

Cite this: *Chem. Sci.*, 2019, 10, 8889

All publication charges for this article have been paid for by the Royal Society of Chemistry

Zinc ion interactions in a two-dimensional covalent organic framework based aqueous zinc ion battery†

Abdul Khayum M,^{ab} Meena Ghosh,^{ab} Vidyanand Vijayakumar,^{ab} Arjun Halder,^{ab} Maryam Nurhuda,^c Sushil Kumar,^b Matthew Addicoat,^c Sreekumar Kurungot^{ab} and Rahul Banerjee^{cd}

The two-dimensional structural features of covalent organic frameworks (COFs) can promote the electrochemical storage of cations like H⁺, Li⁺, and Na⁺ through both faradaic and non-faradaic processes. However, the electrochemical storage of cations like Zn²⁺ ion is still unexplored although it bears a promising divalent charge. Herein, for the first time, we have utilized hydroquinone linked β -ketoenamine COF acting as a Zn²⁺ anchor in an aqueous rechargeable zinc ion battery. The charge-storage mechanism comprises of an efficient reversible interlayer interaction of Zn²⁺ ions with the functional moieties in the adjacent layers of COF (−182.0 kcal mol^{−1}). Notably, due to the well-defined nanopores and structural organization, a constructed full cell, displays a discharge capacity as high as 276 mA h g^{−1} at a current rate of 125 mA g^{−1}.

Received 20th June 2019
Accepted 2nd August 2019

DOI: 10.1039/c9sc03052b

rsc.li/chemical-science

Introduction

Two-dimensional (2D) porous crystalline covalent organic frameworks (COFs) are integrated from pre-designed symmetric organic building units and self-assembled by π - π stacking.¹ Due to the structural tunability, well-defined porosity, and high chemical stability, COFs have been widely explored as electrode materials for electrochemical energy storage.² However, until now, the energy storage applications in COFs were mostly limited to the monovalent ions like H⁺, Li⁺, and Na⁺ that could only carry forward single electron reactions. Although, divalent aqueous zinc ion batteries are observed to be cost-effective and superior in terms of safety and recyclability than the monovalent Li⁺, Na⁺ metal-ion batteries,³ it is extremely challenging to design an organic polymeric electrode that can act as a Zn²⁺ acceptor. A few literature reports deal with small molecules as cathode materials for accepting the electrophilic Zn²⁺ ions through coordination bonds.⁴ However, limited chemical and thermal stabilities of these organic molecules and the undesirable leaching of the active electrode materials during the

electrochemical cycling hampers their performance and creates a need for a polymeric electrode capable of hosting Zn²⁺ ions. Keeping this in perspective, herein, we have demonstrated hydroquinone based covalent organic framework⁵ (**HqTp**) acting as an efficient polymeric cathode material for an aqueous zinc ion battery. We anticipated that a COF must be enriched with several nucleophilic centers so that it can anchor to a significant number of divalent Zn²⁺ ions. Furthermore, the framework must be constructed from such building blocks, which can act as nucleophilic centers for efficient interaction with Zn²⁺ under electrochemical conditions.

It is well documented that the C=O and N-H functionalities in polymers are capable of coordinating Zn²⁺ ions in an electrochemical cell.^{4,6} Both C=O \cdots Zn and Zn \cdots N-H interactions are reversible in nature and can “make and break” during the electrochemical redox process. Thus, an electron-rich backbone in the **HqTp** facilitates efficient coordination with Zn²⁺ ions. **HqTp** exhibits an excellent discharge capacity of 276 mA h g^{−1} at a current of 125 mA g^{−1} in an operating potential of 0.2 to 1.6 V vs. Zn/Zn²⁺ due to the efficient intermolecular O \cdots Zn (\sim 2.0 Å) and N \cdots Zn (\sim 2.0 Å) interactions.^{4a,7} To the best of our knowledge, this is the first report of a COF based aqueous zinc ion battery and puts forward a novel concept based on the zinc ion interaction with a porous crystalline and polymeric organic cathode.

Results and discussion

The **HqTp** COF is synthesized by a solid-state mechano-mixing of 2,5-diaminohydroquinone dihydrochloride (**Hq**) and 1,3,5-triformylphloroglucinol (**Tp**) in the presence of *p*-

^aAcademy of Scientific and Innovative Research (AcSIR), Sector 19, Kamla Nehru Nagar, Ghaziabad, Uttar Pradesh-201002, India

^bPhysical and Materials Chemistry Division, CSIR-National Chemical Laboratory, Dr. Homi Bhabha Road, Pune-411008, India. E-mail: k.sreekumar@ncl.res.in

^cSchool of Science and Technology, Nottingham Trent University, Clifton Lane, NG11 8NS Nottingham, UK

^dDepartment of Chemical Sciences, Indian Institute of Science Education and Research (IISER), Mohanpur Campus, Mohanpur, Kolkata, 741252 India. E-mail: r.banerjee@iiserkol.ac.in

† Electronic supplementary information (ESI) available: Experimental section. See DOI: 10.1039/c9sc03052b



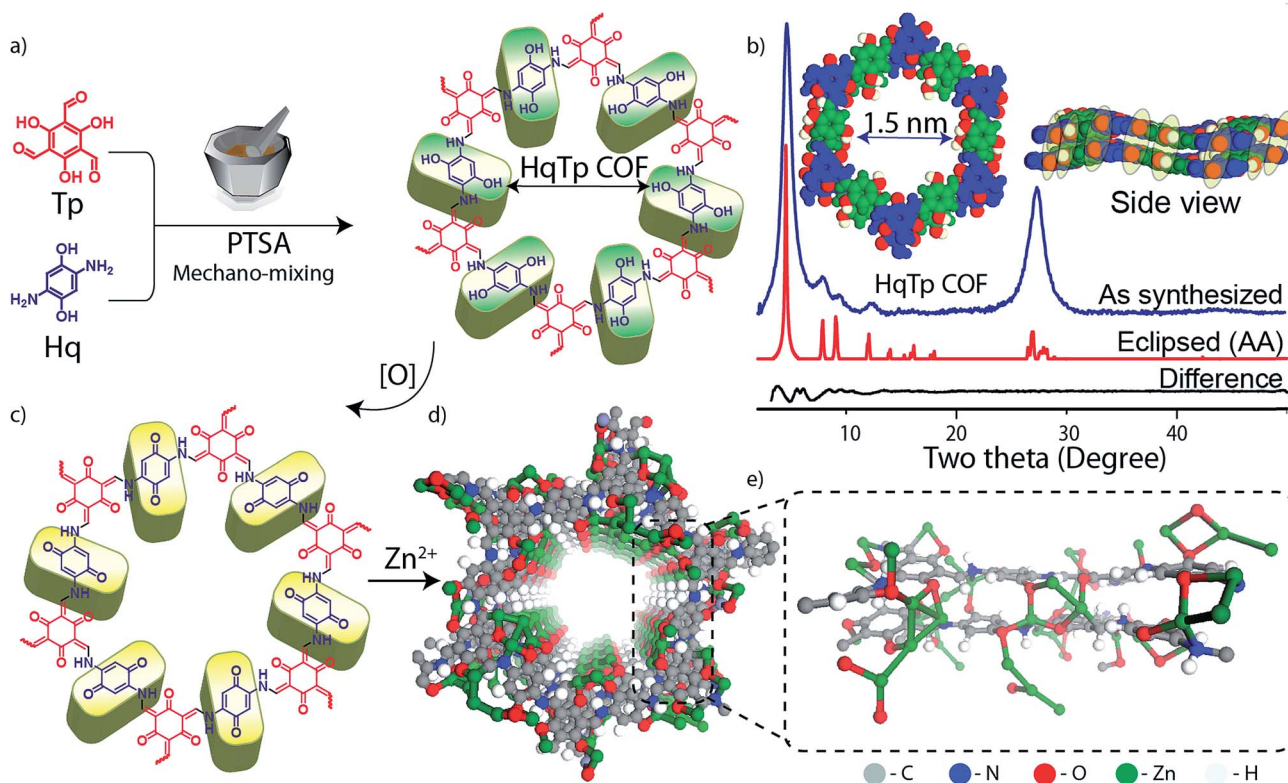


Fig. 1 (a) Schematic representation of the synthesis of **HqTp** (**Hq**-2,5-diaminohydroquinone, **Tp**-1,3,5-triformylphloroglucinol). (b) Powder X-ray diffraction pattern with the AA eclipsed slipped SCC-DFT model. (c) The electrochemical oxidation of hydroquinone to quinone in **HqTp** (d and e). The DFTB model of interlayer interaction of Zn^{2+} cations with the adjacent layers of **HqTp**.

toluenesulfonic acid ($\text{PTSA} \cdot \text{H}_2\text{O}$) as a catalyst (Fig. 1a; ESI, S1†). A black coloured product was obtained after removing the impurities upon wash with water, *N,N*-dimethylacetamide (DMAc) and acetone. The crystalline framework structure and an ordered integration of the building blocks throughout the network is confirmed by the PXRD characterization. The powder X-ray diffraction (PXRD) profile exhibits sharp peaks at the 2θ position of 4.6° , 8.0° , 9.4° , 12.4° and 27.3° (Fig. 1b). The sharp and intense peaks at 4.6 and 27.3° (2θ) correspond to the reflections from 100 and 001 planes, which indicates the existence of a periodic structure. Moreover, the experimental PXRD pattern matches with the simulated PXRD pattern with a slipped eclipsed orientation as modeled using SCC-DFTB (Fig. 1b; ESI S2 and S3†).

The β -ketoenamine framework formation is confirmed by the new peaks in FT-IR spectra at 1583 , 1551 and 1244 cm^{-1} , which correspond to $\text{C}=\text{O}$, $\text{C}=\text{C}$ and $\text{C}-\text{N}$ bonds (all generated after enol–keto tautomerization; ESI; Fig. S6†). Additionally, the ^{13}C solid-state CP-MAS NMR shows enamine and α -enamine carbon peak resonances at 142 and 105 ppm respectively and similarly $\text{C}=\text{O}$ carbon peak at 184 ppm (Fig. 2c; ESI, Fig. S7†). The thermogravimetric analysis (TGA) of **HqTp** displays thermal stability up to 380°C (ESI, Fig. S9†). Brunauer–Emmett–Teller (BET) analysis, using N_2 adsorption at 77 K shows a moderate surface area of $113 \text{ m}^2 \text{ g}^{-1}$ (ESI, Fig. S10†). The non-local density

functional theorem (NLDF) provides a sharp pore size distribution around 1.5 nm (ESI, Fig. S11†).

The presence of $\text{C}=\text{O}$ moieties in **HqTp** COF are principally responsible for the storage of Zn^{2+} ions. These are the C_3 symmetric $\text{C}=\text{O}$ groups resulting from the enol to keto tautomerism and the C_2 symmetric $\text{C}=\text{O}$ group from electrochemically oxidized hydroquinone linker⁸ (Fig. 1a, c, and 3a; ESI, Fig. S8†). The hexagonal 2D lattice enriched with a large number of $\text{C}=\text{O}$ groups, stacked with an interlayer distance of 3.4 \AA , can efficiently host a significant amount of Zn^{2+} in between the two-dimensional layers. Also, the crystalline honeycomb structure of **HqTp** provides a unique pore size of 1.5 nm , which, we believe, further boosts the lucid movement of Zn^{2+} ions through the entire organic cathode without any interruptions.

DFTB calculations were used to probe the nature of the interaction of Zn^{2+} with the COF backbone. They suggest the possibility of the prominent interlayer interactions of Zn^{2+} ions with nucleophilic centers of COF (Fig. 1e, ESI, Fig. S3 and S4†). The $\text{C}=\text{O} \cdots \text{Zn}$ ($2\text{--}2.3 \text{ \AA}$) and $\text{Zn} \cdots \text{N}-\text{H}$ distances ($\sim 2.0 \text{ \AA}$) agree with the possibility of intermolecular interactions. The DFTB models of AA and slipped AA COF $\cdots \text{Zn}^{2+}$ interaction show various structural possibilities including a formation of a Zn^{2+} ion cluster inside the pores. Also, a large enhancement in the per-layer stabilization is observed for the discharged **HqTp** $\cdots \text{Zn}^{2+}$ adduct ($-182.0 \text{ kcal mol}^{-1}$) than the pristine COF





Fig. 2 (a) FT-IR of pristine; charged and discharged cathodes. (b) XPS analysis of pristine; charged and discharged cathodes. (c) ^{13}C CP-MAS NMR of HqTp and the discharged cathode (all the charging and discharge potential are represented vs. Zn/Zn^{2+} reference).

($-40.0 \text{ kcal mol}^{-1}$) which indicates favorable Zn^{2+} ion interaction. To understand the role of electrochemical active sites for Zn^{2+} anchoring in concerned COF, we have performed and compared the electrostatic potential mapping (ESP) of HqTp with benzene linked analog PaTp COF (ESI, Fig. S5 and S30[†]). ESP of the unit cell of HqTp displays a potential range from -0.33 au (red) to 0.061 au (blue). Herein, the negative ESP (red) reflects the affinity of the sites toward the Zn^{2+} uptake. Therefore, from ESP, it is clear that the quinone 'C=O' and Tp 'C=O' in HqTp exhibits more nucleophilicity compared to the 'N-H'

sites. Whereas, the ESP of benzene linked PaTp COF showcases only Tp 'C=O' groups as active sites for the uptake of Zn^{2+} .

We have analyzed the working electrode (cathode) at three different electrochemical conditions to elucidate the probable mechanism of the Zn^{2+} ion interactions within the HqTp COF during the electrochemical charging and discharging process in a zinc ion unit cell; viz: (1) a pristine cathode (an electrode prior to any electrochemical perturbation/cycling); (2) a cathode at a fully discharged state (up to $0.2 \text{ V vs. Zn}/\text{Zn}^{2+}$; where Zn^{2+} do interact with HqTp) and (3) fully charged up to $1.8 \text{ V vs. Zn}/\text{Zn}^{2+}$ (at this stage, Zn^{2+} are withdrawn from the HqTp cathode). Then, we have characterized the electrodes, followed by washing with water.

FT-IR spectroscopy of the pristine organic cathode is found to be similar to the pristine HqTp (Fig. 2a; ESI, Fig. S6b[†]). The C=O group from β -ketoenamine framework appeared at 1584 cm^{-1} . After charging up to $1.8 \text{ V vs. Zn}/\text{Zn}^{2+}$, the hydroquinones are electrochemically oxidized to quinones. This has been confirmed by the observation of a new stretching vibrational peak formed at 1603.0 cm^{-1} . When the system is discharged to $0.2 \text{ V vs. Zn}/\text{Zn}^{2+}$, a simultaneous diminishing of both C=O groups is evident from the FT-IR spectra. Still, a diligent observation shows the presence of a very weak stretching peak at the C=O region of the discharged cathode which indicates that the first cycle of discharging is not sufficient for the full utilization of the C=Os in the framework. Additionally, to figure out the efficiency of the COF based organic cathode, we have strategically performed 500 charge-discharge cycles for a cell. After complete discharging up to $0.2 \text{ V vs. Zn}/\text{Zn}^{2+}$, the cathode was again subjected to FT-IR analysis (ESI, Fig. S6b[†]). The full disappearance of the C=O stretching peaks suggests the prospect of the effective utilization of active groups in HqTp for the interaction with Zn^{2+} .

We have performed ^{13}C CP-MAS NMR spectroscopy of HqTp and discharged ($0.2 \text{ V vs. Zn}/\text{Zn}^{2+}$) HqTp cathode to investigate for any chemical modifications at the atomic level of the framework (Fig. 2c; ESI, Fig. S7[†]). It showcases a down field chemical shift of the C=O peak from 184.0 ppm (for HqTp) to 190.0 ppm (for discharged HqTp). Similarly, the distinct 5–7 ppm downfield chemical shifts have been observed for every peak of the discharged HqTp cathode compared to pristine HqTp COF (ESI, Fig. S7 and S8[†]). It could be due to the appearance of the deshielding effect because of intercalated electropositive Zn^{2+} ions, which decrease the electron density in the framework.⁹

The X-ray photoelectron spectroscopy (XPS) provides information about the chemical state of the concerned elements in the pristine, charged, and discharged conditions of HqTp. In the XPS profile of the discharged HqTp electrode ($0.2 \text{ V vs. Zn}/\text{Zn}^{2+}$), the sharp peaks are visible at the binding energy of 1022.0 and 1045.0 eV , which correspond to $\text{Zn } 2p_{3/2}$ and $\text{Zn } 2p_{1/2}$. Meanwhile, the intensity of the XPS peaks of Zn^{2+} in the charged electrode diminished by half concerning the discharged peak due to the removal of Zn^{2+} ions from the cathode (Fig. 2b; ESI, Fig. S17–S19[†]).

Moreover, the *ex situ* analysis is extended to electronic imaging techniques with elemental mapping. The TEM images



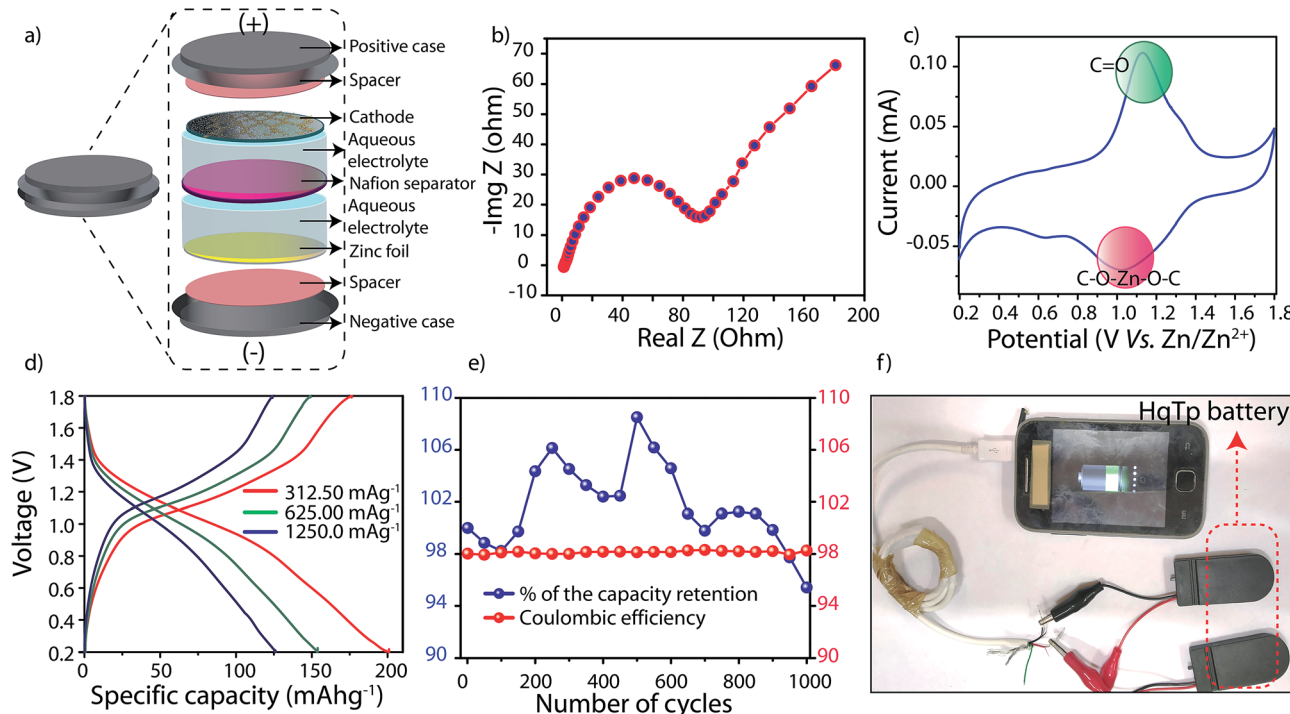


Fig. 4 (a) Diagrammatic representation of the fabrication of lab scale aqueous Zn/HqTp unit cell. Electrochemical characterizations of Zn/HqTp cell. (b) The impedance analysis of Zn/HqTp cell. (c) CV profile of Zn/HqTp cell. (d) Charge–discharge profile of Zn/HqTp cell. (e) Long-life cyclic stability and coulombic efficiency plot at 3750 mA g^{-1} . (f) Digital photograph of charging a smart-phone by Zn/HqTp cell.

materials have been explored for zinc-ion batteries showing charge-storage properties despite the absence of sharp voltage plateaus.¹¹ Similarly, the same trend is observed for COF based Li or Na ion batteries as well.^{2c,e} The lack of a voltage plateau of **HqTp** could be due to the polymeric porous nature of the material and it further indicates a hybrid charge-storage behavior.¹¹ Moreover, it is worth mentioning that **HqTp** exhibits good performance in terms of the obtained discharge capacity as well as cyclic stability as an organic polymer cathode in aqueous zinc ion batteries (Table S4†).^{3,12}

Meanwhile, the **PaTp** COF exhibits only $121.0 \text{ mA h g}^{-1}$ discharge capacity at the same current rate (125 mA g^{-1}) (ESI, Fig. S24†). Considering the discharge capacity of $276.0 \text{ mA h g}^{-1}$, we have found that 7.5 number of Zn^{2+} ions interact in the unit cell of **HqTp** (ESI, S-9†). Moreover, at a current rate of 3750.0 mA g^{-1} where the organic cathode displays the specific capacity of 85.0 mA h g^{-1} , we have carried out a long-term cyclic stability experiment of the **HqTp** zinc ion battery. The cyclic stability exceeded beyond 1000 cycles with 95% retention of its initial capacity (Fig. 3d). Moreover, the coulombic efficiency of the cell is well maintained to 98% throughout the 1000 charge–discharge cycles (Fig. 4e). Although the symmetric shape of reversible peaks are absent in the CV profile, the coulombic efficiency over 1000 cycles indicate the high reversible charge storage characteristics of the **HqTp** COF. Besides, considering the average voltage of 0.87 V (Fig. S27†), a high energy density ($240.0 \text{ W h kg}^{-1}$) has been obtained at the power density of 109.0 W kg^{-1} as provided in the Ragone plot at the energy density of 75.0 W h kg^{-1} , the power density increases

to 3262.0 W kg^{-1} (ESI, Fig. S26†). We could assemble four 1.75 V **HqTp**-zinc ion cells in a series connection and directly used for charging a smart-phone device (Fig. 4f; ESI, Fig. S29†).

Conclusions

In summary, we have demonstrated hydroquinone stitched **HqTp** COF as a new class of organic cathode material for rechargeable aqueous zinc ion batteries. Herein, **HqTp** COF serves as a rich functional platform for binding Zn^{2+} ions. The efficient inter-layer interaction of these divalent Zn^{2+} ions with $\text{C}=\text{O}$ and $\text{N}-\text{H}$ from the adjacent layers provides an excellent discharge capacity ($276.0 \text{ mA h g}^{-1}$ at 125.0 mA g^{-1}). We believe, the utility of COF as an organic cathode in such devices may lead to the further invention of more powerful crystalline polymer organic cathodes in the future.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

A. K. M, V. V, M. G., and A. H acknowledge UGC and CSIR for SRF. M. A. A. thanks the Materials Chemistry Consortium for computational resources on THOMAS (EP/P020194). S. K. acknowledges CSIR for funding through the project TLP003526. R. B. acknowledges IISER-Kolkata start-up research and DST- Swarnajayanti Fellowship grant for funding.



Notes and references

- 1 (a) A. P. Cote, A. I. Benin, N. W. Ockwig, A. J. Matzger, M. O'Keeffe and O. M. Yaghi, *Science*, 2005, **310**, 1166–1170; (b) P. Kuhn, M. Antonietti and A. Thomas, *Angew. Chem., Int. Ed.*, 2008, **47**, 3450–3453; (c) S.-Y. Ding and W. Wang, *Chem. Soc. Rev.*, 2013, **42**, 548–568; (d) S. Wang, Q. Wang, P. Shao, Y. Han, X. Gao, L. Ma, S. Yuan, X. Ma, J. Zhou, X. Feng and B. Wang, *J. Am. Chem. Soc.*, 2017, **139**, 4258–4261; (e) J. W. Crowe, L. A. Baldwin and P. L. McGrier, *J. Am. Chem. Soc.*, 2016, **138**, 10120–10123; (f) A. Sun, B. Aguila, J. Perman, N. Nguyen and S. Ma, *J. Am. Chem. Soc.*, 2016, **138**, 15790–15796; (g) X. Han, Q. Xia, J. Huang, Y. Liu, C. Tan and Y. Cui, *J. Am. Chem. Soc.*, 2017, **139**, 8693–8697; (h) X.-H. Liu, C.-Z. Guan, S.-Y. Ding, W. Wang, H.-J. Yan, D. Wang and L.-J. Wan, *J. Am. Chem. Soc.*, 2013, **135**, 10470–10474; (i) M. Dogru, A. Sonnauer, A. Gavryushin, P. Knochel and T. Bein, *Chem. Commun.*, 2011, **47**, 1707–1709; (j) L. Stegbauer, K. Schwinghammer and B. V. Lotsch, *Chem. Sci.*, 2014, **5**, 2789–2793; (k) E. Q. Jin, M. Asada, Q. Xu, S. Dalapati, M. A. Addicoat, M. A. Brady, H. Xu, T. Nakamura, T. Heine, Q. H. Chen and D. L. Jiang, *Science*, 2017, **357**, 673–676.
- 2 (a) J. Zhou and B. Wang, *Chem. Soc. Rev.*, 2017, **46**, 6927–6945; (b) C. R. DeBlase, K. E. Silberstein, T. T. Truong, H. D. Abruna and W. R. Dichtel, *J. Am. Chem. Soc.*, 2013, **135**, 16821–16824; (c) S. Wang, Q. Wang, P. Shao, Y. Han, X. Gao, L. Ma, S. Yuan, X. Ma, J. Zhou, X. Feng and B. Wang, *J. Am. Chem. Soc.*, 2017, **139**, 4258–4261; (d) M. -S. Kim, W.-J. Lee, S.-M. Paek and J. K. Park, *ACS Appl. Mater. Interfaces*, 2018, **10**, 32102–32111; (e) S. Gu, S. Wu, L. Cao, M. Li, N. Qin, J. Zhu, Z. Wang, Y. Li, Z. Li, J. Chen and Z. Lu, *J. Am. Chem. Soc.*, 2019, **141**(24), 9623–9628.
- 3 (a) G. Fang, J. Zhou, A. Pan and S. Liang, *ACS Energy Lett.*, 2018, **3**, 2480–2501; (b) D. Kundu, B. D. Adams, V. Duffort, S. H. Vajargah and L. F. Nazar, *Nat. Energy*, 2016, **1**, 16119; (c) A. Konarov, N. Voronina, J. H. Jo, Z. Bakenov, Y.-K. Sun and S.-T. Myung, *ACS Energy Lett.*, 2018, **3**, 2620–2640; (d) M. Ghosh, V. Vijayakumar and S. Kurungot, *Energy Technol.*, 2019, 1900442.
- 4 (a) Q. Zhao, W. Huang, Z. Luo, L. Liu, Y. Lu, Y. Li, L. Li, J. Hu, H. Ma and J. Chen, *Sci. Adv.*, 2018, **4**, eaao1761; (b) Z. Guo, Y. Ma, X. Dong, J. Huang, Y. Wang and Y. Xia, *Angew. Chem., Int. Ed.*, 2018, **57**, 11737–11741; (c) D. Kundu, P. Oberholzer, C. Glaros, A. Bouzid, E. Tervoort, A. Pasquarello and M. Niederberger, *Chem. Mater.*, 2018, **30**, 3874–3881; (d) F. Wan, L. Zhang, X. Wang, S. Bi, Z. Niu and J. Chen, *Adv. Funct. Mater.*, 2018, 1804975; (e) L. Ma, S. Chen, H. Li, Z. Ruan, Z. Tang, Z. Liu, Z. Wang, Y. Huang, Z. Pei, J. A. Zapien and C. Zhi, *Energy Environ. Sci.*, 2018, **11**, 2521–2530.
- 5 (a) S. Kandambeth, A. Mallick, B. Lukose, M. V. Mane, T. Heine and R. Banerjee, *J. Am. Chem. Soc.*, 2012, **134**, 19524; (b) S. Chandra, D. R. Chowdhury, M. Addicoat, T. Heine, A. Paul and R. Banerjee, *Chem. Mater.*, 2017, **29**, 2074–2080.
- 6 (a) K. M. Engle, T.-S. Mei, M. Wasa and J.-Q. Yu, *Acc. Chem. Res.*, 2012, **45**, 788–802; (b) S. D. Sarkar, W. Liu, S. I. Kozhushkov and L. Ackermann, *Adv. Synth. Catal.*, 2014, **356**, 1461–1479.
- 7 (a) A. Haaland, *Angew. Chem., Int. Ed.*, 1989, **28**, 992–1007; (b) M. Yamakawa and R. Noyori, *Organometallics*, 1999, **18**, 128–133.
- 8 (a) E. W. L. Chan and M. N. Yousaf, *J. Am. Chem. Soc.*, 2006, **128**, 15542–15546; (b) C. Nicosia and J. Huskens, *Mater. Horiz.*, 2014, **1**, 32–45.
- 9 (a) J. C. Hammel and J. A. S. Smith, *J. Chem. Soc. A*, 1969, 2883–2887; (b) C. A. Wilkie and D. T. Harworth, *J. Inorg. Nucl. Chem.*, 1978, **40**, 195–197.
- 10 (a) M. H. Lee, S. J. Kim, D. Chang, J. Kim, S. Moon, K. Oh, K.-Y. Park, W. M. Seong, H. Park, G. Kwon, B. Lee and K. Kang, *Mater. Today*, 2019, DOI: 10.1016/j.mattod.2019.02.004; (b) L. Suo, O. Borodin, T. Gao, M. Olguin, J. Ho, X. Fan, C. Luo, C. Wang and K. Xu, *Science*, 2015, **350**, 938–943.
- 11 (a) H. Pan, Y. Shao, P. Yan, Y. Cheng, K. S. Han, Z. Nie, C. Wang, J. Yang, X. Li, P. Bhattacharya, K. T. Mueller and J. Liu, *Nat. Energy*, 2016, **1**, 16039; (b) D. Kundu, B. D. Adams, V. Duffort, S. H. Vajargah and L. F. Nazar, *Nat. Energy*, 2016, **1**, 16119–16126; (c) S. Chen, L. Ma, K. Zhang, M. Kamruzzaman, C. Zhia and J. A. Zapien, *J. Mater. Chem. A*, 2019, **7**, 7784–7790; (d) C. Xia, J. Guo, Y. Lei, H. Liang, C. Zhao and H. N. Alshareef, *Adv. Mater.*, 2018, **30**, 1705580; (e) Z. Luo, L. Liu, J. Ning, K. Lei, Y. Lu, F. Li and J. Chen, *Angew. Chem., Int. Ed.*, 2018, **57**, 9443–9446; (f) S. Wang, Q. Wang, P. Shao, Y. Han, X. Gao, L. Ma, S. Yuan, X. Ma, J. Zhou, X. Feng and B. Wang, *J. Am. Chem. Soc.*, 2017, **139**(12), 4258–4261.
- 12 (a) H. Karami, M. F. Mousavi and M. Shamsipur, *J. Power Sources*, 2003, **117**, 255–259; (b) H.-Y. Shi, Y.-J. Ye, K. Liu, Y. Song and X. Sun, *Angew. Chem., Int. Ed.*, 2018, **57**, 16359–16363.

