Although several recent publications describe cathodes for electrochemical energy storage materials made from regrown biomass in aqueous electrolytes, their transfer to lithium–organic batteries is challenging. To gain a deeper understanding, we investigate the influences on charge storage in model systems based on biomass-derived, redox-active compounds and comparable structures. Hybrid materials from these model polymers and porous carbon are compared to determine precisely the causes of exceptional capacity in lithium–organic systems. Besides redox activity, particularly, wettability influences capacity of the composites greatly. Furthermore, in addition to biomass-derived molecules with catechol functionalities, which are described commonly as redox-active species in lithium–bio-organic systems, we further describe guaiacol groups as a promising alternative for the first time and compare the performance of the respective compounds.

Introduction

Guaiacol groups are common in biomolecules, both in small molecules such as vanillin as well as in macromolecules such as lignin. Although they are not redox-active per se, irreversible oxidation in water results in the formation of the o-quinone/o-hydroquinone redox pair. This reaction was first applied prominently by Milczarek and Inganäs in cathode materials from lignin, which has abundant guaiacol units, for green electrochemical energy storage devices. Since then, many endeavors to improve lignin-based cathodes have been undertaken. Additionally, cathodes based on polymers from bio-refined monomers such as guaiacol and syringol phenolic acids and vanillin have been investigated in acidic aqueous electrolytes. Aqueous electrolytes are important for energy storage in these materials as the established process for the formation of quinones requires the presence of water. Recently, a lignin–PEDOT (PEDOT = poly(3,4-ethylenedioxythiophene)) hybrid material was investigated as an electrode material in a lithium–organic system. However, lignin mainly serves to dope PEDOT in this system. PEDOT constitutes the majority of the electrode, which limits the sustainability of such a material.

In polymer-based electrodes, redox-active polymers are always mixed with conductive additives such as carbon nano-tubes, porous carbons, and conductive polymers, all of which contribute significant capacitive energy storage. Both cyclic voltammograms and galvanostatic measurements of the mixed cathode materials are used to show influences of both distinct redox-active groups (high current only at distinct voltages in cyclic voltammetry and plateau-like galvanostatic behavior) and capacitive behavior (rectangular cyclic voltammogram and triangular galvanostatic curve). Consequently, it is debatable whether the devices should be denoted as supercapacitors or batteries.

In studies on organic electrode materials, conductive additives such as carbon materials are often considered passive parts of the electrode composition. To describe charge storage in organic materials more thoroughly, sometimes electrodes without organic active materials are designed, and the resulting capacity is subsequently subtracted from the total capacity. However, such investigations suffer from the possible systematic error when neglecting the modification of charge storage properties of carbon additives by organic active materials.

The rapidly increasing number of publications on bio-derived active electrode materials requires the discussion of such possible influences in more detail. Consequently, we investigate influences of different components in common biopolymer-based materials for electrochemical energy storage. As model systems, we describe hybrid energy storage materials made from model biopolymers and carbon additives. In the former, several low-molecular-weight compounds, which have active functionalities common to biomolecules, are immobilized on a polymer backbone. For immobilization, we make use of the formation of Schiff bases, which can be used to graft aldehydes onto amine-containing polymers, most notably, chitosan and polyallylamine. Vanillin is investigated because it contains guaiacol groups. Protocatechuic aldehyde.
contains a catechol functionality, that is, it may be formed upon the demethylation of the guaiacol unit in vanillin. Finally, 3,5-dihydroxybenzaldehyde is investigated because it is an isomer of protocatechuic aldehyde. However, as the hydroxyl groups are found in the meta instead of the ortho position, it is not redox-active.

The processing of active polymeric materials with conductive additives, for example, by the preparation of a slurry that contains both components, spreading on a current collector, and drying, is a conventional way to create organic batteries. For the formation of electrodes, we chose microporous carbons as conductive additives because of the possibility to synthesize them from biowaste, which makes them an attractive sustainable conductive additive, and because of their intrinsically high capacitance that allows the formation of hybrid electrochemical energy storage devices. The contributions of carbon, the wettability of the electrolyte, and different functional units to the resulting lithium-ion-based energy storage devices are discussed. Firstly, we characterize the polymers under investigation and then their composites with microporous carbon to assess the influence of different physical properties and functionalities on charge storage. Finally, the electrochemical performance of these composites are investigated and compared.

Results and Discussion

Synthesis of polymers and their hybrid materials

Schiff bases of polyallylamine (PAAm) and different aldehydes, namely, vanillin (van), protocatechuic aldehyde (which features two phenolic groups in the ortho position; A-o), and 3,5-dihydroxybenzaldehyde (which features two phenolic groups in the meta position; A-m), were synthesized in ethanolic solutions as described in the Experimental Section and Supporting Information (Scheme 1). The addition of aldehydes to PAAm results in an immediate color change as expected for the formation of Schiff bases, followed by precipitation (Figure S1) caused by the interaction of leftover amine group with acidic phenolic groups. During subsequent drying, the formation of Schiff bases proceeds, and washing with anhydrous ethanol yields the pure modified polymers (denoted P-van, P-o, and P-m, respectively).

The successful formation of Schiff bases is confirmed by using IR spectroscopy (Figure 1), as the resulting polymers contain the functional groups from the respective phenyl ring. Broad bands above $\tilde{\nu} = 3000 \text{ cm}^{-1}$, from hydroxyl groups, are retained both in the spectra of the polymers and aldehydes, which confirms the retention of the hydroxyl groups during synthesis. Two new bands appear in the spectra of polymers at approximately $\tilde{\nu} = 2840$ and $2920 \text{ cm}^{-1}$, which indicates C–H stretching vibrations from the polyallylamine backbone. No aldehyde groups are detected in the spectra of the polymers, but new bands at approximately $\tilde{\nu} = 1680 \text{ cm}^{-1}$ represent imine functionalities in all three samples, which confirms both the successful reaction and the successful removal of any unreacted aldehydes. Intriguingly, the shift between the imine and aldehyde peak is very pronounced in the case of van/P-van and A-m/P-m but is rather subtle in the case of A-o/P-o, only by a few cm$^{-1}$, which originates from the different wave-number of the aldehyde carbonyl vibrations in A-o, van, and A-m. The substitution rates in the samples (details are explained in the Supporting Information) calculated using the amount of N determined by using elemental analysis is sum-

![Scheme 1. Reaction of polyallylamine with different aldehydes.](image1)

![Figure 1. IR spectra of A-o, van, A-m, and their corresponding Schiff bases with PAAm in the range of 1000–4000 cm$^{-1}$.](image2)
Comparison of the electrochemical performance of redox-active and -inactive polymers

C/P-o, C/P-m, and microporous carbon are compared in Figure 2 in terms of their electrochemical performance during cyclic voltammetry (CV) and charge–discharge tests. The redox activity of C/P-o (Scheme S2) is clearly detectable in the CV from peaks at approximately 3.3 V versus Li+/Li (Figure 2a). In the low-voltage region, the voltammograms of C/P-o and C/P-m match fairly well, but there are no redox peaks at approximately 3.3 V versus Li+/Li observable in the trace of C/P-m, as expected for redox-inactive species. Similarly, the absence of redox peaks is characteristic of purely microporous carbon.

The capacity of microporous carbon in galvanostatic charge–discharge experiments is very low. Upon the addition of P-m and even more so upon the addition of P-o (C/P-m and C/P-o, respectively), the capacity increases significantly. Upon prolonged cycling, C and C/P-m retain their capacity, whereas C/P-o shows subsequent decay (Figure 2c), which will be discussed later. The higher capacity of C/P-o than C/P-m can be explained easily by the redox activity of P-o that results in a slightly bell-shaped galvanostatic discharge behavior of C/P-o (Figure 2b) and a significantly higher capacity. The range of the increased discharge capacity in C/P-o compared to that of C/P-m (nonparallel discharge curves shown in Figure 2b) at approximately 2.7–3.8 V versus Li+/Li matches the location of the discharge peaks in the CVs of C/P-o (Figure 2a). Peaks in the latter correspond to the redox reaction between catechol and o-hydroquinone groups, which is assigned to increased capacity (C/P-o compared to C/P-m) in Figure 2b because of the influence of redox reactions. No belly-shaped behavior can be observed for C and C/P-m (Figure 2b). A comparison of the electrochemical impedance spectra (EIS) of C/P-o, C/P-m, and C (Figure S6) shows the similar behavior of the composites. The Nyquist plots show the good conductivity and permeability of all the samples as expected, and C has the fastest kinetics as no polymer blocks its pores.

However, the significantly higher capacity observed if we compare C/P-m and microporous carbon cannot be explained by redox reactions as P-m is not redox-active. A possible explanation for this may be the increased wettability of the

![Graphs](Figure 2. Comparison of the electrochemical performance of C/P-o, C/P-m, and porous carbon (C). The tests were performed in a lithium half-cell setup with lithium as counter electrode and 1 M LiPF₆ in EC/DEC (1:1) as electrolyte. a) CVs at 5 mV s⁻¹, 10⁶th cycle. b) Charge–discharge tests at 0.2 Ag⁻¹, 10⁶th cycle. c) Discharge capacities calculated from the charge–discharge tests at 0.2 Ag⁻¹.)
carbon surfaces by the electrolyte after functionalization with the heteroatom-rich polymer. Consequently, next we discuss the N₂ and water vapor physiosorption behavior to assess the hydrophilicity of the prepared hybrid materials and, therefore, their wettability with solvents of high polarity, such as electrolytes (Figure 3).

During ball-milling, C loses most of its internal surface area, probably caused by a combination of high-energy ball-milling and the blocking of the pores by the binder (cf., physiosorption behavior of the pristine microporous carbon in Figure S7). The Brunauer–Emmett–Teller (BET) surface area of the hybrid materials with the model polymers (C/P-o and C/P-m) are even lower than that of C, possibly because of the relatively low amount of microporous carbon in the hybrid material and because of the additional blocking of the pores by the active polymers. All samples exhibit high external surface areas because of the loose packing of particles.

Water vapor physiosorption does not follow the same trend as N₂ physiosorption. The water vapor uptake in C/P-o and C/P-m is similarly high, and the carbon material adsorbs significantly less water vapor than C/P-o and C/P-m over the entire range of relative pressure. As the elemental composition of P-o and P-m is the same, this behavior may be ascribed to abundant hydroxyl groups in the polymers, which results in mixed carbon-polymer materials with a similar hydrophilicity. Hydrophilicity is significantly higher in the hybrid materials than in the carbon material, and subsequently, as a result of the high polarity of water and electrolytes, so is the wettability by polar electrolytes. As the electrolyte reaches a larger part of the internal surface of the microporous carbon, the contribution to energy storage by the formation of the electric double layer increases. More charge can be stored on the surface for both C/P-o and C/P-m compared to carbon, which leads to a higher capacity. Similar behavior was described previously in aqueous solutions of polypyrrole with and without chitosan, in which the redox-inactive chitosan increased the capacity of polypyrrole significantly. Notably, ball-milling tends to increase the hydrophilicity of carbon materials, although to a low extent.

Comparison of the electrochemical performance of catechol and guaiacol groups

Catechol groups in hybrid materials with microporous carbons contribute to the overall capacity both through the increased wettability of carbon surfaces by the electrolytes and redox activity. Still, catechol groups are rather rare in nature, notable exceptions are dopamine and tannic acid, which have been described previously for electrochemical energy storage.

In contrast, guaiacol groups (which are chemically similar; cf. Scheme 1) are abundant in nature, for example, in low-value biomass materials such as lignin. To our knowledge, such materials have never been used in secondary lithium-ion batteries without additional redox active polymers to date. Thus, we will focus on such materials by comparing the electrochemical performance of C/P-o and C/P-van (Figure 4). Initially, the discharge curves differ greatly, both in terms of overall capacity as well as shape, but become more similar with the increase of the number of cycles. Unlike that of C/P-o, the discharge curve of C/P-van shows no belly shape at the beginning. With the increase of the number of cycles, the shape changes, and ultimately, almost matches that of the C/P-o discharge curve after approximately 50 charge–discharge cycles. The same behavior is apparent in the CV curves (Figure S8). Although no clear redox peaks are observed in the early cycles of C/P-van, such peaks at similar potential as seen in the CV of C/P-o appear with continuing cycling, and the CVs of C/P-van and C/P-o start to resemble each other. We performed EIS for both samples (Figure S6), which shows the good conductivity of the materials, and C/P-van exhibits slightly slower kinetics possibly because of the presence of methoxy groups.

In aqueous systems, demethylation occurs in the first oxidation step, expressed by the high current in the CV curves at voltages approximately 0.2 V higher than the reversible redox peak that evolves in the next reduction cycle. A quinone–hydroquinone redox pair thus can be formed easily also in molecules that initially contain guaiacol groups. As this mechanism requires the presence of water, it is impossible in a lithium-ion-containing setup in water-free organic electrolytes. Consistently, no such first irreversible oxidation peak is observable by using CV in our setup. Still, galvanostatic and CV curves of C/P-van indicate the formation of the same quinone–hydroquinone redox couple as that in C/P-o with prolonged cycling (Figure 4 and Figure S8) probably because of the slow demethylation that exposes redox-active functionalities. The mechanism for...
the involved demethylation process is unclear; however, we suggest a similar mechanism to that found in aqueous systems, with the difference that fluoride anions serve as nucleophiles instead of water and form fluoromethane in the process. As no significant irreversible oxidation peak is observed in a distinct early CV cycle (Figure S9 and b), unlike in the case of similar polymers in aqueous systems (Figure S9 c), we conclude that demethylation requires several charge–discharge cycles in a lithium half-cell setup. Irreversible oxidation, which usually results in diminishing capacity with prolonged cycling, is common in organic battery materials.

The different charge storage behavior with continuous cycling may be explained by differences in the sleeping phases between both materials. The oxidation of hydroquinone groups happens much faster than that of guaiacol groups, which results in the complete transition of C/P-o from the sleeping phase to the living phase within a few cycles. The subsequent fading capacity indicates a slow transition to the dead phase. In contrast, the constant capacity of C/P-van after the maximum is reached, which may be ascribed to the continuous slow transition of some guaiacol groups from the sleeping to living phase at a similar rate as the transition of other active groups from the living phase to the dead phase. Consequently, this does not necessarily indicate the higher stability of the material during cycling. Ultimately, the maximum capacity is lower in C/P-van than C/P-o as partial demethylation (formation of the living species) continues after the maximum capacity is reached. Only after approximately 50 cycles do the capacities of C/P-m and C/P-o become similar because of the formation of quinone groups in C/P-van and the irreversible oxidation of C/P-o. The final apparent specific capacity of C/P-van is still slightly lower than that of C/P-o because the specific capacity is calculated from the initial mass of the hybrid material, which includes the methyl groups in P-van. The demethylation of guaiacol groups reduces the weight of the polymer material slightly, which makes the assumed gravimetric capacity an underestimation.

New promising sustainable cathode material: C/P-o

We next evaluate the electrochemical behavior of the best-performing material C/P-o in detail. If we compare the charge storage behavior of C/P-o, C/P-m, and C (Figure 2b), the influences of carbon, redox activity, and changes in wettability may be estimated as discussed in more detail in the Supporting Information. According to the galvanostatic charge–discharge experiments, after 10 cycles, reversible redox reactions contribute to 18.1% of the charge storage of the total electrode material. With only 13.0% derived from capacitive charge storage...
on the surface of the unmodified carbon material, changes in wettability of the surface have a major influence on charge storage (68.9%). Abundant phenolic functionalities and imines result in significantly enhanced hydrophilicity, which facilitates capacitive charge storage using polar electrolytes. These results point to the importance of the selection of appropriate binders and carbons in the design of electrode materials from renewable resources.

Further information on the electrochemical behavior of C/P-o is given in Figure 5. We performed charge–discharge tests at different current densities (Figure 5a) between 0.05 and 0.80 A g\(^{-1}\) (see Experimental Section for details). Within the first three cycles, the capacity increases and starts to decrease slowly afterwards, as discussed above. At higher charge–discharge rates up to 0.80 A g\(^{-1}\), the capacity decreases only moderately and is restored to almost initial values after the current density returns to 0.05 A g\(^{-1}\), which prominently supports diffusion- and surface-controlled charge storage. Discharge curves at different current densities (Figure 5b) generally appear as almost straight lines with a slight indication of a belly shape, which indicates dominant capacitive behavior with some added distinct quinone–hydroquinone redox reactions, respectively.

CV curves of C/P-o at different rates are presented in Figure 5c. As the integrated area of the CV curve increases slowly over some tens of cycles (Figure S10), for better comparability the hybrid material was cycled firstly for 100 cycles at 25 mV s\(^{-1}\) followed by cycling at 20, 15, 10, 5, and 0.5 mV s\(^{-1}\) for one cycle each. CV curves show clear oxidation and reduction peaks at 3.6 and 2.8 V, respectively, which match the range of the belly-shaped behavior in the galvanostatic charge–discharge curves. From the performance at different rates at 5 mV s\(^{-1}\) as calculated from the CV at different rates (see Supporting Information for details). e) Long-term stability test at 0.1 A g\(^{-1}\) (every third data point is shown).
cycling speeds, the ratio of Faradaic charge storage and double layer capacitance was calculated (details can be found in the Supporting Information). Except for the limits between which the CV experiments were performed, the resulting curve for the capacitive contribution to charge storage (Figure 5d, black curve) resembles the CV of C/P-m (Figure 2a, blue curve). In both cases, the capacity is approximately 80% as high as that of C/P-o (84.8% cf. Figure 2a, red curve and 74.8% cf. Figure 5d green curve, respectively), which confirms the importance of hydrophilicity on the charge storage and matching the results of galvanostatic experiments (Faradaic reactions in C/P-o are responsible for 18.1% of the charge storage). Clearly, the addition of P-o to carbon not only introduces the possibility of Faradaic charge storage but also facilitates capacitive charge storage. The notable differences between the calculated capacitive contribution to charge storage shown in Figure 5d and the charge storage in C/P-m (Figure 2a, blue curve) at low and high voltages can be ascribed to the limitations of the assumed model, expressed by a low coefficient of determination ($R^2$; Figure S11) caused by the sudden increase or decrease in current density at the edges of the CVs. Slight differences between the capacitive charge storage in C/P-m and calculated capacitive contribution to C/P-o as denoted above can be ascribed primarily to the aforementioned collapse of the model at the edges of the CVs.

During long-term cycling at 0.1 Ag$^{-1}$ (Figure 5e), the capacity decreases slowly as is observed generally for organic systems (see discussion above). However, a remarkable retention of 73.5% is observed after 200 cycles (on average 0.13% decay of capacity per cycle). The high Coulombic efficiency of approximately 97% makes P-o a promising polymer for electrochemical energy storage applications, comparable to other biomass-derived batteries.[46, 47]

Conclusions

Polymers made from polyallylamine and redox-active and in-active aldehydes were used as model compounds for bioenergetic polymers to elucidate the different contributions to electrochemical energy storage in future bio-based batteries and lithium ion capacitors. In the systems under investigation, which included carbon, binder, and the (in)active polymer, only 18.1% of the capacity could be ascribed to distinct redox reactions. All samples showed a significantly higher capacity than microporous carbon itself, and the difference was ascribed to the increased hydrophilicity of the capacitive hybrid material that causes better wettability by the electrolyte and better lithium transport behavior. Therefore, we conclude that a significant portion of the capacity of electrochemical energy storage devices based on natural polyphenols stems from capacitive charge storage on hydrophilic surfaces in addition to redox activity. Furthermore, the electrochemical performance in a lithium-ion-containing setting for the model polymers that contain redox-active catechol groups was compared to that with naturally abundant guaiacol groups. Both showed a similar performance after prolonged cycling, with a slower initial increase in capacity for the guaiacol-containing polymers because of their slow demethylation processes. From these findings, a biomass-derived hybrid material was prepared that showed excellent properties for electrochemical energy storage in lithium-organic systems with a capacity of over 100 mA h g$^{-1}$ at 0.05 Ag$^{-1}$ and capacity retention of 73.5% after 200 cycles at 0.1 Ag$^{-1}$.

Experimental Section

Materials

Protocatechuic aldehyde (A-o, Carl Roth), vanillin (van, Sigma–Aldrich), 3,5-dihydroxybenzaldehyde (A-m, Fischer), an aqueous solution (15%) of polyallylamine with an average molecular weight of 15 000 g mol$^{-1}$ (PAAm, Polysciences, Inc.), carbon EQ-AB-520Y (C, MTI Corporation), carbon paper Spectracarb 2050A-0550 (Fuel Cell Store), and the solvents ethanol (Fischer) and N-methyl-2-pyrroldione (NMP, PanReac AppliChem) were used as received.

Synthesis of P-o$^{[24]}$

An aqueous solution of PAAm (0.33 g, 0.87 mmol of repeating units) was added to ethanol (10 mL) and stirred for 10 min. After the addition of A-o (0.12 g, 0.87 mmol) dissolved in ethanol (10 mL), the mixture was stirred for 1 h. Afterwards the solvent was removed at 60 °C at a pressure of 180 mbar, which was lowered to 150 mbar shortly after the beginning of the solvent removal. After the obtained polymer was dried at 80 °C under vacuum for 2 h, it was washed with ethanol by centrifugation (4 times with 50 mL ethanol, 4000 rpm, for 5 min). The sample was dried in a vacuum oven overnight. More information on the synthesis in included in the Supporting Information.

TGA and combustive elemental analysis

TGA measurements were performed by using a NETZSCH TG 209F1 Libra machine under N$_2$ at a heating rate of 10 K min$^{-1}$. Combustive elemental analysis was performed by using a varioMicro CHNS machine.

Morphology characterization

The morphology at the surface was investigated by SEM by using a Zeiss Leo Gemini 1550 microscope, and the atomic distribution of C, O, and N was mapped by using energy-dispersive X-ray spectroscopy (EDX, X-Max, Oxford instruments).

Physiosorption

Before the physiosorption measurements, the samples were degassed under vacuum at 80 °C for 16 h. N$_2$ physiosorption measurements were performed by using a Quantachrome Quadrasorb SI physisorption instrument at 77 K, whereas water vapor physiosorption was measured by using a Quantachrome Autosorb IQ physiosorption instrument at 298 K.

Spectroscopy

FTIR spectroscopy was performed by using a Nicolet iS 5 FTIR spectrometer (ThermoFisher Scientific).
Electrochemical measurements

All electrochemical measurements were performed in Swagelok-type cells and tested using a BioLogic MPG2 potentiostat. Lithium foil, Celgard 2325 (13 mm in diameter, 25 μm thick), and 1 M LiPF₆ in ethylene carbonate (EC) and diethyl carbonate (DEC; volumetric ratio 1:1, 100 μl; solution used as obtained from Sigma–Aldrich) were used as counter electrode, membrane, and electrolyte, respectively. A circular carbon current collector covered with the active material was used as the working electrode. The electrolyte was chosen as it is one of the most common choices for carbonyl-based organic lithium-ion batteries. All Swagelok-type cells were assembled in a glovebox with low water and O₂ levels. Charge-discharge measurements were performed at a constant current density (0.2 Ag⁻¹ for 50 cycles and 0.1 Ag⁻¹ for 200 cycles) or at varying current density (20 cycles at 0.05 Ag⁻¹, 10 cycles at 0.10 Ag⁻¹, 10 cycles at 0.20 Ag⁻¹, 10 cycles at 0.40 Ag⁻¹, 10 cycles at 0.80 Ag⁻¹) in triplicate. CV was performed by cycling at 25 mVs⁻¹ for 100 cycles and at 20, 15, 10, 5, and 0.5 mVs⁻¹ subsequently for one cycle each. In the displayed CV curves, the last cycle at the respective speed is shown unless mentioned otherwise.

Acknowledgements

We acknowledge help in the laboratory by Jessica Brandt. We furthermore thank Antje Völkel for help with elemental analysis and TGA measurements, and Markus Antonietti, Nieves Lopez Salas, and Martin Oschatz for fruitful discussions. We appreciate financial support by the German Research Foundation (DFG; grant: LI 2526/4-1), the Max Planck Society, and the Max Planck Institute of Colloids and Interfaces. Lastly, we would like to thank the Chez Briel-Institution for substantial support.

Conflict of interest

The authors declare no conflict of interest.

Keywords: biomass · electrochemistry · energy conversion · polymers · redox chemistry

[1] D. Esposito, M. Antonietti, Chem. Soc. Rev. 2015, 44, 5821 – 5835.
[2] G. Milczarek, Langmuir 2009, 25, 10345–10353.
[3] G. Milczarek, O. Inganäs, Science 2012, 335, 1468 – 1471.
[4] S. Admassie, T. Y. Nilsson, O. Inganäs, Phys. Chem. Chem. Phys. 2014, 16, 24681 – 24684.
[5] D. H. Nagaraju, T. Rebis, R. Gabrielson, A. Elfwing, G. Milczarek, O. Inganäs, Adv. Energy Mater. 2014, 4, 1300443.
[6] H. Xu, H. Jiang, X. Li, G. Wang, RSC Adv. 2015, 5, 76116 – 76121.
[7] T. Y. Nilsson, M. Wagner, O. Inganäs, ChemSusChem 2015, 8, 4081 – 4085.
[8] T. Rebis, G. Milczarek, Electrochem. Acta 2016, 204, 108 – 117.
[9] S. Chaleawlert-umporn, T. Berthold, X. Wang, M. Antonietti, C. Liedel, Adv. Mater. Interfaces 2017, 4, 1700989.
[10] S. Chaleawlert-umporn, C. Liedel, J. Mater. Chem. A 2017, 5, 24344 – 24352.
[11] S. K. Kim, Y. K. Kim, H. Lee, S. B. Lee, H. S. Park, ChemSusChem 2014, 7, 1094 – 1101.