

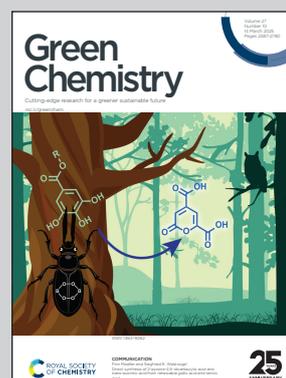
Showcasing research from Dr. Alexis Bordet's laboratory, Max Planck Institute for Chemical Energy Conversion, Mülheim an der Ruhr, Germany.

Photo-induced enhancement of hydrogenation activity for ruthenium nanoparticles immobilized on carbon dots

The present work demonstrates the capability of irradiated carbon dots to act as electron reservoirs for immobilized ruthenium nanoparticles, thereby greatly enhancing their hydrogenation activity. The strong synergistic effects arising from the combination of carbon dots and ruthenium nanoparticles under UV are potentially more broadly accessible with other carbon dots structures, metal nanoparticles compositions, and chemical transformations.

Image reproduced by permission of Alexis Bordet from *Green Chem.*, 2025, **27**, 2666.

As featured in:



See Alexis Bordet *et al.*, *Green Chem.*, 2025, **27**, 2666.



Cite this: *Green Chem.*, 2025, **27**, 2666

Received 30th October 2024,
 Accepted 30th December 2024

DOI: 10.1039/d4gc05468g

rsc.li/greenchem

Photo-induced enhancement of hydrogenation activity for ruthenium nanoparticles immobilized on carbon dots†

Carlotta Campalani,^a Manisha Durai,^{a,b} Walter Leitner^{a,b} and Alexis Bordet^{a*}

Ruthenium nanoparticles immobilized on environmentally benign luminescent carbon dots act as hydrogenation catalysts, whereby light irradiation results in greatly enhanced activity. Nitrogen-doped carbon dots were prepared by hydrothermal treatment, and served as supports for the organometallic synthesis of 2.1 nm Ru NPs. UV-irradiated Ru@CDs proved 1.5–13 times more active for C=C hydrogenation of α - β unsaturated ketones under mild

conditions (3.5 bar H₂, 60 °C, in water/butan-1-ol) than when used in the dark. Reference experiments and mechanistic investigations rationalize this activity enhancement by a transfer of electron density from the irradiated CDs to the Ru NPs. The strong synergistic effects arising from the combination of CDs and Ru NPs under UV are potentially more broadly accessible with other CDs structures and metal NPs compositions.

Green foundation

1. We investigate the synthesis, characterization, and application in catalysis of photo-responsive multifunctional catalytic systems composed of metal nanoparticles immobilized on carbon dots.
2. We demonstrate a photo-induced enhancement of hydrogenation activity (up to $\times 13$) for ruthenium nanoparticles immobilized on biodegradable, non-toxic, and luminescent carbon dots. Mechanistic investigations involving reference experiments and CO-IR measurements revealed a transfer of electron density from CDs to supported Ru nanoparticles resulting in a boost of the metal nanoparticles' hydrogenation activity
3. The recyclability of NPs@CDs catalysts should be improved, and the demonstrated capability of CDs to act as electron reservoirs for Ru nanoparticles should be explored for other CDs and nanoparticles compositions, in particular non-noble metals.

Catalytic hydrogenation with molecular hydrogen (H₂) is a crucial atom economic transformation for the chemical industry that is applied in the production of most chemicals such as fuels, fine chemicals, agrochemicals, and pharmaceuticals.¹ The development of sustainable catalysts with enhanced and tailor-made reactivity is thus a long-lasting goal in the field.² In heterogeneous catalysis, the modulation of catalytic activity can be achieved through the alteration of the structural and electronic properties of catalytic surfaces.³ For example, traditional strategies involve the control of particle size, structure and dis-

persion,⁴ the preparation of doped/multimetallic particles,^{5,6} the use of promoters,⁷ and metal support interactions.^{8,9} Photo-irradiation is also attractive in this context, as it offers sustainable means for activating or boosting performances of appropriately designed catalytic systems under mild conditions.¹⁰ In particular, photocatalysis is considered as an inherently sustainable approach, as it leverages light as a renewable energy source to drive chemical transformations, thereby reducing reliance on energy-intensive chemicals and processes.¹¹ Promising advances have been made in the use of irradiation in photocatalysis for the valorization of biomass and plastics to renewable chemicals in so-called “photorefineries”.¹² TiO₂-based catalysts are typical and extensively studied examples of photocatalysts, for which ultraviolet (UV) irradiations generate surface reactive electron-hole pairs that facilitate redox reactions.¹³

In recent years, carbon-based nanomaterials emerged as potentially more versatile and environmentally benign alterna-

^aMax Planck Institute for Chemical Energy Conversion, Stiftstraße 34–36, 45470 Mülheim an der Ruhr, Germany. E-mail: alexis.bordet@cec.mpg.de

^bInstitut für Technische und Makromolekulare Chemie, RWTH Aachen University, Worringerweg 2, 52074 Aachen, Germany

† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d4gc05468g>



tives to TiO₂ and classical semi-conductors for photocatalytic applications.^{14,15} In particular, carbon dots (CDs) possessing excellent photo-responsive properties are attracting increasing interest among the scientific community for applications in catalysis, and also in optoelectronic, bioimaging, drug delivery, and energy storage.^{16–21} CDs are biodegradable, non-toxic and luminescent carbon nanoparticles (NPs) with a spherical shape comprising a carbogenic core and a surface rich in organic functionalities such as hydroxyls, carbonyls and carboxylic groups.^{22–24} They can be prepared from various abundant and renewable carbon feedstock such as fish scales, waste paper, algae, *etc.*²⁵ Their ability to absorb/emit light at wavelengths ranging from UV to visible (Vis) and infrared (IR) allows their use as both electron donor and acceptor photocatalysts.²⁶ Their photoredox properties can be easily tailored through doping with various heteroatoms (*e.g.* nitrogen, sulphur, boron).^{27,28} The combination of CDs with metal NPs (NPs@CDs) can generate complementary interaction resulting in materials with enhanced thermo/photo-catalytic properties.^{29–32} For example, CDs have been used as supports for rhodium NPs, improving their stability in thermocatalytic alkene hydrogenation.³³ CDs can also act as e[−] reservoirs,³⁴ and the potential of photo-induced e[−] density transfer from CDs to metal NPs has been hypothesized as a way to boost the catalytic performance of metal NPs.^{29,32} Experimental demonstrations and fundamental understanding of such effects are stills lacking, however.

Herein, we present a novel photo-responsive hydrogenation catalyst composed of ruthenium (Ru) NPs supported on nitrogen-doped CDs (Ru@CDs). Ru@CDs are prepared, characterized, and applied to the hydrogenation of various α - β unsaturated ketones (Fig. 1). The focus is placed on the impact of light irradiation on catalytic performance, and on the understanding of underlying mechanisms.

Nitrogen-doped CDs were, indeed, prepared following a previously reported simple bottom-up hydrothermal treatment, using citric acid and diethylenetriamine as widely available precursors and microwave as a green heating method (see ESI† for experimental procedures).^{35–37}

Scanning Transmission Electron Microscopy (STEM) showed fairly spherical CDs with a diameter of 86.7 ± 21.5 nm (Fig. S1†), consistent with literature data.^{38–40} CDs' surface properties were probed by Fourier Transform infrared spectroscopy (FT-IR) in ATR mode, revealing the presence of various organic functionalities including hydroxyls, carbonyls, carboxyls, but also aminic and amidic groups, consistent with the desired nitrogen doping (Fig. S2†). Successful N-doping of the CDs was confirmed by elemental analysis that gave the following weight composition: 48% C, 27% O, 18% N, and 7% H. CDs' optical characteristics were studied by Ultraviolet-Visible (UV-Vis) and Photoluminescence-Photoluminescence Emission (PL-PLE) spectroscopies, evidencing two main absorption peaks (Fig. S3†). The first at 250 nm was attributed to the π - π^* transition of the aromatic carbogenic core, while the second at 350 nm was assigned to the n - π^* transition of the functional groups on the surface. PLE spectra showed a

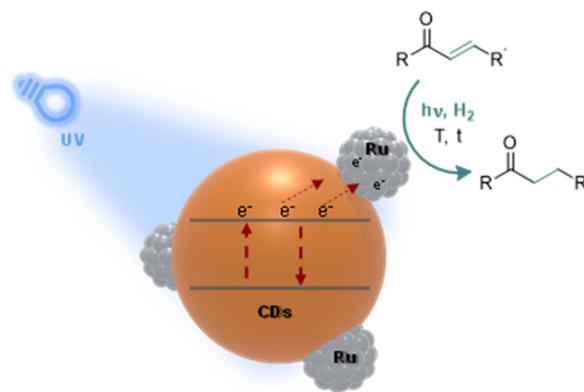


Fig. 1 Illustration of our approach: study of photo-responsive Ru@CDs applied to the model hydrogenation of α - β unsaturated ketones.

maximum emission in the blue region of the visible spectrum at 440 nm (Fig. S4†). The absence of blue or red shifts as a function of the excitation wavelength indicates that the emission originates from well-defined electronic states (*i.e.* functional groups or fluorophores) and not from surface defects.

The obtained N-doped CDs were used as photo-responsive supports for the preparation of Ru NPs by adapting an organometallic approach using [Ru(2-methylallyl)₂(cyclooctadiene)] as precursor (see ESI† for experimental details).^{41,42} The resulting Ru@CDs material contained 5 wt% of Ru as determined by Inductively-Coupled Plasma Mass Spectrometry (ICP-MS) analyses, and retained the chemical and optical properties of the CDs (Fig. S5† for FT-IR spectrum and Fig. S6† for UV-Vis spectrum). STEM analysis showed the presence of spherical Ru NPs with an average diameter of 2.1 ± 0.5 nm and narrow size distribution on the CDs (Fig. 2a and b). The blue emission at 440 nm observed for the pristine CDs was conserved also in Ru@CDs (Fig. 2c), confirming the retention of CDs' luminescent properties upon deposition of Ru NPs at their surface.

Potential effects of light irradiation on the catalytic properties of Ru@CDs were investigated using the hydrogenation of biomass-derived furfuralacetone (FFA, **1**) as a model reaction (Scheme 1) under mild conditions.

FFA was selected as the model substrate because it contains several reducible moieties that allow probing easily changes in catalytic activity and selectivity. Standard reaction conditions were set at 3.5 bar H₂, 60 °C, 4 h, 700 rpm, water/butan-1-ol as a green solvent mixture, without or with light irradiation at $\lambda_{\text{irradiation}} = 365$ nm (see Fig. S7–S10 in the ESI† for optimization steps).

Without irradiation, Ru@CDs was found poorly active under these conditions, giving only 28% conversion of **1** toward product **1a** (Table 1, entry 1). Light irradiation resulted in strikingly higher activity, with a conversion reaching 82% and the following distribution of products: 72% of **1a**, 7% of **1b**, 3% of **1c** (Table 1, entry 2). Time profiles revealed a five-fold increase in the initial C=C hydrogenation rate upon light exposure ($r_{0(\text{C}=\text{C})} = 0.06$ s^{−1} under irradiation vs. 0.012 s^{−1} in the dark Fig. S11 and S12†).



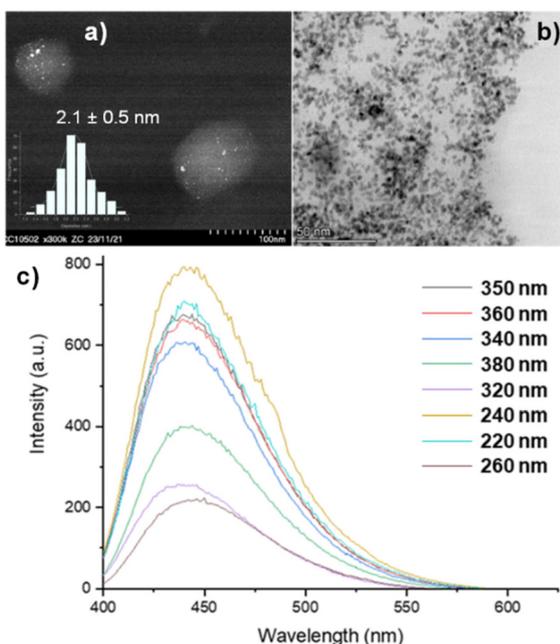
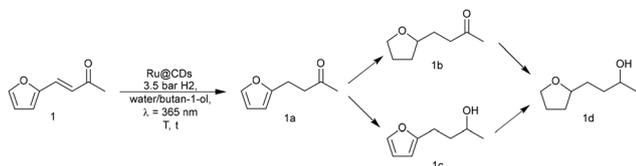


Fig. 2 Characterization of Ru@CDs by (a) STEM-HAADF, (b) STEM-BF, (c) PLE at different excitation wavelengths.



Scheme 1 Reaction sequence for the hydrogenation of furfuralacetone (**1**) using Ru@CDs in the presence of hydrogen and light irradiation.

Potential photothermal contributions to this substantial enhancement were investigated by performing blank experiments at a fixed temperature (30 °C) and monitoring any variation in the measured values of the reaction media (Table S1†). In all experiments, temperature variations

remained within measurement error, suggesting poor photothermal properties of CDs at the used excitation wavelength.^{43,44} These reference experiments indicate that the contribution of thermal effects to catalytic performance is negligible.

Reactions performed with irradiation and only CDs or without any catalyst gave negligible conversion (Table 1, entries 3 and 4), confirming the need for Ru NPs and the absence of spontaneous photo-induced hydrogenation of the substrate. Changing the irradiation wavelength to 450 nm led to lower conversion with Ru@CDs, and still no activity in the absence of catalyst (Table 1, entries 5 and 6). Interestingly, using white light from a simple torchlight gave 63% yield of **1a** (Table 1, entry 7), significantly higher than for a reaction in the dark. However, the enhancement is lower than for an irradiation at 365 nm due to the limited ability of our CDs to absorb the visible and infrared portions of the white light. Most importantly, no significant effect of light irradiation was observed when using a reference non-luminescent Ru/C catalyst (Table 1, entries 8 and 9), indicating that photothermal effects from the carbon support or plasmonic effects coming from the Ru NPs are negligible. Taken together, these results reveal a strong synergistic effect arising from the combination of Ru NPs and luminescent CDs. Satisfyingly, the reaction (conditions of Table 1, entry 2) could be scaled up by a factor of 40 (1.1 g (8 mmol) of substrate **1**, 800 mg of catalyst) while maintaining the same level of performance (84% conversion, 82% yield of **1a**, Scheme S1†). Recyclability was found challenging (Fig. S13†), although no substantial Ru NPs size increase (2.3 nm, Fig. S14†) nor Ru leaching (XRF) were detected. Thus, the observed deactivation is presumably due to the aggregation of the catalyst during washing steps, consistent with the known challenge associated with CDs separation and aggregation.^{45,46}

To gather additional insight, the systematic activity enhancement of Ru@CDs under light irradiation was investigated by FT-IR using carbon monoxide (CO) as a molecular probe (Fig. 3, see ESI† for experimental procedure). FT-IR spectra show CO absorption bands at 2289 cm⁻¹ for Ru@CDs in the dark (black line), and at 2216 cm⁻¹ for the irradiated

Table 1 Conversion and product yields for FFA hydrogenation using 3.5 bar of H₂, 20 mg catalyst (0.01 mmol Ru), 0.2 mmol FFA, water/butan-1-ol, 60 °C, 700 rpm

Entry	Catalyst	$h\nu$ (nm)	Time (h)	Conv. ^a (%)	Product yields ^a (%)			
					1a	1b	1c	1d
1	Ru@CDs	—	4	28	28	0	0	0
2	Ru@CDs	365	4	82	72	7	3	0
3	CDs	365	4	0	0	0	0	0
4	None	365	4	0	0	0	0	0
5	Ru@CDs	450	4	30	30	0	0	0
6	None	450	4	0	0	0	0	0
7	Ru@CDs	400–700	4	63	63	0	0	0
8	Ru/C	—	2	86	18	0	68	0
9	Ru/C	365	2	87	22	0	65	0

^a Determined by GC-FID using tetradecane as internal standard.



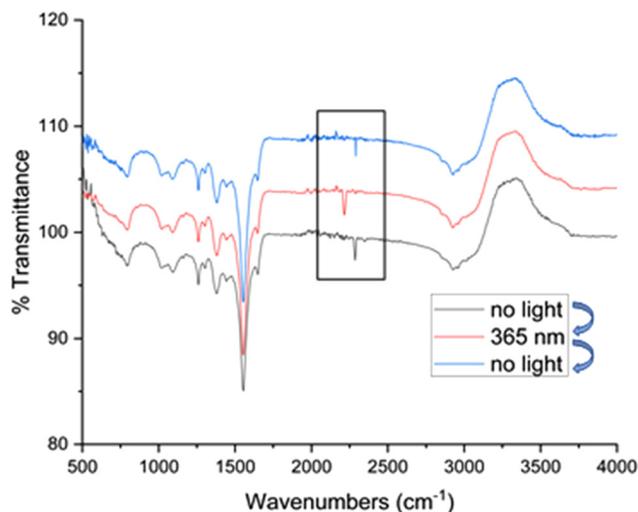


Fig. 3 Comparison between the FT-IR spectra of CO-adsorbed Ru@CDs in water under irradiation (365 nm, red spectrum) and in the dark (blue and black spectra). The black rectangle highlights the region with CO absorption bands.

sample (red line). Such a shift toward lower wavenumbers is characteristic of electron density being transferred from a metal center to the CO bond, weakening it and lowering the vibration frequency.^{47,48} This data evidences that irradiated CDs act as electron donors increasing the electron density on the surface of Ru NPs, which can then be transferred to CO or other molecules, thereby facilitating their reaction. Interestingly, the CO band shift was found fully reversible upon turning on or off light irradiation. Most importantly, such behavior was not observed when using reference Ru/C or metal-free CDs (Fig. S15 and S16[†]), demonstrating that intimate contact between CDs and metal NPs is required to observed synergistic effects in hydrogenation. Thus, the enhancement of hydrogenation activity observed for Ru@CDs under light irradiation is attributed to a transfer of electron density from CDs to the Ru NPs, resulting in facilitated activation and/or transfer of hydrogen to the C=C bond of FFA.

The optical properties of the catalyst, assessed *via* UV-Vis spectroscopy, were retained after reaction (Fig. S17[†]). On-off experiments were conducted to highlight the dependence of the reaction on the presence of UV irradiation and the adaptivity of the catalyst to intermittent electricity supply. As can be seen in Fig. 4, when the light source was switched off the reaction became slow and then started again with the restart of the irradiation.

The versatility of Ru@CDs as photo-enhanced hydrogenation catalysts was tested for other α - β unsaturated ketones (Table 2). Reactions were conducted under conditions adapted for each substrate, at incomplete conversion to facilitate the observation of irradiation effect (for more conditions, see Tables S2–S5[†]). With benzylideneacetone (3), conversion reached 70% under irradiation, and only 46% in the absence of light (Table 2, entries 3 and 4). The conversion levels of the

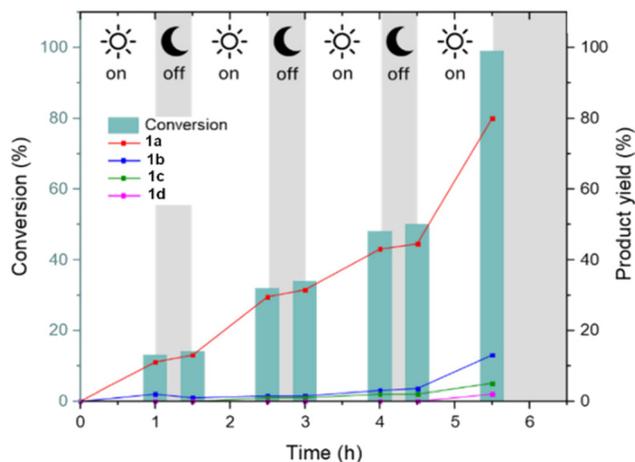


Fig. 4 Conversion and product yields versus time for on-off experiments for the hydrogenation of FFA with Ru@CDs. White areas represent the time frame conducted with irradiation at 365 nm, grey areas are in the dark.

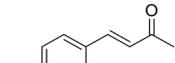
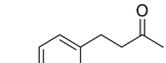
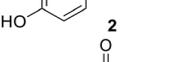
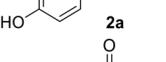
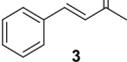
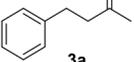
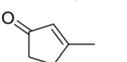
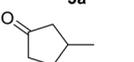
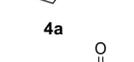
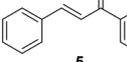
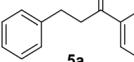
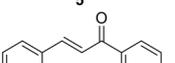
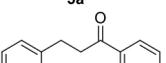
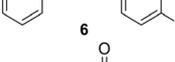
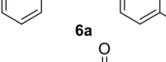
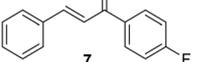
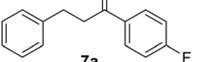
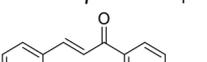
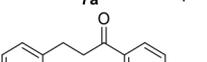
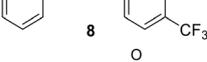
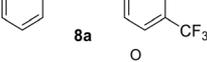
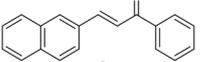
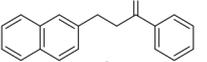
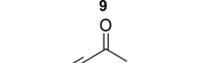
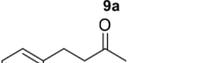
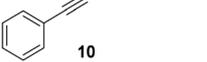
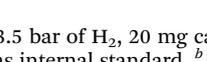
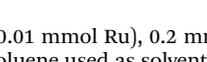
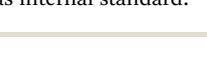
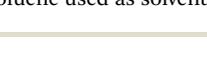
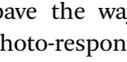
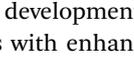
cyclic α - β unsaturated ketone methylcyclopentenone (4) remained moderate, but still substantially higher under irradiation (31%) than in the dark (9%) (Table 2, entries 5 and 6). Similar photo-induced enhancements were observed for chalcone (5) and its methoxy (6) and fluorinated (7) derivatives (Table 2, entries 7–12). Conversion reached 54–66% under light irradiation, and remained under 10% in the dark.

Trifluoromethyl- (8) and naphthyl-substituted (9) chalcones were less reactive under these reaction conditions (Table 2, entries 13–16), showing respectively 21% and 11% conversion. However, these substrates were not converted in the dark. Finally, the ability of Ru@CDs to reduce triple C–C bonds was assessed using phenylbutynone (10) as a substrate reaching 88% conversion under irradiation (with 50% yield of 10a, 22% of the corresponding alkene and 16% of the equivalent alcohol, Table 2, entry 17 and Table S6[†] entry 3) and 27% conversion without light (11% yield of 10a, Table 2, entry 18). No arene hydrogenation was observed for these substrates. Light-induced activity enhancement was observed irrespective of the substrates' absorption at 365 nm (Fig. S18–S27[†]), indicating the absence of correlation between catalytic performance and substrates' response to UV irradiations. No coupling product resulting from potential radical pathways was observed.

In conclusion, a multifunctional catalytic system composed of Ru NPs immobilized on luminescent and environmentally benign CDs has been prepared and characterized. The performances of the Ru@CDs catalyst in the hydrogenation of α - β unsaturated ketones under mild conditions were found substantially enhanced by UV irradiation. Mechanistic investigations involving references experiments and CO-IR measurements revealed a transfer of electron density from CDs to supported Ru NPs resulting in a boost of the metal NPs' hydrogenation activity. The present demonstration of the capability of irradiated CDs to act as electron reservoirs for Ru NPs is potentially applicable to other CDs compositions and metal NPs,



Table 2 Hydrogenation of various α - β unsaturated ketones using Ru@CDs with and without UV irradiation

Entry	Substrate	Product Xa	<i>t</i> (h)	<i>T</i> (°C)	λ	Conversion ^a (%)	Yield _{Xa} ^a (%)
1			4	60	Yes	100	100
2			4	60	No	13	13
3			4	80	Yes	70	67
4			4	80	No	46	43
5			6	80	Yes	31	27
6			6	80	No	9	7
7			6	80	Yes	63	63
8			6	80	No	8	8
9			6	80	Yes	66	66
10			6	80	No	5	5
11			6	80	Yes	54	54
12			6	80	No	6	6
13			6	80	Yes	21	21
14			6	80	No	2	2
15 ^b			6	80	Yes	11	11
16 ^b			6	80	No	0	0
17			4	80	Yes	88	50
18			4	80	No	27	11

Conditions: 3.5 bar of H₂, 20 mg catalyst (0.01 mmol Ru), 0.2 mmol substrate, water/butan-1-ol, 700 rpm, 365 nm. ^a Calculated *via* GC-FID using tetradecane as internal standard. ^b Water/toluene used as solvent.

and may pave the way toward the development of a wide variety of photo-responsive NPs@CDs with enhanced catalytic performance.

Author contributions

A. B. and C. C. conceptualized and developed the project. C. C. and M. D. performed experiments, data analysis. C. C., W. L. and A. B. wrote and edited the manuscript. All authors contributed and agreed on the final version of the manuscript.

Data availability

The data supporting this article have been included as part of the ESI.†

Conflicts of interest

The authors declare no conflicts of interest.

Acknowledgements

The authors acknowledge financial support by the Max Planck Society and by the Deutsche Forschungsgemeinschaft (DFG, German Research Foundation) under Germany's Excellence Strategy – Exzellenzcluster 2186 “The Fuel Science Center” ID: 390919832. We thank Alina Jakubowski, Annika Gurowski and Justus Werkmeister for GC, GC-MS and elemental analysis. The authors would like to thank Alin Benice Schöne and Norbert Pfänder for TEM and STEM/EDX measurements. Open Access funding provided by the Max Planck Society.

References

- H. U. Blaser, C. Malan, B. Pugin, F. Spindler, H. Steiner and M. Studer, *Adv. Synth. Catal.*, 2003, **345**, 103–151.
- K. Liu, R. Qin and N. Zheng, *J. Am. Chem. Soc.*, 2021, **143**, 4483–4499.
- A. T. Bell, *Science*, 2003, **299**(5613), 1688–1691.
- G. A. Somorjai and J. Y. Park, *Angew. Chem., Int. Ed.*, 2008, **47**, 9212–9228.



- 5 P. Buchwalter, J. Rose and P. Braunstein, *Chem. Rev.*, 2015, **115**, 28–126.
- 6 L. He, F. Weniger, H. Neumann and M. Beller, *Angew. Chem., Int. Ed.*, 2016, **55**, 12582–12594.
- 7 G. J. Hutchings, *Catal. Lett.*, 2001, **75**, 1–12.
- 8 P. Liu, R. Qin, G. Fu and N. Zheng, *J. Am. Chem. Soc.*, 2017, **139**, 2122–2131.
- 9 W. Karim, C. Spreafico, A. Kleibert, J. Gobrecht, J. VandeVondele, Y. Ekinici and J. A. van Bokhoven, *Nature*, 2017, **541**, 68–71.
- 10 M. Tarr and P. Zito, in *Photochemistry of Nanomaterials*, ed. American Chemical Society, ACS In Focus, 2022, vol. 15.
- 11 D. Ravelli, D. Dondi, M. Fagnoni and A. Albini, *Chem. Soc. Rev.*, 2009, **38**, 1999–2011.
- 12 S. Feng, P. T. T. Nguyen, X. Ma and N. Yan, *Angew. Chem., Int. Ed.*, 2024, **63**, e202408504.
- 13 X. Chen and S. S. Mao, *Chem. Rev.*, 2007, **107**, 2891–2959.
- 14 J. Ackermann, J. T. Metternich, S. Herbertz and S. Kruss, *Angew. Chem., Int. Ed.*, 2022, **61**, e202112372.
- 15 Y. Liang, Y. Li, H. Wang and H. Dai, *J. Am. Chem. Soc.*, 2013, **135**, 2013–2036.
- 16 S. N. Baker and G. A. Baker, *Angew. Chem., Int. Ed.*, 2010, **49**, 6726–6744.
- 17 G. A. M. Hutton, B. C. M. Martindale and E. Reisner, *Chem. Soc. Rev.*, 2017, **46**, 6111–6123.
- 18 S. Cailotto, E. Amadio, M. Facchin, M. Selva, E. Pontoglio, F. Rizzolio, P. Riello, G. Toffoli, A. Benedetti and A. Perosa, *ACS Med. Chem. Lett.*, 2018, **9**, 832–837.
- 19 H. Yu, R. Shi, Y. Zhao, G. I. Waterhouse, L. Z. Wu, C. H. Tung and T. Zhang, *Adv. Mater.*, 2016, **28**, 9454–9477.
- 20 C. Xia, S. Zhu, T. Feng, M. Yang and B. Yang, *Adv. Sci.*, 2019, **6**, 1901316.
- 21 J. Liu, R. Li and B. Yang, *ACS Cent. Sci.*, 2020, **6**, 2179–2195.
- 22 T. C. Wareing, P. Gentile and A. N. Phan, *ACS Nano*, 2021, **15**, 15471–15501.
- 23 V. Mishra, A. Patil, S. Thakur and P. Kesharwani, *Drug Discovery Today*, 2018, **23**, 1219–1232.
- 24 C. Campalani, E. Cattaruzza, S. Zorzi, A. Vomiero, S. You, L. Matthews, M. Capron, C. Mondelli, M. Selva and A. Perosa, *Nanomaterials*, 2021, **11**, 524.
- 25 Z. L. Wu, Z. X. Liu and Y. H. Yuan, *J. Mater. Chem. B*, 2017, **5**, 3794–3809.
- 26 (a) X. Wang, L. Cao, F. Lu, M. J. Mezziani, H. Li, G. Qi, B. Zhou, B. A. Harruf, F. Kermarrec and Y.-P. Sun, *Chem. Commun.*, 2009, 3774–3776; (b) C. Campalani, G. Petit, J. C. M. Monbaliu, M. Selva and A. Perosa, *ChemPhotoChem*, 2022, e202200234.
- 27 S. Cailotto, R. Mazzaro, F. Enrichi, A. Vomiero, M. Selva, E. Cattaruzza, D. Cristofori, E. Amadio and A. Perosa, *ACS Appl. Mater. Interfaces*, 2018, **10**, 40560–40567.
- 28 W. K. Li, J. T. Feng and Z. Q. Ma, *Carbon*, 2020, **161**, 685–693.
- 29 C. Lu, Q. Zu, X. Zhang, H. Ji, Y. Zhou, H. Wang, Q. Liu, J. Nie, W. Han and X. Li, *ACS Sustainable Chem. Eng.*, 2019, **7**, 8542–8553.
- 30 S. Sadjadi, M. M. Heravi, L. Mohammadi and M. Malmir, *ChemistrySelect*, 2019, **4**, 7300–7307.
- 31 J. Shen, W. Chen, G. Lv, Z. Yang, J. Yan, X. Liu and Z. Dai, *Int. J. Hydrogen Energy*, 2021, **46**, 796–805.
- 32 Y. Ren, C. Hao, Q. Chang, N. Li, J. Yang and S. Hu, *Green Chem.*, 2021, **23**, 2938–2943.
- 33 J. Zhang, Y. Chen, J. Tan, H. Sang, L. Zhang and D. Yue, *Appl. Surf. Sci.*, 2017, **396**, 1138–1145.
- 34 S. Deshmukh, A. Deore and S. Mondal, *ACS Appl. Nano Mater.*, 2021, **4**, 7587–7606.
- 35 K. Jiang, Y. Wang, X. Gao, C. Cai and H. Lin, *Angew. Chem., Int. Ed.*, 2018, **57**, 6216–6220.
- 36 T. V. de Medeiros, J. Manjoudakis, F. Noun, J.-R. Macairan, F. Victoria and R. Naccache, *J. Mater. Chem. C*, 2019, **7**, 7175–7195.
- 37 L. Vallan and H. Himahori, *ACS Appl. Electron. Mater.*, 2022, **4**, 4231–4257.
- 38 Y. Liu, S. Roy, S. Sarkar, J. Xu, Y. Zhao and J. Zhang, *Carbon Energy*, 2021, **23**, 795–826.
- 39 S.-C. Wei, Y.-W. Lin and H.-T. Chang, *J. Food Drug Anal.*, 2020, **28**, 558–574.
- 40 J. C. G. Esteves da Silva and H. M. R. Gonçalves, *TrAC, Trends Anal. Chem.*, 2011, **30**(8), 1327–1336.
- 41 A. Bordet and W. Leitner, *Acc. Chem. Res.*, 2021, **54**, 2144–2157.
- 42 A. Bordet, G. Moos, C. Welsh, P. Licence, K. L. Luska and W. Leitner, *ACS Catal.*, 2020, **10**, 13904–13912.
- 43 S. Balou, P. Shandilya and A. Priye, *Front. Chem.*, 2022, **10**, 1023602.
- 44 T. Zhang, J. Wu, Z. Tang and S. Qu, *Mater. Chem. Front.*, 2023, **7**, 2359–2372.
- 45 J. Gao, M. Zhu, H. Huang, Y. Liu and Z. Kang, *Inorg. Chem. Front.*, 2017, **4**, 1963.
- 46 L. L. Mokoloko, R. P. Forbes and N. J. Coville, *Catalysts*, 2023, **13**, 1201.
- 47 J. A. Anderson, C. H. Rochester and Z. Wang, *J. Mol. Catal. A: Chem.*, 1999, **139**, 285–303.
- 48 S. Xu, S. Chansai, S. Xu, C. E. Stere, Y. Jiao, S. Yang, C. Hardacre and X. Fan, *ACS Catal.*, 2020, **10**, 12828–12840.

