–87</sup> on the structurally optimized S<sub>1</sub> state of **Me-BD-Me** in THF solution with the CAM-B3LYP<sup>88</sup> functional and def2-TZVP<sup>89,90</sup> basis set, which predicted two vertical transitions with pronounced oscillator strength, *i.e.* S<sub>1</sub> → S<sub>2</sub> (1.31 eV/946 nm) and S<sub>1</sub> → S<sub>6</sub> (2.73 eV/454 nm). The latter is reasonably consistent with the experimentally observed peak at 2.98 eV/416 nm, while the S<sub>1</sub> → S<sub>2</sub> transition is beyond our probe wavelength range. Note that the calculated (454 nm) and experimentally observed wavelength (416 nm) show a discrepancy of 2.98 eV (416 nm) – 2.73 eV (454 nm) = 0.25 eV, which is still in the expected confidence range of DFT calculations of electronic transition energies that are conventionally associated with errors of ±0.3 eV. A clear rise in the transient spectra can be observed at the red edge (Fig. 2a, b, S1a, b, d and e†), which shows a decay process similar to band 1, and implies the existence of an extra positive band in the near-infrared regime. Such an observation is consistent with the calculated S<sub>1</sub> → S<sub>2</sub> ESA band. The peaks at 3.14 eV/395 nm and 3.44 eV/360 nm observed in band 1 may originate from vibrational progressions or excitations to higher states.

The disappearance of band 1 indicates excited-state deactivation that subsequently leads to the rise of a strong and narrow band (band 2, peak at 3.35 eV/370 nm) in the 350–400 nm region

(Fig. 2b, S1b and e†) that builds up within hundreds of picoseconds. Band 2 is slightly red-shifted from the 3.44 eV/360 nm peak of band 1; apparently bands 1 and 2 are strongly overlapped with each other in the 350–450 nm regime. It is difficult to conclude whether band 2 is newly formed but overlapped with band 1, or band 2 is subsequently transformed from the 350–400 nm part of band 1, which we attempt to clarify by the quantitative target analysis in the following sections. We postulate that this corresponds to the photochemical generation of a reactive intermediate formed from the excited bis-diyne, namely the *o*-benzynes (**Me-BZ-Me**), discussed further and confirmed below. The TD-DFT calculations also predicted the vertical excitation energy of *o*-benzynes **Me-BZ-Me** to be 3.33 eV/372 nm (S<sub>0</sub> → S<sub>1</sub>), which is consistent with the experimentally observed energy of band 2, *i.e.* 3.35 eV/370 nm.

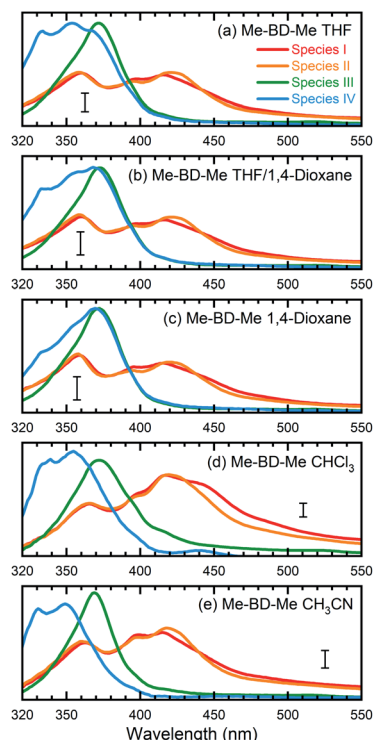
In the following step (Fig. 2c, S1c and f†), band 2 slowly decays with the concurrent rise of a new band (band 3) at even shorter wavelengths (320–370 nm), visible as a shoulder of band 2. Several peaks can be observed within band 3, which may originate from vibrational progressions. The sequential nature of the photoreaction sequence (band 1 → band 2 → band 3) is nicely observed by the occurrence of isosbestic points at 393 nm (Fig. 2a, b, S1a and b†), 368 nm (Fig. 2c) for **Me-BD-Me** in THF and CHCl<sub>3</sub>, and at 388 nm (Fig. S1d and e†) and 356 nm (Fig. S1ff†) for **Me-BD-Me** in CH<sub>3</sub>CN.

We performed a target analysis of the transient UV/VIS absorption spectra to confirm the sequential process of reactive steps, obtain time constants, and assign the associated species. Singular-value decomposition indicated the presence of four linearly independent components (species I, II, III, IV) above the experimental noise level, and three time constants [ $\tau_1$ (I → II),  $\tau_2$ (II → III),  $\tau_3$ (III → IV)] describing their temporal evolution. The target analysis, therefore, employed the assumption of a sequential kinetic model consisting of three steps (I → II → III → IV) with any possible decay of species IV occurring beyond the maximum time delay and, thus, not observable. The set of species-associated spectra (SAS) obtained and corresponding fitted time constants of this analysis, as well as others further discussed below, are shown in Fig. 3a and summarized in Table 1. The quality of the fit is shown in Fig. S4† with time traces at selected probe wavelengths, while the target analysis fitted concentration evolution of each transient species of **Me-BD-Me** and perylene in CHCl<sub>3</sub> are displayed in Fig. S5.†

As shown in Fig. 3a, d and e (red and orange curves), the target analysis successfully isolated the bi-exponential decay (I → II → ) of band 1 observed in the transient absorption spectra (Fig. 2). Such a bi-exponential decay of an ESA band can be explained by a typical non-radiative decay of the excited state *via* a pronounced potential energy barrier,<sup>91–93</sup> *i.e.*, the initially populated excited state (species I) relaxes [ $\tau_1$ (I → II)] within the S<sub>1</sub> potential energy surface until it reaches a local minimum (species II). Then, the trapped wave packet must overcome an energy barrier [ $\tau_2$ (II → III)] to reach a conical intersection (CI) associated with further photochemical conversion.

Regarding process I → II, we firstly exclude two potential assignments that might in principle show up on a several





**Fig. 3** Species-associated spectra (SAS) obtained from target analysis of transient UV/VIS spectra. Data are shown for **Me-BD-Me** in (a) THF, (b) a 50/50 mixture of THF and 1,4-dioxane, (c) 1,4-dioxane, (d)  $\text{CHCl}_3$ , and (e)  $\text{CH}_3\text{CN}$ . The sequential model ( $\text{I} \rightarrow \text{II} \rightarrow \text{III} \rightarrow \text{IV}$ ) was applied for the target analysis to estimate the SAS and corresponding time constants which are listed in Table 1. See text for detailed assignments. The scale bar in each subplot corresponds to an absorbance change of 5 mOD.

picoseconds time scale in the early stages of excited-state evolution following optical excitation: (1) vibrational relaxation in the excited state, *i.e.*, the wave packet escaping from the initially populated Franck-Condon area, can be excluded as we observed neither red-shift nor spectral narrowing during the  $\text{I} \rightarrow \text{II}$  process; (2) we also exclude ultrafast solvation, because

solvation is usually highly dependent on solvent polarity whereas, in our case, the fitted time constants of  $\text{I} \rightarrow \text{II}$  show no significant change upon varying the solvent, from high-polarity acetonitrile ( $\epsilon = 35.95$ ) to low-polarity 1,4-dioxane ( $\epsilon = 2.21$ ). Next, the DFT/TD-DFT calculations indicate that both the ground ( $\text{S}_0$ ) and excited state ( $\text{S}_1$ ) of **Me-BD-Me** have two conformers with comparable energies: the “open” conformation in which the two  $-\text{C}\equiv\text{C}-\text{C}\equiv\text{C}-$  units are relatively far from each other and the “closed” conformation in which the two  $-\text{C}\equiv\text{C}-\text{C}\equiv\text{C}-$  units are close to each other (see Fig. S6†). The “open” conformation was shown to be the structure of ground-state **Me-BD-Me** in the solid state by single-crystal X-ray diffraction,<sup>94</sup> and our DFT calculations on **Me-BD-Me** also indicate a preference for the open conformation with  $\Delta G(\text{Open}) - \Delta G(\text{Closed}) = -3.3 \text{ kJ mol}^{-1}$  in favor of the open conformer. However, it is logical to assume that the “closed” conformation leads to an increased HDDA reaction probability between the two  $-\text{C}\equiv\text{C}-\text{C}\equiv\text{C}-$  units in the excited state because the two units have to approach each other for the reaction to occur. Thus, we tentatively attribute the process  $\text{I} \rightarrow \text{II}$  to the excited-state isomerization of **Me-BD-Me** from “open” to “closed” conformation or other structural alteration for assisting the subsequent HDDA reaction, although this particular assignment cannot be proved unambiguously with the available data. As the electronic absorption spectrum is normally insensitive to such structural conformational changes, the SAS of species I and II are similar to one another and differ only by their associated time scales. These time constants of several picoseconds are consistent with the typical time scales of intramolecular excited-state isomerization.<sup>95,96</sup> We note that the TD-DFT calculations are not accurate enough (at a  $\sim 0.3 \text{ eV}$  confidence level) to predict the small differences between the transient absorption spectra of species I and II. The isomerization process includes no drastic changes in geometry and the electronic states should be able to follow the geometry adiabatically. Larger differences in the spectrum might be expected if the geometry change during the isomerization gave rise to new electronic couplings. This does not seem to be the case,

**Table 1** Time constants (with  $2\sigma$  error limits) obtained by target analysis with a sequential kinetic model of **Me-BD-Me** in various solvents and with co-reactants

	BD <sup>a</sup> in THF <sup>b</sup>	BD <sup>a</sup> in THF <sup>b</sup> /D <sup>c</sup>	BD <sup>a</sup> in D <sup>c</sup>	BD in CH <sub>3</sub> CN
$\tau_1(\text{I} \rightarrow \text{II})/\text{ps}$	$3.17 \pm 0.28$	$2.77 \pm 0.31$	$2.62 \pm 0.24$	$3.03 \pm 0.32$
$\tau_2(\text{II} \rightarrow \text{III})/\text{ps}$	$48.5 \pm 5.0$	$50.5 \pm 4.7$	$51.3 \pm 4.5$	$63.2 \pm 5.9$
$\tau_3(\text{III} \rightarrow \text{IV})/\text{ns}$	$4.03 \pm 0.78$	$4.15 \pm 0.76$	$4.64 \pm 0.82$	$2.47 \pm 0.30$
	BD <sup>a</sup> in CHCl <sub>3</sub>	BD <sup>a</sup> + P <sup>d</sup> in CHCl <sub>3</sub>	P <sup>d</sup> in CHCl <sub>3</sub>	
$\tau_1(\text{I} \rightarrow \text{II})/\text{ps}$	$5.27 \pm 0.45$	$6.58 \pm 0.59$	$1.92 \pm 0.15$	
$\tau_2(\text{II} \rightarrow \text{III})/\text{ps}$	$70.2 \pm 5.8$	$74.1 \pm 7.8$	$24.8 \pm 2.6$	
$\tau_3(\text{III} \rightarrow \text{IV})/\text{ns}$	$7.47 \pm 1.21$	$6.45 \pm 1.10$	$5.46 \pm 1.09^e$	

<sup>a</sup> BD: bis-diyne, *i.e.*, **Me-BD-Me**. <sup>b</sup> THF: tetrahydrofuran. <sup>c</sup> D: 1,4-dioxane. <sup>d</sup> P: perylene. <sup>e</sup> The transient absorption spectra of perylene were fitted with a three-step model ( $\text{I} \rightarrow \text{II} \rightarrow \text{III} \rightarrow \text{IV}$ ), wherein the third time constant indicates the decay of species III, *i.e.*  $\tau_3(\text{III} \rightarrow \text{IV})$ , instead of  $\tau_3(\text{III} \rightarrow \text{IV})$ . See Fig. S5 for details.





however, according to the experimental evidence and the calculations.

The process II  $\rightarrow$  III produces pronounced absorption bands centered at 370 nm with formation time constants of 48.5 ps (in THF), 70.2 ps (in  $\text{CHCl}_3$ ) and 63.2 ps (in  $\text{CH}_3\text{CN}$ ), respectively, visible in the green curves in Fig. 3a, d and e. Species III then slowly converts [ $\tau_3(\text{III} \rightarrow \text{IV}) = 3\text{--}5\text{ ns}$ ] to species IV, visible as a new band with a vibrational progression at shorter wavelengths (left peak(s) in blue curves of Fig. 3). While we assign species I and II to be the initially populated  $\text{S}_1$  state in the “open” form and its relaxed “closed” conformer, respectively, the assignment of species III and IV requires more consideration. As we mentioned above, the vertical excitation energy of *o*-benzyne (**Me-BZ-Me**) is 3.33 eV ( $\text{S}_0 \rightarrow \text{S}_1$ ), which is consistent with the experimentally observed energy of band 2 (3.35 eV). However, such consistency alone is not sufficient to assign band 2 to the benzyne species. Considering the fact that an experimental electronic spectrum of benzyne species in solution has not been reported previously, direct comparison with other experimental data is difficult. Therefore, we turned to an indirect procedure by performing a series of intra and intermolecular trapping experiments on benzyne that, together with the consistent DFT description, makes the assignment possible, which is discussed in the following sections.

If we assume the observed sequence (I  $\rightarrow$  II  $\rightarrow$  III  $\rightarrow$  IV) to be the  $h\nu$ -HDDA reaction, species III, with a strong absorption band (band 2) at  $\sim 370\text{ nm}$ , might belong to the *o*-benzyne derivative (denoted as **Me-BZ-Me**) generated. Then, species IV (band 3) can be assigned to secondary products of reactions of the *o*-benzyne with abundant possibilities, highly depending on the chemical environments. In order to identify the photo-products, and assign the transient absorption features, we carried out different time-resolved intermolecular trapping experiments as described in the following paragraphs. The strategies were designed to confirm the existence of the transient *o*-benzyne spectrum in our transient absorption observations. Our objective is the assignment of the slowly rising band 3 observed in transient absorption spectra (see Fig. 2c, S1c and f†) of **Me-BD-Me** in THF,  $\text{CHCl}_3$ , and  $\text{CH}_3\text{CN}$ , *i.e.*, species IV from the target analysis (see Fig. 3).

### Double-hydrogen transfer

As a first possibility, we consider the scenario that double-hydrogen transfer from an alkane solvent generates the benzene species **Me-H<sub>2</sub>BZ-Me** (Fig. 1, path 1). This is plausible because THF was reported to be a relatively good 2H donor (with a yield of  $\sim 60\%$ ) for reaction with *o*-benzyne.<sup>17</sup> Thus, we carried out transient absorption measurements on **Me-BD-Me** in 1,4-dioxane and a THF/1,4-dioxane mixture because the 2H transfer yield is almost zero in 1,4-dioxane.<sup>17</sup>

The results of transient absorption measurements on **Me-BD-Me** are shown for a 3.8 ns delay in Fig. 4a for THF (red), a 50/50 mixture of THF and 1,4-dioxane (pink), and pure 1,4-dioxane solvent (light pink), and the results of the corresponding target analysis are displayed in Fig. 3b for the 50/50 mixture, and in Fig. 3c for pure 1,4-dioxane. It is clear that band 3 builds

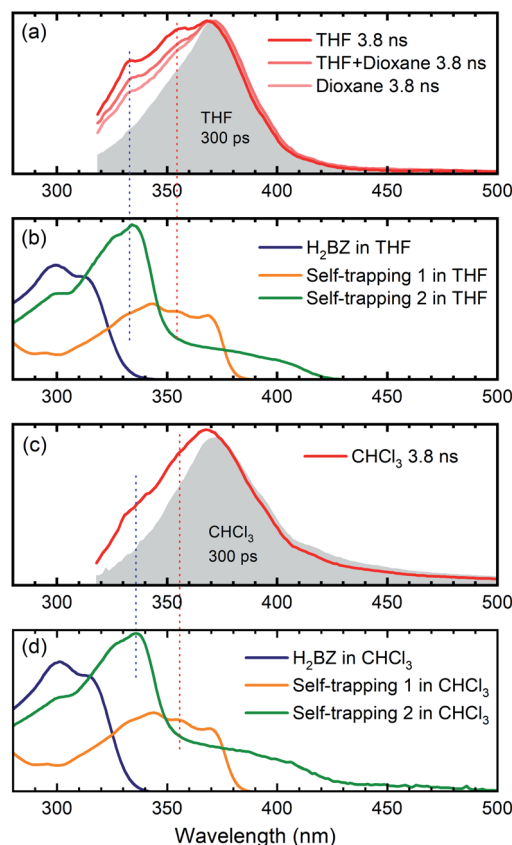


Fig. 4 Comparison between transient absorption spectra of **Me-BD-Me** and spectra of chemically isolated trapping products. (a and c) Transient absorption spectra in the 320–500 nm region at 300 ps (gray filled) and 3800 ps (colored solid lines) delay of **Me-BD-Me** in THF, THF/1,4-dioxane, 1,4-dioxane (a) and  $\text{CHCl}_3$  (c). (b and d) Static absorption spectra of chemically isolated **Me-H<sub>2</sub>BZ-Me** (navy line) and of two self-trapping species (orange and green lines) in THF (b) and  $\text{CHCl}_3$  (d). Several dashed lines are used for assisting the comparison. See Fig. 1 for structural formulae, and text for details.

substantially even in 1,4-dioxane, although it is weaker than in THF. Thus, it can be tentatively excluded that band 3 is dominated by the **Me-H<sub>2</sub>BZ-Me** species. In order to provide more direct evidence, chemical synthesis and purification of **Me-H<sub>2</sub>BZ-Me** was conducted together with comprehensive structural characterization (ESI Note 2†). The measured absorption spectrum of **Me-H<sub>2</sub>BZ-Me** (Fig. 4b, navy line) is located mostly at wavelengths shorter than 320 nm, *i.e.*, outside the probe window of our transient absorption measurements. Therefore, although the double-hydrogen transfer is well-known as an important secondary reaction following benzyne formation in solution, any **Me-H<sub>2</sub>BZ-Me** contribution to band 3 must be very small even if the species is formed. The **Me-H<sub>2</sub>BZ-Me** species can be safely excluded as a potential assignment of the slowly formed band 3 following band 2 at 370 nm.

### Intermolecular self-trapping by bis-diyne

With **Me-H<sub>2</sub>BZ-Me** excluded to be responsible for transient absorption band 3, we performed chemical separation of a thermal reaction of **Me-BD-Me**. Two dominating and stable



products were separated using flash chromatography. The structural characterization confirmed a product with a dimeric structure, *i.e.*, a naphthalene derivative,<sup>21,94</sup> denoted Self-trapping-1 with its structure illustrated in Fig. 1 (path 2, green). In addition, an indane product (Self-trapping-2, Fig. 1, path 3, orange) and a fragment (Fig. 1, path 3 gray) were also isolated and identified.<sup>94</sup> Although the isolated products arose from a thermal reaction, it has been reported that the HDDA products arising from both the thermally and photochemically generated *o*-benzynes intermediate are the same.<sup>83</sup> Thus, it was assumed that the electronic state of the *o*-benzynes intermediate is the same in both cases. Fig. 4b and d show the absorption spectra in THF and CHCl<sub>3</sub>, respectively, of the two products Self-trapping-1 (orange) and Self-trapping-2 (green). It can be seen that these spectra match the transient spectral features of band 3 for both THF and CHCl<sub>3</sub> (vertical dashed lines in Fig. 4). This correspondence indicates that HDDA-generated *o*-benzynes is probably self-trapped by an extra aryl butadiyne moiety of **Me-BD-Me** (for a plausible reaction mechanism see ref. 94). With the most plausible assignment of band 3 thus in place, band 2 at ~370 nm can be concluded to be the *o*-benzynes species.

In addition to the direct interpretation of the transient spectra in Fig. 4, we compare the steady-state absorption spectra of the isolated products with the SAS from target analysis in Fig. S10.† It can be seen that the SAS of species IV in THF and CHCl<sub>3</sub> are consistent with spectra of the isolated products, *i.e.*, Self-trapping-1 (at ~335 nm) and Self-trapping-2 (at ~354 nm and ~368 nm). The trapping experiment thus provides evidence that the observed band 3 can be tentatively attributed to the self-trapping product of *o*-benzynes with **Me-BD-Me** itself, and that, in turn, band 2 probably belongs to the prior *o*-benzynes intermediate from which the self-trapping species emerges.

### Intermolecular trapping by perylene

An alternative strategy for trapping *o*-benzynes is its reaction with species containing  $\text{C}=\text{C}$  bonds such as in a  $\pi$ -conjugated system reported recently (without time resolution).<sup>12,15,26,47,83</sup> We also made use of this reaction, but again employing femtosecond time-resolved techniques. The trapping product is expected to absorb in the >400 nm wavelength region where the transient absorption of the reactant bis-diynes is sufficiently weak to allow the largely unhindered observation of the trapping products. Among the reported options, perylene, with its absorption in the 400–450 nm region,<sup>47</sup> is ideal for our application. In such cases, we have to consider that the intermolecular trapping process occurs as a bimolecular reaction with diffusion-controlled kinetics.<sup>97–100</sup> To promote the diffusion and enable us to observe the trapping product within our maximum delay range, we employed low-viscosity CHCl<sub>3</sub> as a solvent for the perylene trapping experiments. Meanwhile, according to our target analysis, **Me-BD-Me** shows the slowest self-trapping reaction in CHCl<sub>3</sub> compared with reactions in THF and CH<sub>3</sub>CN (see Table 1). Thus, utilizing CHCl<sub>3</sub> minimizes the interference of the self-trapping reaction in the delay range employed. Although perylene can also be excited directly at our pump wavelength (295 nm), the remaining ground-state concentration should be sufficient for

trapping *o*-benzynes, as the molar extinction coefficient of perylene at 295 nm is small (Fig. S11†).

We carried out transient absorption measurements on **Me-BD-Me** and on **Me-BD-Me**/perylene (1 : 5 concentration ratio), referred to as “trapping mixture”, both in CHCl<sub>3</sub> solvent under otherwise similar conditions as described in the sections above. Again, we performed a target analysis using a sequential model on the temporal-spectral maps measured, and the resulting SAS are displayed in Fig. 5a (trapping mixture) and Fig. 5b (pure **Me-BD-Me**). For comparison, we also measured the transient absorption spectra of pure perylene in CHCl<sub>3</sub> under identical concentration and excitation conditions, such as excitation wavelength and power, which produced the transient absorption spectra shown in Fig. S12a and b,† and the target analysis results shown in Fig. 5, S12c and d.†

The transient spectra of **Me-BD-Me** in CHCl<sub>3</sub> display features similar to the ones in THF or 1,4-dioxane, while UV-excited perylene contributes multi-peak structures originating from ground-state bleach and stimulated emission in the 380–520 nm wavelength range (Fig. S12†) which persist until the maximum delay

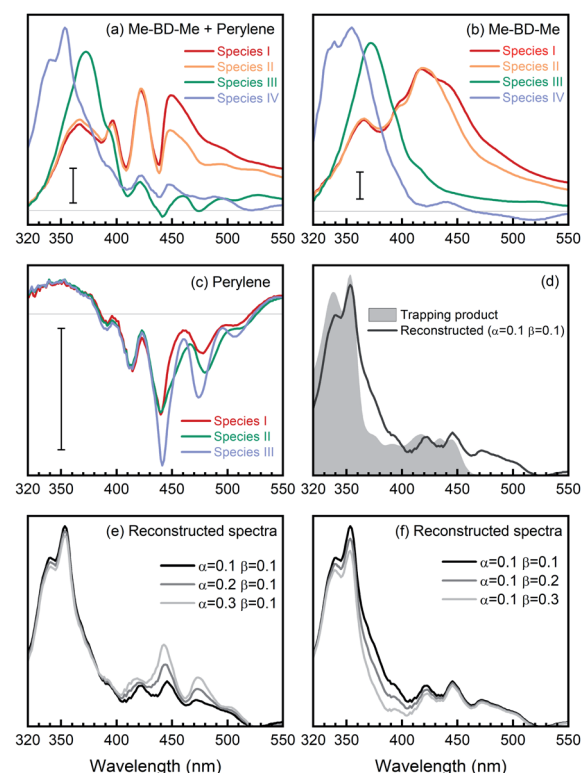


Fig. 5 Reconstruction of *o*-benzynes trapping with perylene in CHCl<sub>3</sub> solvent using SAS. (a) SAS from target analysis of transient absorption of **Me-BD-Me**/perylene mixture [ $S_M(N, \lambda)$ ], wherein  $N$  stands for the numbering of species from the target analysis. (b) SAS of pure **Me-BD-Me** [ $S_B(N, \lambda)$ ]. (c) SAS of pure perylene [ $S_P(N, \lambda)$ ]. (d) Reconstructed spectrum of trapping product [ $S_T(\lambda)$ , pink line] calculated by  $S_T(\lambda) = S_M(4, \lambda) - \alpha S_P(3, \lambda) - \beta S_B(3, \lambda)$ , and steady-state absorption spectrum of chemically isolated trapping product (gray filled). (e and f) Reconstructed spectra of trapping products using different weighting parameters  $\alpha$  (e) and  $\beta$  (f). The scale bars correspond to an absorbance change of 5 mOD.



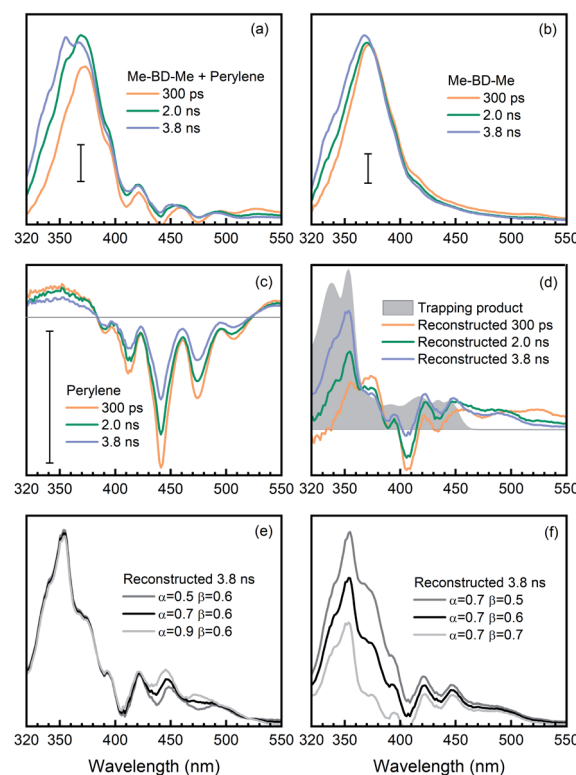
**Table 2** Possible species and their representative SAS and transient absorption spectra in the trapping mixture of Me-BD-Me and pure perylene in CHCl<sub>3</sub> upon UV photolysis

Species	Representative SAS	Transient spectra
Mixture	Mixture SAS4, $S_M(4, \lambda)$	Mixture, $A_M(\lambda, t)$
Perylene	Perylene SAS3, $S_P(3, \lambda)$	Perylene, $A_P(\lambda, t)$
<i>o</i> -Benzynes	<b>Me-BD-Me</b> SAS3, $S_B(3, \lambda)$	<b>Me-BD-Me</b> , $A_B(\lambda, t)$
Trapping product	Reconstructed, $S_T(\lambda)$	Reconstructed, $A_T(\lambda, t)$
Self-trapping products	Of minor importance	

time (3.8 ns). Such a structural signature also appears in the transient absorption spectra of the trapping mixture (Fig. S13†), *i.e.*, they are superimposed on the weak signals of the expected trapping products. Upon UV photolysis, both bis-diyne (**Me-BD-Me**) and perylene can be excited to their respective  $S_1$  states. Hence, as listed in Table 2, several components might contribute to the transient absorption of the trapping mixture after several nanoseconds of electronic deactivation and reaction: the excited state of perylene,  $h\nu$ -HDDA-generated *o*-benzynes, *o*-benzynes-perylene trapping product, and self-trapping products which are, however, expected to be of minor importance. In order to extract the desired trapping signal from the mixture of all possible contributions, we employed two spectral reconstruction strategies to remove the perylene and *o*-benzynes contributions as explained further below. As the most reliable reference, we isolated the *o*-benzynes-perylene trapping product (ESI Note 3†) and measured its absorption spectrum (see Fig. 5d). Although quantitative dynamics of bimolecular reactions in solution have been extensively reported for other species, the precondition for these studies is that only one of the reactants can be excited by the pump pulses.<sup>101–106</sup> Thus, we limit our analysis to a qualitative discussion due to the existence of both excited bis-dynes and excited perylene.

Target analysis with a sequential model ( $I \rightarrow II \rightarrow III \rightarrow IV$ ) successfully reproduced the transient absorption spectra of bis-dynes and their corresponding mixture with perylene in CHCl<sub>3</sub>, as can be inferred by the quality of the fit (Fig. S16†). For the first strategy, we removed the contribution of unreacted *o*-benzynes and perylene from the SAS of the trapping mixture and, thus, “manually” reconstructed the “spectral signature” of the trapping product. As summarized in Table 2, the SAS4 of the mixture [ $S_M(4, \lambda)$ ] arises from unreacted perylene [corresponds to SAS3 of perylene,  $S_P(3, \lambda)$ ], unreacted *o*-benzynes [SAS3 of **Me-BD-Me**,  $S_B(3, \lambda)$ ], trapping product [ $S_T(\lambda)$ ], and ignorable self-trapping products. With proper proportionality constants, the spectrum of the trapping product can thus be expressed as  $S_T(\lambda) \approx S_M(4, \lambda) - \alpha S_P(3, \lambda) - \beta S_B(3, \lambda)$ . Considering the highly complicated nature of the system, accurate estimation of factors  $\alpha$  and  $\beta$  is challenging. In the first step, the factor  $\alpha = 0.1$  was fitted as it minimizes the spectral modulation in the range of 380–520 nm. The factor  $\beta = 0.1$  was subsequently used to reach a convincing spectral shape without an unreasonable negative signal. The resulting reconstructed spectrum of the trapping product from transient absorption (Fig. 5d, black) is consistent with the spectrum of the chemically isolated product (gray). Nevertheless, an additional band can be observed in the longer

wavelength regime (460–510 nm), which cannot be attributed to the simple 1 : 1 trapping product, but might belong to a double (2 : 1) adduct structure with extended conjugation.<sup>47</sup> Meanwhile, although values of the factors  $\alpha$  and  $\beta$  are determined with some uncertainty, we found that alternations of  $\alpha$  (Fig. 5e) or  $\beta$  (Fig. 5f) do not substantially change the shape of reconstructed spectra. Therefore, we assign the reconstructed spectral feature to the expected trapping product.



**Fig. 6** Reconstruction of *o*-benzynes trapping with perylene in CHCl<sub>3</sub> solvent by using transient absorption spectra at selected delay times (300 ps, 2.0 ns and 3.8 ns). (a) Time-resolved transient absorption spectra of **Me-BD-Me**/perylene mixture [ $A_M(\lambda, t)$ ]. (b) Time-resolved transient absorption spectra of **Me-BD-Me** [ $A_B(\lambda, t)$ ]. (c) Time-resolved transient absorption spectra of perylene [ $A_P(\lambda, t)$ ]. (d) Reconstructed transient absorption spectra of the trapping product [ $A_T(\lambda, t)$ , colored lines] calculated by  $A_T(\lambda, t) = A_M(\lambda, t) - \alpha A_P(\lambda, t) - \beta A_B(\lambda, t)$ , and the absorption spectrum of the chemically isolated trapping product (gray filled). (e and f) Reconstructed transient spectra of trapping products at 3.8 ns delay time by using different weighted parameters  $\alpha$  (e) and  $\beta$  (f). The scale bars correspond to an absorbance change of 5 mOD.



In addition, as a second method to obtain the trapping product signature, we removed the contribution of pure perylene from the transient spectra of the trapping mixture and, thus, “manually” reconstructed the time-resolved spectra of the trapping product during its formation. This was done by subtracting the transient absorption spectra of pure perylene [ $A_P(\lambda, t)$ , Fig. 6c] and bis-diyne [ $A_B(\lambda, t)$ , Fig. 6b], with suitable proportionality constants, from the spectra of the trapping mixture [ $A_M(\lambda, t)$ , Fig. 6a] at selected delay times,  $A_T(\lambda, t) \approx A_M(\lambda, t) - \alpha A_P(\lambda, t) - \beta A_B(\lambda, t)$ , to obtain the spectral contribution of the trapping product [ $A_T(\lambda, t)$ ].

Again, the factor  $\alpha$  was optimized to  $\alpha = 0.6$ , so that the modulation of perylene in the reconstructed spectra can be minimized in the 380–520 nm regime. Meanwhile, as illustrated in Fig. 6d (gray filled) the trapping products have no absorption in the spectral region  $>550$  nm, *i.e.*,  $A_T(\lambda, t) \approx 0$ , and the factor for the *o*-benzyne contribution was optimized to  $\beta = 0.6$  so as to make the reconstructed spectra [ $A_M(\lambda, t) - \alpha A_P(\lambda, t) - \beta A_B(\lambda, t)$ ] as close as possible to the spectral baseline ( $\Delta A = 0$ ). The reconstructed spectra at selected delay times are shown in Fig. 6d (colored lines) in comparison with the isolated spectrum (gray). One can clearly see the rise of a positive band at 400–510 nm from 0.3 ns to 3.8 ns time delay, which is very similar in spectral range and shape to the reconstructed SAS (see Fig. 5d, black line) obtained from target analysis. Analogously to the discussion above, the precise values of the factors  $\alpha$  (Fig. 6e) or  $\beta$  (Fig. 6f) do not affect significantly the spectral feature of reconstructed spectra at 3.8 ns delay time. This provides further evidence that indeed the trapping product of HDDA-generated *o*-benzyne with perylene was observed in our experiments.

Considering both trapping experiments together, we have successfully observed and confirmed the existence of trapping products of *o*-benzyne intermediates. By backtracking the reaction sequence, we believe that the most plausible assignment for the strong transient absorption band centered at 372 nm is the  $h\nu$ -HDDA reaction-produced *o*-benzyne intermediate, which arises from excited-state bis-diyne (**Me-BD-Me**, 400–450 nm) with a time constant of  $\sim 50$  ps, and decays on

a nanosecond time scale *via* self-trapping by an additional bis-diyne or by perylene trapping.

Lastly, the thermal-HDDA reaction was reported to occur *via* a stepwise(-like) mechanism with a diradical intermediate,<sup>26</sup> which is a reaction from the electronic ground state in contrast to the photoreaction we describe in the present manuscript (see a comparative illustration of reaction pathways in Fig. 7). The  $h\nu$ -HDDA reaction relies on the UV excitation to populate the  $S_1$  state directly, from which the wave packet might decay *via* a  $S_1/S_0$  conical intersection leading to a highly efficient reaction.

## Conclusions

In this work, the UV-induced photochemistry of the bis-diyne compound **Me-BD-Me**, where Me is  $-\text{CH}_3$  and BD is  $-\text{C}_6\text{H}_4-(\text{C}\equiv\text{C})_2-(\text{CH}_2)_3-(\text{C}\equiv\text{C})_2-\text{C}_6\text{H}_4-$ , was measured by femto-second transient absorption spectroscopy in the UV/VIS region and in a 100 fs to 3.8 ns time range. We deduced multi-sequential kinetics for which, in the first step, the UV-populated first excited singlet state of **Me-BD-Me** isomerizes from an “open” to a “closed” conformation [ $\tau_1(\text{I} \rightarrow \text{II}) = 2\text{--}5$  ps]. The following processes are the formation [ $\tau_2(\text{II} \rightarrow \text{III}) = 50\text{--}70$  ps] of an intermediate and its slow decay [ $\tau_3(\text{III} \rightarrow \text{IV}) > 3$  ns]. By employing intermolecular trapping strategies *via* trapping reactions with a second **Me-BD-Me** molecule, as well as reaction with perylene, we assigned the process  $\text{II} \rightarrow \text{III}$ , observed experimentally, to be the formation of the *o*-benzyne intermediate in the  $h\nu$ -HDDA reaction. Use of the trapping strategies allowed us to circumvent the general insensitivity of UV/VIS transient absorption spectroscopy to chemical structure. The HDDA-generated *o*-benzyne species from **Me-BD-Me** displayed an absorption at  $\sim 372$  nm. To the best of our knowledge, this is the first report which spectroscopically confirms the existence of an *o*-benzyne species in solution as a product of the  $h\nu$ -HDDA reaction. The HDDA-generated *o*-benzyne has seen rapidly increased general interest and application in organic syntheses during the last eight years.<sup>15</sup> A wide variety of substrates have been investigated for reaction with HDDA-generated *o*-benzyne. They range from small molecules such as furan<sup>107</sup> to highly complex natural products.<sup>20</sup>

## Experimental section

### Sample preparation

**Me-BD-Me** was synthesized according to the literature.<sup>94</sup> The general information on chemical synthesis and isolation is summarized in ESI Note 1.† In all transient absorption spectroscopic experiments, samples were dissolved in spectroscopic-grade solvents (Sigma-Aldrich) as received. The static absorption spectra for sample preparation were measured with a JASCO V-670 UV/VIS spectrometer.

### Spectroscopic measurements

The details of the ultrafast transient absorption setup have been described elsewhere.<sup>108,109</sup> Briefly, a commercial 1 kHz Ti:sapphire laser system (Solstice, Spectra-Physics) delivered 120 fs

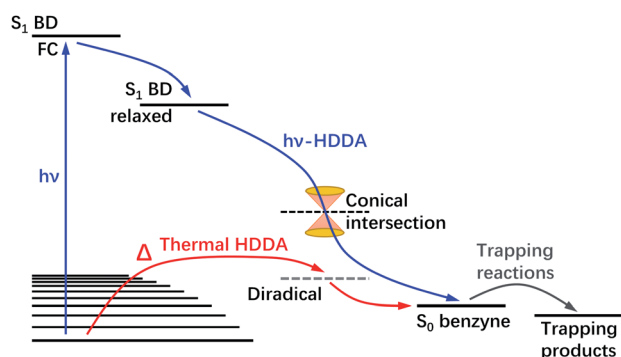


Fig. 7 Schematic illustration of the mechanism of the photochemical ( $h\nu$ ) and thermal HDDA reactions. The  $h\nu$ -HDDA mechanism (blue path) is revealed by the present work, while the thermal HDDA mechanism was reported by Wang *et al.*<sup>26</sup> and included here for comparison.





pulses centered at 800 nm. The  $\sim 100$  fs excitation pulses ( $\lambda_{\text{pump}} = 295$  nm) were derived from the second-harmonic output of a commercial nonlinear optical parametric amplifier (TOPAS White, Light Conversion). Approximately 0.5–1.0  $\mu\text{J}$  pulse energy was measured at the sample position. The broadband UV/VIS probe pulses were generated by focusing a small portion of the Ti:sapphire laser fundamental into a linearly moving  $\text{CaF}_2$  window, resulting in a white-light spectrum between 320 nm and 670 nm. Pump and probe beams were spatially overlapped in a flow cuvette with a sample thickness of 200  $\mu\text{m}$  which, at the chosen sample concentration, led to an optical density of 0.5 at 295 nm. The polarizations of the pump and probe beams were set to the magic angle of  $54.7^\circ$ . A cross correlation of  $\sim 120$  fs between pump and probe pulses was achieved. The pump–probe delay time was varied up to 3.8 ns by delaying the probe beam with a mechanical translation stage (M-IMS600, Newport). Every second pump pulse was blocked by a chopper driven at 500 Hz. After passing the sample, the probe pulses were dispersed in a spectrometer (Acton SP2500i, Princeton Instruments) and detected shot-to-shot by a CCD camera (Pixis 2K, Princeton Instruments). During the transient absorption measurements, the sample solution was circulated through a 0.1 mm thick flow-cell (48/UTWA2/Q/0.1, Starna GmbH) by using a micro-gear pump (mzr-4605, HNP Mikrosysteme). We conducted a control experiment to verify the reliability of our transient absorption measurements when dealing with irreversible photochemical reactions (Fig. S17†). We measured the transient absorption spectra at 350 ps (orange line) and then repeated the measurement (blue line) after constant illumination, for 1 h, with the 295 nm pump beam and, as can be seen in Fig. S17,† we observed no significant alteration of the transient absorption spectra, *i.e.*, any irreversible photodegradation of the sample does not create any uncertainty in the transient spectra. The transient data were evaluated *via* target analysis<sup>110</sup> with the software package Glo-taran based on the R-package TIMP.<sup>111</sup>

### Theoretical calculations

All TD-DFT calculations were executed with the Gaussian 09 program package,<sup>112</sup> using the CAM-B3LYP functional combined with the def2-TZVP basis set. Oscillator strengths among excited states were calculated using the wavefunction analyzer program Multiwfn.<sup>113</sup> Dispersion effects were incorporated by the use of Grimme's dispersion correction with Becke–Johnson damping<sup>114</sup> and solvation was treated with the polarizable continuum model.<sup>115</sup>

### Conflicts of interest

There are no conflicts to declare.

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