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Direct grafting-from of PEDOT from a photoreactive Zr-based MOF – a novel route to electrically conductive composite materials[†]

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The postsynthetic potential of the two-dimensional metal-organic framework Zr-*bzpdc*-MOF which is based on the photoreactive molecule benzophenone-4,4'-dicarboxylic acid (H_2bzpdc) is used here to selectively functionalize the MOF surface. We report the direct radical-induced oxidative grafting-from polymerization of the precursor EDOT on Zr-*bzpdc*-MOF, leading to an electrically conductive composite material and opening the road to a variety of applications.

Metal–organic frameworks (MOFs) feature an ordered structure with permanent porosity which is suitable for a variety of possible applications¹ like catalysis,^{2,3} gas separation and storage,^{4,5} sensing,^{6,7} in biomedicine^{8,9} or in energy-storage devices.¹⁰ However, the direct synthesis of materials with desired yet sensitive functionalities is limited due to the synthesis conditions of MOFs. Postsynthetic modification (PSM) is a strategy to alleviate such problems. PSM summarizes different approaches like linker exchange, postsynthetic metalation or organic reactions at the linker molecules.^{11–15}

The synthetic possibilities of organic reactions at linker molecules are in principle limitless.¹⁶ Usually, grafting procedures^{17–20} are carried out as grafting-to reactions. Especially for the modification of crystal surfaces with polymers this approach may be problematic as the first grafted-to polymer strands may hinder further access to the reactive functionalities at the surface, resulting in a low degree of functionalization. Grafting-from reactions, where polymer chains grow in a simultaneous and parallel fashion from surface-standing polymerization-initiating entities are usually more effective. Only few grafting-from MOF polymerization reactions have been described in the literature. These usually involve additional preparatory steps as fixing polymerization-initiating groups on the surface^{21,22} or adding a reactive MOF shell to an innocent MOF core.²² Here, we describe grafting-from polymerizations from the surface of MOF particles directly as they are obtained after synthesis and purification. We photochemically initiated a polymerization of 3,4-ethylenedioxythiophene (EDOT) to the electrically conductive polythiophene derivative poly-ethylenedioxythiophene (PEDOT), in this way directly synthesizing an electrically conductive MOF–polymer composite (illustrated in Scheme 1).

Electrical conductivity is currently a much sought-after property in metal–organic frameworks for applications like sensing, energy storage or electronics.^{23–29} Also applications in coatings of neuronal electrodes could be of interest, as they would allow electrical stimulation of nerve cells/registration of neuronal signals combined with drug delivery from the pores of the MOF. However, only very few MOFs exhibit intrinsic electrical conductivity. The rare examples rely on the formation of π -stacked pathways through threedimensional MOFs,^{30,31} delocalization in two-dimensional coordination polymers^{32–36} or *via* sulfur-based linker molecules³⁷ as well as the special properties of iron-based MOFs.^{38–40}

Alternatively, the introduction of guest molecules like tetracyanoquinododimethane,⁴¹ fullerene,⁴² polyaniline and polythiophene derivatives,^{40,43–49} as well as the formation of composites with conducting carbon modifications^{42,50–54} or conducting polymers^{55,56} can lead to conductive MOF-based materials. Generally, the introduction of guests into the pore space can compromise porosity; on the contrary, polymerization of EDOT in the pore system of MIL-101(Cr) leads to a conductive material with only slightly reduced porosity.⁵⁵

Especially for Zr-based MOFs there have been only a few studies to impart electrical conductivity on these compounds,^{42,46,57} although this family has been in the spotlight of research during the last years due to a controlled synthetic access using the modulation method,^{58,59} to their chemical and topological versatility and to their high chemical and thermal stability, as compared to other MOFs.^{60,61} Recently, we have described the Zr-*bzpdc*-MOF, which contains linkers derived from benzophenone-4,4'-dicarboxylic acid (H₂*bzpdc*).⁶² This MOF has a two-dimensional structure, a moderate porosity of about 650 m² g⁻¹, and can be delaminated into very thin sheets. The most striking feature of the Zr-*bzpdc*-MOF,

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Scheme 1 Schematic illustration of the direct grafting-from approach of PEDOT (orange chain) from photochemically reactive sites at the surface of a Zr-*bzpdc*-MOF crystal, starting from EDOT monomer units (orange dot).

however, is the possibility to use the photoreactive keto groups of the linker molecules for PSM reactions. Upon excitation with light, benzophenone moieties react with practically any compound containing a C–H bond,^{63–65} according to Scheme 2. We have exemplarily shown that such reactions can also proceed when the *bzpdc* linker is part of a MOF by grafting-to reactions of the Zr-*bzpdc*-MOF with decane and polyethylene glycol.⁶² Meanwhile, we have extended these grafting-to reactions to other organic molecules and have descried that molecules with more than four carbon atoms react pre-dominantly at the outer surface of the crystals whereas smaller molecules can enter the pore system so that the reaction can take place throughout the whole crystal.⁶⁶ Here, we use the ketyl radicals formed by photoexcitation of the keto groups of the Zr-*bzpdc*-MOF to start polymerization reactions.

The Zr-*bzpdc*-MOF was synthesized as described in the literature using a slightly modified procedure (Section S2.1, ESI⁺).⁶² This synthesis approach leads to rhombic shaped Zr-*bzpdc*-MOF crystals with only a very low amount of amorphous byproduct (Fig. S1, ESI⁺). The resulting colorless powder was Soxhlet-extracted with acetone for 24 h and dried under reduced pressure overnight to remove solvent molecules.

For the grafting procedure, Zr-*bzpdc*-MOF crystals were dispersed in neat EDOT in a quartz vessel, purged with argon for one hour and then irradiated under argon purging for a specific time with a UV LED at a wavelength of 365 nm to induce the biradical formation at the keto group of the benzophenone dicarboxylate linkers in the framework. To remove excess EDOT and non-attached oligo/ polymers, the products were Soxhlet-extracted with acetone for 24 h and dried under reduced pressure (further details on the synthesis and characterization methods are given in Sections S2.1, S3.1 and S3.2 of the ESI[†]).

The most obvious and visible result to confirm a successful polymerization of EDOT is the strong color change of the samples after irradiation. With increasing reaction time the color of the sample changes from colorless to dark brownish (Fig. 1a). The crystallinity of the MOFs, as determined *via* powder X-ray diffraction, is not affected by the PSM (Fig. 1a). All reflections are present and the intensity ratios are comparable.

The investigation of the postsynthetically modified samples with energy-dispersive X-ray spectroscopy (EDX) showed a sulfur signal associated with the rhombic-shaped MOF crystals after postsynthetic modification with PEDOT (Section S3.3, ESI[†]). To quantify the amount of EDOT attached on the MOF surface, thermogravimetric and elemental (carbon and sulfur, C-S) analysis were performed. Thermogravimetric analysis (TGA) indicates a mass fraction of up to 2.8% of additional organic species (for the sample UV-irradiated for 72 h) (Section S3.4, ESI[†]). C-S analysis (Section S3.5, ESI[†]) gives a sulfur content of 0.76%, corresponding to a somewhat larger additional organic part of 3.4%. On the other hand, results from NMR spectroscopy (Section S3.6, ESI⁺) on acid-digested samples show that the vast majority of the keto groups - those located in the inner pores of the crystals - has not participated in the reaction, i.e. only the keto groups exposed on the outer surface have reacted. Using the size of the crystals and information from the crystal structure, the number of these exposed keto groups can be calculated. With the results from TGA and C-S analysis and using some assumptions we can estimate that the grafted-from PEDOT chains have lengths of at least ca. 100 EDOT units (for details of the calculations see Section S3.7, ESI[†]).

We postulate that the polymerization of EDOT proceeds *via* initiation by surface-standing ketyl radicals and subsequent oxidative polymerization (which is the common polymerization mechanism for EDOT; a corresponding reaction scheme presenting a postulated mechanism is described in Section S4 of the ESI†). Within this mechanism, the ethylene moiety of an EDOT molecule is used for its covalent attachment to the MOF surface. Subsequently, photo-excited benzophenone units of the linkers of



Scheme 2 Schematic mechanism for the photoreaction of the keto group of benzophenone units in the framework of Zr-*bzpdc*-MOF: formation of a ketyl radical which then reacts with a C–H bond-containing molecule (in this study EDOT), resulting in a covalent bond.



Fig. 1 Characterization of Zr-bzpdc-MOF and of samples postsynthetically modified using different UV irradiation times. (a) PXRD patterns with corresponding pictures of the powders. (b) $N_2@77$ K physisorption isotherms. (c) Electrical conductivity values.

the MOF oxidize further EDOT molecules, whereby the keto groups are reduced to alcohols. Usually, iron(m) salts or other oxidizing agents are used for the oxidative polymerization of EDOT.⁶⁷ However, the photo-oxidative potential of irradiated benzophenone is well-known in literature^{68–70} and presents a possible route to oxidative polymerization of EDOT to PEDOT without the necessity for other oxidizing agents (which are not present in our system). The polymerization of EDOT in its neat state without prior attachment to the MOF surface cannot be ruled out, but the resulting polymer chains would not be covalently bonded to the MOF crystals and should be removed by the extended extraction performed on the samples. Therefore, we propose that the PEDOT chains are covalently bonded on the MOF surface, leading to an electrically conductive composite material.

Composites between MOFs and polymers have been described extensively.^{71,72} In our case, the combination of porosity and electrical conductivity is the most exciting aspect of the composite materials described here. Physisorption measurements (Fig. 1b) show a slight decrease in pore volume and surface area which cannot be explained solely by the weight increase due to the mass of the attached polymer (Table S1, ESI[†]). Possibly, PEDOT molecules partially block the pores resulting in a slightly decreased pore volume. The isotherms also show a slight hysteresis which becomes more pronounced with longer irradiation time. Such hysteretic sorption behavior is indicative of the presence of a soft polymer which is deformed during adsorption.^{73,74} It can therefore be taken as an additional indication for a layer of PEDOT molecules at the MOF surface.

The electrical conductivity of the materials was measured using a van der Pauw setup on pressed pellets. The crystallinity of the framework is not affected by the pressure used to obtain the pressed pellets (Section S3.8, ESI†). The results of the electrical conductivity measurements are shown in Fig. 1c. Surprisingly, the non-modified Zr-*bzpdc*-MOF shows an electrical conductivity in a range of about 10^{-6} S cm⁻¹. Unexpectedly, this result places the Zr-*bzpdc*-MOF in the small group of intrinsically conducting MOFs. A possible explanation could be the generation of radicals at the keto groups, *e.g.* by former exposure to light (the actual measurement was performed in the dark under dry argon) or by the application of an electric potential during the measurements (the benzophenone moiety is known to be electrochemically reactive).⁷⁵ A measurable conductivity could then result from radical delocalization within the benzophenone units⁷⁶ and electron hopping between those. In fact, when we first deactivate the radical-forming benzophenone units by reacting them photochemically with ethanol (all benzophenone units within the crystals can be accessed in this way, Fig. S9, ESI[†]),⁶⁶ there is no measurable conductivity (Section S3.8, ESI[†]).

The samples irradiated in EDOT show much higher conductivities than the pristine Zr-*bzpdc*-MOF (Table S5, ESI†). The electrical conductivity increases during the first 24 h of irradiation by more than two orders of magnitude and then saturates in the range of 10^{-3} S cm⁻¹, with the highest value of 0.007 S cm⁻¹ observed for samples which had been irradiated for 72 h (Fig. 1c). This value is in the range of semiconducting materials and fits well for various applications, *e.g.* sensing.

In summary, we have demonstrated the postsynthetic photochemical modification of a MOF by a grafting-from reaction, which does not need any further preparatory steps, but can be carried out directly on the MOF crystals using the surfacestanding linker molecules. The resulting composite materials of the Zr-*bzpdc*-MOF with PEDOT show an interesting combination of moderate porosities and good electrical conductivities (Fig. S13, ESI[†]) so that applications as sensor material or as a biomaterial in neuronal electrodes appear feasible. This work thus reveals great potential for postsynthetic photochemical modifications reactions of the Zr-*bzpdc*-MOF and other MOFs containing photoreactive linkers.

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Conflicts of interest

There are no conflicts to declare.

Notes and references

- 1 U. Mueller, M. Schubert, F. Teich, H. Puetter, K. Schierle-Arndt and J. Pastré, *J. Mater. Chem.*, 2006, **16**, 626–636.
- 2 D. Farrusseng, S. Aguado and C. Pinel, Angew. Chem., Int. Ed., 2009, 48, 7502–7513.
- 3 J. Lee, O. K. Farha, J. Roberts, K. A. Scheidt, S. T. Nguyen and J. T. Hupp, *Chem. Soc. Rev.*, 2009, **38**, 1450–1459.
- 4 Z. Kang, L. Fan and D. Sun, J. Mater. Chem. A, 2017, 5, 10073-10091.

- 5 B. Li, H.-M. Wen, W. Zhou and B. Chen, J. Phys. Chem. Lett., 2014, 5, 3468–3479.
- 6 P. Kumar, A. Deep and K.-H. Kim, *Trends Anal. Chem.*, 2015, 73, 39–53.
- 7 M. Schulz, A. Gehl, J. Schlenkrich, H. A. Schulze, S. Zimmermann and A. Schaate, *Angew. Chem.*, *Int. Ed.*, 2018, 57, 12961–12965.
- 8 T. Simon-Yarza, A. Mielcarek, P. Couvreur and C. Serre, *Adv. Mater.*, 2018, **30**, 1870281.
- 9 M. Giménez-Marqués, T. Hidalgo, C. Serre and P. Horcajada, *Coord. Chem. Rev.*, 2016, **307**, 342–360.
- 10 V. Bon, Curr. Opin. Green Sustainable Chem., 2017, 4, 44-49.
- 11 K. K. Tanabe and S. M. Cohen, Chem. Soc. Rev., 2011, 40, 498-519.
- 12 T. Islamoglu, S. Goswami, Z. Li, A. J. Howarth, O. K. Farha and J. T. Hupp, Acc. Chem. Res., 2017, 50, 805–813.
- 13 P. Roy, A. Schaate, P. Behrens and A. Godt, *Chem. Eur. J.*, 2012, **18**, 6979–6985.
- 14 T. von Zons, L. Brokmann, J. Lippke, T. Preuße, M. Hülsmann, A. Schaate, P. Behrens and A. Godt, *Inorg. Chem.*, 2018, 57, 3348–3359.
- 15 B. Liu, M. Ma, D. Zacher, A. Bétard, K. Yusenko, N. Metzler-Nolte, C. Wöll and R. A. Fischer, J. Am. Chem. Soc., 2011, 133, 1734–1737.
- 16 S. M. Cohen, J. Am. Chem. Soc., 2017, 139, 2855-2863.
- 17 C. Chen, J. Kim, D.-W. Park and W.-S. Ahn, *Mater. Lett.*, 2013, **106**, 344–347.
- 18 K. Leus, Y.-Y. Liu, M. Meledina, S. Turner, G. van Tendeloo and P. van der Voort, *J. Catal.*, 2014, **316**, 201–209.
- 19 N. E. Thornburg, Y. Liu, P. Li, J. T. Hupp, O. K. Farha and J. M. Notestein, *Catal. Sci. Technol.*, 2016, 6, 6480–6484.
- 20 X. Wang, H. Li and X.-J. Hou, J. Phys. Chem. C, 2012, 116, 19814-19821.
- 21 H. Liu, H. Zhu and S. Zhu, Macromol. Mater. Eng., 2015, 300, 191-197.
- 22 K. A. McDonald, J. I. Feldblyum, K. Koh, A. G. Wong-Foy and A. J. Matzger, *Chem. Commun.*, 2015, **51**, 11994–11996.
- 23 L. Sun, M. G. Campbell and M. Dincă, Angew. Chem., Int. Ed., 2016, 55, 3566–3579.
- 24 C.-W. Kung, Y.-S. Li, M.-H. Lee, S.-Y. Wang, W.-H. Chiang and K.-C. Ho, J. Mater. Chem. A, 2016, 4, 10673–10682.
- 25 S. Lin, Y. Pineda-Galvan, W. A. Maza, C. C. Epley, J. Zhu, M. C. Kessinger, Y. Pushkar and A. J. Morris, *ChemSusChem*, 2017, 10, 514–522.
 26 M. G. Campbell and M. Dincă, *Sensors*, 2017, 17, 1108.
- 27 M. D. Allendorf, A. Schwartzberg, V. Stavila and A. A. Talin, *Chem. Eur. J.*, 2011, 17, 11372–11388.
- 28 S. K. Bhardwaj, N. Bhardwaj, R. Kaur, J. Mehta, A. L. Sharma, K.-H. Kim and A. Deep, *J. Mater. Chem. A*, 2018, **6**, 14992–15009.
- 29 P. Li and Bo Wang, Isr. J. Chem., 2018, 58, 1010-1018.
- 30 C. F. Leong, B. Chan, T. B. Faust and D. M. D'Alessandro, *Chem. Sci.*, 2014, 5, 4724–4728.
- 31 T. C. Narayan, T. Miyakai, S. Seki and M. Dincă, J. Am. Chem. Soc., 2012, 134, 12932–12935.
- 32 M. G. Campbell, S. F. Liu, T. M. Swager and M. Dincă, J. Am. Chem. Soc., 2015, 137, 13780–13783.
- 33 M. G. Campbell, D. Sheberla, S. F. Liu, T. M. Swager and M. Dincă, Angew. Chem., Int. Ed., 2015, 54, 4349–4352.
- 34 M. Hmadeh, Z. Lu, Z. Liu, F. Gándara, H. Furukawa, S. Wan, V. Augustyn, R. Chang, L. Liao, F. Zhou, E. Perre, V. Ozolins, K. Suenaga, X. Duan, B. Dunn, Y. Yamamto, O. Terasaki and O. M. Yaghi, *Chem. Mater.*, 2012, 24, 3511–3513.
- 35 X. Huang, P. Sheng, Z. Tu, F. Zhang, J. Wang, H. Geng, Y. Zou, C.A. Di, Y. Yi, Y. Sun, W. Xu and D. Zhu, *Nat. Commun.*, 2015, 6, 7408.
- 36 B. Hoppe, K. D. J. Hindricks, D. P. Warwas, H. A. Schulze, A. Mohmeyer, T. J. Pinkvos, S. Zailskas, M. R. Krey, C. Belke, S. König, M. Fröba, D. L. Haur and P. Pabrara, Contractional Conference on Contractional Contraction of Contraction of
- R. J. Haug and P. Behrens, *CrystEngComm*, 2018, 20, 6458–6471.
 F. Li, X. Zhang, X. Liu and M. Zhao, *ACS Appl. Mater. Interfaces*, 2018, 10, 15012–15020.
- 38 L. Sun, C. H. Hendon, S. S. Park, Y. Tulchinsky, R. Wan, F. Wang, A. Walsh and M. Dincă, *Chem. Sci.*, 2017, 8, 4450–4457.
- 39 J. G. Park, M. L. Aubrey, J. Oktawiec, K. Chakarawet, L. E. Darago, F. Grandjean, G. J. Long and Jeffrey R. Long, *J. Am. Chem. Soc.*, 2018, 140, 8526–8534.
- 40 M. L. Aubrey, B. M. Wiers, S. C. Andrews, T. Sakurai, S. E. Reyes-Lillo, S. M. Hamed, C.-J. Yu, L. E. Darago, J. A. Mason, J.-O. Baeg, F. Grandjean, G. J. Long, S. Seki, J. B. Neaton, P. Yang and J. R. Long, *Nat. Mater.*, 2018, 17, 625–632.

- 41 A. A. Talin, A. Centrone, A. C. Ford, M. E. Foster, V. Stavila, P. Haney, R. A. Kinney, V. Szalai, F. El Gabaly, H. P. Yoon, F. Léonard and M. D. Allendorf, *Science*, 2014, 343, 66–69.
- 42 S. Goswami, D. Ray, K.-I. Otake, C.-W. Kung, S. J. Garibay, T. Islamoglu, A. Atilgan, Y. Cui, C. J. Cramer, O. K. Farha and J. T. Hupp, *Chem. Sci.*, 2018, **9**, 4477–4482.
- 43 C. Lu, T. Ben, S. Xu and S. Qiu, Angew. Chem., Int. Ed., 2014, 53, 6454-6458.
- 44 H. Shiozawa, B. C. Bayer, H. Peterlik, J. C. Meyer, W. Lang and T. Pichler, *Sci. Rep.*, 2017, 7, 2439.
- 45 Y. Wang, L. Wang, W. Huang, T. Zhang, X. Hu, J. A. Perman and S. Ma, *J. Mater. Chem. A*, 2017, *5*, 8385–8393.
- 46 T. Wang, M. Farajollahi, S. Henke, T. Zhu, S. R. Bajpe, S. Sun, J. S. Barnard, J. S. Lee, J. D. W. Madden, A. K. Cheetham and S. K. Smoukov, *Mater. Horiz.*, 2017, 4, 64–71.
- 47 D. Fu, H. Zhou, X.-M. Zhang, G. Han, Y. Chang and H. Li, *ChemistrySelect*, 2016, 1, 285–289.
- 48 T.-Y. Huang, C.-W. Kung, Y.-T. Liao, S.-Y. Kao, M. Cheng, T.-H. Chang, J. Henzie, H. R. Alamri, Z. A. Alothman, Y. Yamauchi, K.-C. Ho and K. C.-W. Wu, *Adv. Sci.*, 2017, 4, 1700261.
- 49 C.-W. Kung, K. Otake, C. T. Buru, S. Goswami, Y. Cui, J. T. Hupp, A. M. Spokoyny and O. K. Farha, *J. Am. Chem. Soc.*, 2018, **140**, 3871–3875.
- 50 I. Ahmed and S. H. Jhung, Mater. Today, 2014, 17, 136-146.
- 51 O. Fleker, A. Borenstein, R. Lavi, L. Benisvy, S. Ruthstein and D. Aurbach, *Langmuir*, 2016, 32, 4935–4944.
- 52 P. Freund, I. Senkovska and S. Kaskel, *ACS Appl. Mater. Interfaces*, 2017, 9, 43782–43789.
- 53 Y. Mao, G. Li, Y. Guo, Z. Li, C. Liang, X. Peng and Z. Lin, *Nat. Commun.*, 2017, **8**, 14628.
- 54 X. Xu, W. Shi, P. Li, S. Ye, C. Ye, H. Ye, T. Lu, A. Zheng, J. Zhu, L. Xu, M. Zhong and X. Cao, *Chem. Mater.*, 2017, **29**, 6058–6065.
- 55 B. Le Ouay, M. Boudot, T. Kitao, T. Yanagida, S. Kitagawa and T. Uemura, *J. Am. Chem. Soc.*, 2016, **138**, 10088–10091.
- 56 R. Haldar, B. Sen, S. Hurrle, T. Kitao, R. Sankhla, B. Kühl, A. Welle, S. Heissler, G. Brenner-Weiß, P. Thissen, T. Uemura, H. Gliemann, C. Barner-Kowollik and C. Wöll, *Eur. Polym. J.*, 2018, **109**, 162–168.
- 57 T. C. Wang, I. Hod, C. O. Audu, N. A. Vermeulen, S. T. Nguyen, O. K. Farha and J. T. Hupp, ACS Appl. Mater. Interfaces, 2017, 9, 12584–12591.
- 58 G. Zahn, H. A. Schulze, J. Lippke, S. König, U. Sazama, M. Fröba and P. Behrens, *Microporous Mesoporous Mater.*, 2015, 203, 186–194.
- 59 A. Schaate, P. Roy, A. Godt, J. Lippke, F. Waltz, M. Wiebcke and P. Behrens, *Chem. – Eur. J.*, 2011, **17**, 6643–6651.
- 60 J. E. Mondloch, M. J. Katz, N. Planas, D. Semrouni, L. Gagliardi, J. T. Hupp and O. K. Farha, *Chem. Commun.*, 2014, 50, 8944–8946.
- 61 Y. Bai, Y. Dou, L.-H. Xie, W. Rutledge, J.-R. Li and H.-C. Zhou, *Chem. Soc. Rev.*, 2016, **45**, 2327–2367.
- 62 A. Mohmeyer, A. Schaate, B. Brechtken, J. C. Rode, D. P. Warwas, G. Zahn, R. J. Haug and P. Behrens, *Chem. – Eur. J.*, 2018, 24, 12848–12855.
- 63 D. Karaca Balta, Ö. Karahan, D. Avci and N. Arsu, Prog. Org. Coat., 2015, 78, 200-207.
- 64 M. A. Winnik and U. Maharaj, Macromolecules, 1979, 12, 902-905.
- 65 O. Prucker, C. A. Naumann, J. Rühe, W. Knoll and C. W. Frank, J. Am. Chem. Soc., 1999, 121, 8766–8770.
- 66 A. Mohmeyer, M. Schäfer, A. Schaate, S. Locmelis and P. Behrens, unpublished results.
- 67 T. Horii, H. Hikawa, M. Katsunuma and H. Okuzaki, *Polymer*, 2018, 140, 33–38.
- 68 N. Filipescu and F. L. Minn, J. Am. Chem. Soc., 1968, 90, 1544–1547.
- 69 B. Qu, Y. Xu, L. Ding and B. Rånby, J. Polym. Sci., Part A: Polym. Chem., 2000, 38, 999–1005.
- 70 A. Demeter, K. Horváth, K. Böör, L. Molnár, T. Soós and G. Lendvay, J. Phys. Chem. A, 2013, 117, 10196–10210.
- 71 T. Kitao, Y. Zhang, S. Kitagawa, B. Wang and T. Uemura, *Chem. Soc. Rev.*, 2017, **46**, 3108–3133.
- 72 S. Mochizuki, T. Kitao and T. Uemura, Chem. Commun., 2018, 54, 11843–11856.
- 73 Y. Tsujita, Prog. Polym. Sci., 2003, 28, 1377-1401.
- 74 J. Weber, M. Antonietti and A. Thomas, *Macromolecules*, 2008, **41**, 2880–2885.
- 75 N. G. Tsierkezos and U. Ritter, Phys. Chem. Liq., 2011, 49, 729-742.
- 76 Y. Du, J. Xue, M. Li and D. L. Phillips, J. Phys. Chem. A, 2009, 113, 3344–3352.

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