MOF derived carbon based nanocomposite materials as efficient electrocatalysts for oxygen reduction and oxygen and hydrogen evolution reactions

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The escalating global energy demands and the formidable risks posed by fossil fuels coupled with their rapid depletion have inspired researchers to embark on a quest for sustainable clean energy. Electrochemistry based technologies, e.g., fuel cells, Zn–air batteries or water splitting, are some of the frontrunners of this green energy revolution. The primary concern of such sustainable energy technologies is the efficient conversion and storage of clean energy. Most of these technologies are based on half-cell reactions like oxygen reduction, oxygen and hydrogen evolution reactions, which in turn depend on noble metal based catalysts for their efficient functioning. In order to make such green energy technologies economically viable, the need of the hour is to develop new noble metal free catalysts. Porous carbon, with some assistance from heteroatoms like N or S or earth abundant transition metal or metal oxide nanoparticles, has shown excellent potential in the catalysis of such electrochemical reactions. Metal–organic frameworks (MOFs) containing metal nodes and organic linkers in an ordered morphology with inherent porosity are ideal self-sacrificial templates for such carbon materials. There has been a recent spurt in reports on such MOF-derived carbon based materials as electrocatalysts. In this review, we have presented some of this research work and also discussed the practical reasons behind choosing MOFs for this purpose. Different approaches for synthesizing such carbonaceous materials with unique morphologies and doping, targeted towards superior electrochemical activity, have been documented in this review.

1. Introduction

The rapid progress in technology and industrialization has caused an enormous increase in global energy requirement. To date, the world relies almost entirely on non-renewable fossil

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fuels to cater to the daily energy requirements.\textsuperscript{1} However, the immense adverse effects caused by these fossil fuels on the environment in the form of global warming as well as their swift depletion rates makes it essential for us to regulate their use immediately.\textsuperscript{2–4} Thus the world is in the process of a gradual shift towards a renewable energy based economy, namely solar, wind, and hydro power.\textsuperscript{5–9} However, these sources do not always ensure a steady flow of power and are often sporadic in their outputs. The efficient usage of such sources thus requires effective assistance from conversion and storage technologies such as fuel cells, batteries and capacitors.\textsuperscript{10–19} Fuel cells and metal-air batteries are typical representatives of energy conversion and storage systems with high energy density, high efficiency and no greenhouse emissions.\textsuperscript{20–23} The competence of these devices is primarily determined by the functional materials used as electrocatalysts in their components. Thus the major concern of scientists worldwide is the development of cost and energy efficient sustainable electrocatalysts for better conversion, storage and utilization of renewable energy. While fuel cells involve the combination of a steady stream of O\textsubscript{2} at the cathode and H\textsubscript{2} at the anode to form water, regenerative fuel cells rely on the reverse process of splitting water to generate O\textsubscript{2} and H\textsubscript{2}.\textsuperscript{18,12,14–18} The former is concerned mainly with oxygen reduction reaction (ORR) at the cathode and hydrogen oxidation reaction (HOR) at the anode, and the latter includes hydrogen and oxygen evolution reactions (HER and OER, respectively). Similarly, metal-air batteries perform OER while charging and ORR during discharging cycle.\textsuperscript{20,22–25,30,31,29–33} Many of the above mentioned electrochemical reactions are extremely sluggish in nature, making the use of catalysts mandatory. However, the most efficient catalysts for these processes are often expensive and scarce noble metals like Pt, Ir and Ru.\textsuperscript{45–61} The exorbitant prices of these metals are the biggest obstacles hindering the widespread commercialization of electrochemical process dependant energy technologies. Hence one of the biggest challenges of the materials science world currently involves the development of low-cost, sustainable, energy efficient functional materials that can compete with these noble metals in their activity and thereby replace them. The alternative materials envisaged for this purpose are carbon based materials or benign earth abundant metal based counterparts.\textsuperscript{27–29,57–75} An absolute necessity in these materials is the presence of easily accessible redox-active centres, that will trap the particular element and reduce them promptly. Hence redox active, semiconducting, porous materials like MOFs, COFs or CMPs seem to be the ideal candidates for such catalysis.\textsuperscript{64,124,166,167} Among the cost-effective catalysts explored in recent times, porous carbon materials have emerged as one of the flag bearers owing to their stability, high conductivity, excellent surface area, tunable porosity and surface chemistry.\textsuperscript{64} A few recent reports have also exhibited the efficacy of doping heteroatoms (e.g., N, S and B) in a porous carbon matrix in elevating electrocatalytic activity. These heteroatoms are present in the carbon matrix as defects and brings about electronic and structural modifications (Scheme 1).\textsuperscript{65–75}

One of the most celebrated functional materials of the past decade has been Metal–organic frameworks (MOFs) comprising ordered arrays of metal nodes connecting organic linkers.\textsuperscript{76–79} MOFs have clinched the attention of chemists owing to their light weight, low density, ultra high surface area, porosity and tunable pore surface. Traditionally, MOFs have found application in gas separation, storage and capture of specific molecules.\textsuperscript{80–85} They have also been utilized for catalysis, sensing and for various optoelectronic applications.\textsuperscript{85–90} Of late, MOFs and their derivatives have been explored as electrocatalysts. MOFs, owing to their innate porosity, large carbon content and ordered structures are the ideal templates for porous carbon based materials. Many of the recent research work on electrocatalysis have upheld MOFs and their derived materials as efficient electrocatalysts. In this review, we have documented some of them which stand out and establish that the tailored synthesis of MOFs and their derived materials can provide some of the best materials for electrocatalysis. For this purpose, we have first discussed the design principles of such materials, followed by their activity in common electrochemical processes, e.g., ORR, HER, OER and also bifunctional activity. The systematic arrangement in this review will hopefully be helpful to researchers working in this field and in the future it will help towards the development of supreme MOF-derived carbon based electrocatalysts.

2. Why MOFs as templates for porous carbon based electrocatalysts?

Taking a step away from the conventional applications of MOFs, e.g., storage and separation of gases, drug delivery, magnetism and optoelectronics, recent research has been aimed at designing MOF based electrocatalysts. The easily tunable pore structures and accessible metal nodes are the biggest advantages of using MOFs as electrocatalysts. The permanent porosity of MOFs helps in the swift diffusion of substrates throughout the material, thus helping the substrates reach catalytic sites.
comfortably. By careful selection of metal and linkers and altering the stoichiometry, one can achieve structures that have supreme electrochemical activity. Frameworks containing earth abundant redox-active first row transition metals (e.g., Fe, Co, Ni, Mn, etc.) are particularly suitable for the job. Upon being coupled with a redox active linker, MOFs can show excellent electrochemical activity.\(^\text{168}\) Moreover the isolated metal nodes in MOFs enclosing nanoconfinies are ideal for molecular catalysis. In spite of several advantages, the biggest drawback in using MOFs as electrocatalysts is their poor conductivity that limits the electrochemical activity.\(^\text{91}\) The most common way to counter this is to introduce catalytic nanoparticles or conductive guest species in the pores of the MOF, e.g., TCNQ or TTF.\(^\text{92}\) Sometimes the formation of composites with highly conductive materials like graphene is also advantageous.\(^\text{169}\) A composite of a Zr-MOF, NU-1000 with Ni-S films has shown excellent activity as a catalyst for HER.\(^\text{93}\) Growing MOFs as thin films is another strategy for generating productive electrocatalysts, as the control over film thickness is crucial to facilitate electron transport through the active catalytic sites. Yang \textit{et al.} used thin films of Coporphyrin nanoMOF for the electrochemical reduction of CO\(_2\).\(^\text{94}\) However, the most effective strategy to fabricate electrocatalysts from MOFs is the synthesis of MOF derived carbonaceous materials.

Porous carbon when doped with heteroatoms, e.g., N, S or B, have been reported to show enhanced electrocatalytic activities, comparable to some of the noble metal-based benchmark catalysts. Also, non-precious metal and metal oxide nanoparticles (NPs) supported on carbon matrix (MNPC) with high porosity offer numerous exposed and easily accessible active sites, thereby improving overall electrocatalytic activity.\(^\text{67,68,70,75,95,96}\) However, the traditional methods for synthesis of such porous carbon materials involves simple pyrolysis followed by physical or chemical activation of organic precursors, which often gives rise to highly disordered structures with broad pore size distribution. This paved the way for nanocasting or templating approaches, initiated using soft templates, e.g., resols and hard templates, e.g., zeolites.\(^\text{97,98}\) These were again found to be unsuitable for bulk scale production. In an innovative approach, Xu \textit{et al.} first proposed MOFs as self sacrificial templates for effective synthesis of porous carbon.\(^\text{99}\) The inherent porosity of MOF, high surface area, large carbon content and ordered structure makes them ideal candidates as templates for carbon.

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Scheme 1  Development of electrocatalysts from MOF derived carbon materials.
materials. They also ensure facile heteroatom and metal doping in the carbon matrix, thereby producing novel materials with enhanced properties. In many cases, precursors like furfuryl alcohol, sucrose, glycerol, phenolic resin, etc. are introduced in the MOF templates as a carbon source. However, it is also possible to make such materials without the use of any external precursor and the original MOF linker is treated as a source for carbon as well as heteroatoms like N or S. Because of their electron rich surfaces, such heteroatom doped carbon materials have the potential to manifest excellent electrocatalytic activity. In addition, if the original electroactive metal atom of the MOF is retained in the form of metal or metal oxide nanoparticles, the electrochemical activity can be further enhanced.

To put it in layman’s terms, electronic properties of a particular material dictates its efficiency as an electrocatalyst. A pure graphitic carbon matrix consisting of sp2 bonded carbon atoms with an uniformly delocalized electron cloud lacks polarization and consequently cannot afford to accommodate the existing electronic cloud of intermediates. Heteroatoms, e.g., N, B, P or S, with different electronegativities when introduced into this graphitic matrix, a redistribution of the electronic cloud takes place, thereby creating a polarization which can now accommodate the foreign invasion of intermediates. The most commonly doped heteroatom is nitrogen which is more electronegative and hence these doping sites are potent for metal functional group coordination and act as active sites for the electrocatalysis.

The heteroatom doped carbon matrix have the potential to act as efficient metal free electrocatalysts. One of the primary reasons to choose MOFs as templates for carbonization is the ease of including heteroatoms in the carbon matrix, by simply selecting a MOF with a heteroatom containing linker. For example, MOFs with imidazolate framework are often pyrolyzed at inert atmosphere to get N-doped carbon matrix with low porosity. Other N-containing linkers, e.g., pyridine derivatives may also be used. We showed the formation for Co doped carbon nanotubes (Co-CNTs) by the pyrolysis of dicyanamide based nitrogen rich MOF \( \text{[Co(bpe)}_2(N(CN))_2\{N(CN)_2\}]/\text{Co-MOF-1, bpe}=1,2\text{-bis(4-pyridyl)ethane, } N(CN)_2^- \text{ = dicyanamide}. \)

However, in most cases, scientists opt for external N-sources like urea, melamine, etc. to generate N containing C materials. Kurungot et al. performed the pyrolysis MOF5 in the presence of nitrogen rich melamine to generate g-C_3N_4 for electrocatalytic ORR (Fig. 1(a)). Guest molecules impregnated within the MOF pores may also be used as a source of heteroatom. In an interesting piece of work, we have utilized a guest dimethyl amine cation of an anionic MOF as a pre-bonded C–N–C moiety to generate N-doped C-dots. Also, co-doping two or more elements with reverse electronegativity results in greater number of defects, generating more active sites for an elevated electrocatalytic activity. Thus, compared to single heteroatom doping, multiple heteroatom doping could be beneficial. For this, either the ligand must be selected such that it has the choice of heteroatoms or the MOF can be impregnated with heteroatom containing moieties.

It has already been established that metals/metal oxides and their nanoparticles (NPs) play an integral role in the electrocatalysis. The more recent literature presents that metal/metal oxide NPs encapsulated inside carbon matrix are also effective in improving the electrocatalytic activity. MOFs, containing redox-active metal nodes and carbon rich organic ligands thus become the automatic choice of sacrificial templates for preparing metal NPs embedded in a carbon matrix. Introduction of more than one transition metal in the matrix has also been proven beneficial in enhancing the electrocatalytic activity. Post-synthetic metal exchanged MOF, metal cores coated with MOF, bimetallic MOFs are some of the possible sacrificial precursor for the preparation of multiple transition metal electrocatalysts. Depending upon the carbonization atmosphere, encapsulations and type of post treatments, metal carbides, nitrides, oxides, sulphides or phosphides are formed. Shim et al. pyrolysed a 1D coordination polymer synthesized from Zn and dithiooxamide linker doped with hemoglobin to generate Fe, S and N doped carbon matrix (Fe-SNC) which showed excellent activity as a supercapacitor and electrocatalyst for ORR (Fig. 1(c)).

Morphology of the MOF or the derived material also plays a crucial role in determining its electrochemical activity. The porosity of the composite depicts the active site accessibility and consequently the efficacy. However, high temperature annealing sometimes causes the framework to shrink and lose porosity, surface area and active site accessibility. Although this may be addressed by controlling the annealing temperature, however there is no clear solution to counter it and thus morphology control still remains a challenge. An effective and desirable electrocatalyst should have highly accessible active catalytic sites, facile electron transfer all over the matrix in order to get good conductivity and usage of earth abundant transition metals to curtail cost. A cautious balance of these features is the key to generate effective electrocatalysts for sustained use. In summary, MOFs are one of the best suited templates for the generation of such porous carbon based matrices for electrochemical applications because of the following reasons:

(a) The nanocasting approach of using MOF templates is extremely effective because of the ordered structure and innate porosity. Thus MOFs ensure facile encapsulation of carbon or heteroatom containing guest species.

(b) The various nanoscale morphologies of MOFs ensure similar morphologies in their carbon counterparts, exposing active sites.

(c) MOFs can be efficiently grown over conductive surfaces or substrates as nanomorphologies or thin films, ensuring better activity.

In the following paragraphs, few electrocatalysts prepared using the above strategies are discussed categorically as ORR, OER, HER and bifunctional catalysts.
3. MOF derived carbon nanocomposites in ORR, OER and HER

3.1 MOF derived materials for oxygen reduction reaction (ORR)

Reduction of oxygen is not only one of the most important reactions in animal physiology, but also important industrially, particularly in fuel cells and metal–air batteries. In this process, oxygen may either gain two electrons to form H$_2$O$_2$ or four electrons to form H$_2$O. The kinetics of this reaction is extremely sluggish in nature and requires catalysts for its progress. Till date, the best known catalyst for this has been platinum, which because of its scarcity and immense expenses, has kept all attempts of commercializing ORR based technologies at bay. Thus over the past few decades, one of the major topics of research in materials science has been the incessant quest for cheaper, sustainable, environmentally benign catalyst for ORR. Some of the nanoporous carbons have shown tremendous potential as electrocatalysts for ORR. As mentioned previously, heteroatom doping plays an important role in creating defects in the carbon matrix which in turn reflects in the improvement of the catalytic activity.

The most common approach to prepare N-doped MOF derived electrocatalyst is to select a MOF with a N-rich ligand, viz., imidazole (IM), in particular the ZIF series of MOFs.
et al. showed that direct pyrolysis of ZIF-67 (Co(MIM)$_2$, MIM = methylimidazole) produced a highly N-doped carbonaceous material with Co impregnation.$^{156}$ Metal free, N-doped porous carbon material was prepared by Cao and group where they treated ZIF-7 (Zn[PhIM]$_2$·(H$_2$O)$_3$; PhIM = benzimidazole) with glucose as an external excess carbon source.$^{140}$ The carbonized material was tested as an electrocatalyst for ORR and it showed an onset potential of 0.7 V vs. RHE, selective H$_2$O production without any methanol crossover effect.

A series of bimetallic ZIFs (BMZIFs) with varied Zn/Co ratio were also studied for preparation of porous carbon materials where N was doped uniformly by Jiang and group (Fig. 2(a)).$^{141}$ Zn got evaporated leaving Co decorated N-doped carbon matrix (CNCo-$_n$) ($n$ indicating the Zn/Co molar ratio in the precursor BMZIF). Later CNCo-$_n$ was doped with P by annealing it with triphenylphosphine under N$_2$ atmosphere (Fig. 2(b–d)). CNCo-20 when studied as an ORR electrocatalyst showed an onset potential of $-0.08$ vs. Ag/AgCl, on P-doping (P–CNCo-20) the onset potential shifted towards more positive potential ($-0.04$ V vs. Ag/AgCl) (Fig. 2(e–g)). These catalysts also showed long term stability and good methanol tolerance. Tang and co-workers post-synthetically treated a Co-MOF (ZIF-67) with NiCl$_2$ solution to get Ni/Co-MOF which on pyrolysis transformed into a highly porous carbon matrix with uniformly distributed Co, Ni and N centres, ensuring a high density of active sites with superior ORR activity in electrolytic solution of neutral to higher pH.$^{142}$

Carbonization of self-adjusted Zn and Co bimetal–organic frameworks (bi-MOF) [Zn$_{1-x}$Co$_x$(MeIM)$_2$; MeIM = methyl imidazole] resulted in Co–N–C$_x$ composites. Among these materials, Co–N–C-0.8 was found to be the most active toward the ORR due to the presence of pyridinic N (47%), pyrrolic N (43%), and oxidized N (10%) in the carbon matrix.$^{143}$

In an attempt to understand the effect of annealing temperature on the morphology of MOF derived materials, recently our group documented synthesis of nanoporous carbon and nitrogen doped carbon dots from Mg(II) based anionic MOF (Mg-MOF) {[Mg$_3$(ndc)$_2$·(HCO$_2$)$_2$(H$_2$O)]·[NH$_2$Me$_2$]·2H$_2$O·DMF} (ndc = naphthalene dicarboxylate) by varying the carbonizing temperature.$^{129}$ The nanoporous carbon (NPC) obtained by carbonizing Mg-MOF at 900 °C and subsequently washing with a concentrated acid mixture (HCl : HNO$_3$ in 1 : 1 v/v). N-doped carbon dots (NCD) were obtained at 500 °C treating in similar conditions (Fig. 3(a)). Both NPC and NCD were characterized by IR, Raman, XPS, XRD and several microscopic techniques. NPC showed BET surface area of 3690 m$^2$ g$^{-1}$, which is one of the highest surface areas of a porous

![Fig. 2](image_url) (a) Schematic representation of the imidazole based frameworks used for carbonization. SEM images of (b) BMZIF-20 and its pyrolysed counterpart (c) CNCo-20. (d) TEM image showing Co nanoparticles embedded in layers of graphitized carbon. (e) CV of P–CNCo-20 in 0.1 M KOH in Ar and O$_2$ saturated conditions. (f) LSV curves for the different carbonized samples in 0.1 M KOH at 1600 rpm. (g) LSV curves at different rotation speeds for P–CNCo-20 in 0.1 M KOH. Reproduced with permission from ref. 141. Copyright 2015 Wiley.
carbon obtained from directly carbonizing a MOF. The presence of DMA (NH₂Me₂) containing prebonded C–N moieties in the pore of the anionic Mg-MOF helped in the synthesis of N-doped carbon dots. The diameter of each dot is well below 10 nm and HRTEM showed graphitic lattice fringes with an interplanar distance of 0.34 nm (Fig. 3(b)). Surface area of NCD is about 95 m² g⁻¹. NCD has structural and compositional features similar to g-C₃N₄ which is known to show good catalytic activity for ORR. Linear sweep voltammetry (LSV) at different rotation speeds was performed and NCD showed moderate catalytic activity for ORR (Fig. 3(c)). To further improve the catalytic activity, Co atoms were doped in the NCD. This was achieved by mixing Co–porphyrin with NCD and the mixture was pyrolyzed at 800 °C and a new composite was obtained. PXRD, XPS and TEM analysis suggest the presence of metallic Co-nanoparticle on the N-doped graphitic carbon (Fig. 3(e)). XRD also suggested the tris-triazine units in carbon nitride, remain intact after Co-doping. TEM study showed that metallic Co-nanoparticles (less than 5 nm) embedded in a carbon matrix and lattice spacing of 0.24 nm correspond to fcc Co. ICP analysis suggested about 1.5 wt% of Co in the Co@CN nano-composite material. ORR activity was studied using Co@CN sample and a significant difference was observed with respect to NCD. LSV of Co@CN showed significantly enhanced cathodic current as a function of rotation speed and affording a current density of −1 mA cm⁻² at low overpotentials (−0.8 V) as compared to 0.36 V for NCD (Fig. 3(f)). K–L plot revealed that linearity and the number of electrons transferred during the ORR is 3.87, suggesting 4 electron transfer pathway to produce H₂O from O₂ reduction. Further, rotating ring disk electrode (RRDE) measurement showed that H₂O₂ production is decreased from 40.8% for NCD to 2.3% for Co@CN (Fig. 3(d)). The Tafel slopes for Co@CN were 63 mV per decade and 141 mV per decade at lower and higher current density respectively, suggesting faster reaction kinetics. Chronopotentiometric (CP) electrolysis study were performed for both NCD and Co@CN at a constant voltage for 48 h at a rotation speed of 1600 rpm (Fig. 3(g)). Both the catalysts showed excellent stability in alkaline solution during ORR. Overall, these results demonstrated that NCD catalyze the reduction of O₂ to H₂O₂ and

Fig. 3  (a) Schematic representation of the pyrolysis of 1 at 500 °C to generate N-doped C-dots and further Co doping in it. (b) TEM image of the N-doped C-dots formed by carbonizing 1. (c) LSV curves for ORR by NCD-1. (d) RRDE voltammograms collected at 5 mV s⁻¹ and 1600 rpm showing ORR in 0.1 M KOH. (e) TEM image of Co@CN. (f) LSV curve for ORR in 0.1 M KOH for Co@CN. (g) Chronoamperometric response for NCD-1 and Co@CN. Reproduced from ref. 129. Copyright 2017 Royal Society of Chemistry.
metallic Co-nanoparticles catalyze the decomposition of the formed \( \text{H}_2\text{O}_2 \). Such enhanced catalytic activity in Co@CN can be attributed to the electronic interaction between the metal nanoparticles and graphitic carbon, which accelerates the kinetics of the ORR.

A similar attempt by Liang et al., yielded N and S doped mesoporous carbon by annealing an inherently N and S containing MOF \( [\text{Zn}_2(\text{TDC})_2(\text{DABCO})] \cdot 4\text{DMF} \) (SCUT-12, TDC = 2,5-thiophenedicarboxylic acid, DABCO = 1,4-diazabicyclo[2.2.2]octane) without the use of any external dopant. Owing to different pyrolysis temperatures, the S content and type varied, giving different ORR activities. Overall the material showed excellent ORR activity comparable to Pt/C catalyst.\(^{144}\)

Zou and co-workers studied the effect of particle size of nano MOF and the corresponding derived material on the catalytic activity.\(^{145}\) They synthesized ZIF-67 (Fig. 4(a)) in various conditions to achieve different particle sizes ranging from bulk to 300 nm. The polyhedral morphology of the MOF nanoparticles were retained in the MOF derived carbon materials (MDCs) by carefully choosing the pyrolyzing temperature under Ar atmosphere, which is ascertained from TEM and SEM studies (Fig. 4(b and c)). TEM also proves the presence of Co in the carbon matrix (Fig. 4(c)). The effect of size of the particles directly affects the electrocatalytic activity, as observed by recording the LSV curves for the different samples (Fig. 4(d)). The LSV shows a lesser overpotential for onset of ORR (0.86 V vs. RHE) for the lowest size MDC (300 nm) than for the MDC derived from bulk MOF (0.80 V vs. RHE). The electron transfer pathway was also affected by the particle size, as the 300 nm MDC shows 4-electron transfer with the highest stability (Fig. 4(e)). This study establishes that with smaller particle size, the active sites are much more exposed and easily accessible which promotes faster mass and electron transfer.

The effect of morphology on electrochemical activity was presented in an intriguing work by Yu et al.,\(^{146}\) where tellurium nanowires (TeNWs) were used as templates for growing ZIF-8 nanofibers (Fig. 5(a and b)). Carbonization of the resultant high aspect ratio ZIF-8 nanofibers at 1000 °C yields highly porous N-doped carbon nanofibers (Z8-Te1000) which retain the hollow fibrous morphology with uniform diameter of 50 nm (Fig. 5(c)). The electrocatalytic activity of Z8-Te1000 was primarily shown in the CVs recorded in \( \text{O}_2 \) and \( \text{N}_2 \) saturated 0.1 M KOH solution separately (Fig. 5(d)). The Z8-Te1000 showed good catalytic activity towards ORR which is apparent from the LSVs recorded at different rotation speeds and the corresponding K–L plot (Fig. 5(e,f)). To further increase the catalytic activity reannealing with triphenylphosphine was done which resulted in P-doping (P-Z8-Te1000). P-Z8-Te1000 shows excellent activity as an electrocatalyst for ORR, even superior to the benchmark Pt/C catalyst (Fig. 5(g)). However, the carbon material obtained by the direct pyrolysis of bulk ZIF-8 does not show good electrocatalytic activity (Fig. 5(g)).

A remarkable piece of work by Yu et al., utilizes the bifunctionality observed in a MOF-derived porous carbon for direct use in \( \text{Zn}–\text{Air} \) and \( \text{Li}–\text{S} \) batteries. In this case, a dual templating approach was followed where initially an ordered silica photonic crystal template was produced from 200 nm silica particles, followed by the infiltration by ZIF-8 precursor solution (Fig. 6(a)). Complete solvent evaporation yields uniform ZIF-8 crystals on the silica microsphere surface and in the interstitial voids (Fig. 6(b)). Next pyrolysis at high temperature of (800–1100 °C) converted the composite into carbon morphologies supported on a silica matrix (Fig. 6(c)). A macroporous carbon texture with interconnected narrow walls between the special voids was obtained upon removing the silica matrix (Fig. 6(d)). The sample carbonized at 950 °C (BHPC-950) has a high total pore volume of 13.42 cm\(^3\) g\(^{-1}\), large specific surface area of 2546 m\(^2\) g\(^{-1}\) and a high nitrogen dopant content of 7.6 atom%.

BHPC-950 shows an excellent ORR activity with an onset potential of 0.93 V and a limiting current density of 5.34 mA cm\(^{-2}\) and a 4-electron pathway (Fig. 6(e)). Not only does it surpass the benchmark Pt/C catalyst in its activity, but also it shows a very small Tafel slope of 74 mV dec\(^{-1}\), signifying a large number of exposed active sites and swift ORR kinetics. However, a sample prepared by the direct carbonization of ZIF-8 nanocrystals under the same conditions without macropores does not show appreciable ORR activity. Since both the samples have equivalent N content, the reduced activity can only be attributed to the low specific surface area, due to the lack of macropores, less number of exposed sites and high mass-transfer barrier. BHPC-950 was assembled by loading on carbon cloth as an air cathode along with a Zn-foil as the anode in a \( \text{Zn}–\text{air} \) battery. BHPC-950 shows a high discharge current density of 319 mA cm\(^{-2}\) and peak power density of 197 mW cm\(^{-2}\) at catalyst loading of merely 0.5 mg cm\(^{-2}\) (Fig. 6(f)). The BHPC-950 battery shows a specific capacity of 797 mA h g\(_{\text{Zn}}^{-1}\) corresponding to an energy density of 963 W h kg\(_{\text{Zn}}^{-1}\) at 20 mA cm\(^{-2}\), which corresponds to 95% utilization of theoretical capacity. The outstanding stability of BHPC-950 electrode is exhibited by the continuous lighting of a LED pattern comprising 30 2.2 V LED bulbs by two such Zn–air batteries in series for more than 12 h without brightness decay (Fig. 6(g)).\(^{147}\)

Several other simplistic approaches have been undertaken to synthesize economically viable electrocatalysts for ORR with supreme activity. In fact, among the three electrochemical reactions mentioned here, the most number of MOF-derived electrocatalysts have been developed for ORR, upholding a new vista of possibilities.

### 3.2 MOF derived electrocatalysts for oxygen evolution reaction (OER)

Electrochemical water splitting has been highly regarded as one of the most important reactions in modern times in order to generate \( \text{H}_2 \) for storing energy in the form of chemical bonds. One half of the water splitting reaction involves the evolution of oxygen from water also regarded as oxygen evolution reaction (OER). The reaction involved is as follows:

\[
\text{H}_2\text{O} + 2(\text{h}^+) \rightarrow \frac{1}{2}\text{O}_2 + 2\text{H}^+ 
\]

The efficiency of this reaction is not only essential for the overall water splitting reaction, but this is also an important
anode reaction in fuel cells and metal–air batteries. However, similar to ORR, OER is also extensively dependent on expensive noble metal based catalysts, e.g., IrO$_2$ or RuO$_2$. Hence for the overall development of electrochemical technologies, it is important to develop cheaper alternatives for catalysis of OER.

In an interesting work, Lee and group used a layered Co–Mo-MOF (Fig. 7(a)) as a template along with polyvinylpyrrolidone (PVP) as a structure directing agent as well as a N source for generating N-doped carbon matrix. Here, the PVP is expected to lower the surface energy of the ultrathin Co–Mo-MOF layers and contribute in the formation of integrated 2D and 3D structures. Field effect scanning electron microscopic (FESEM) images show formation of hierarchical microstructure composed of the hybrid MOF (Co–Mo-MOF) nanowires (Fig. 7(b)). On pyrolyzing Co–Mo-MOF at 600 °C, CoO$_x$–MoC/NC were formed in the form of 3D nanowire assemblies, similar to the template. The FESEM images of the pyrolyzed material shows 3D micro-ribbon structures with evenly distributed metal nanoparticles (Fig. 7(c and d)). To realize an optimum Co/Mo ratio, MOFs with varied Co/Mo ratio viz. 35.8 : 1, 21 : 1, 10.3 : 1 was prepared and the corresponding pyrolyzed products formed are CoO$_x$–MoC/NC-1, CoO$_x$–MoC/NC-2 and CoO$_x$–MoC/NC-3, respectively. The LSV of all the CoO$_x$–MoC/NC reveals an optimum electrocatalytic activity of CoO$_x$–MoC/NC-1 with
lowest onset potential of 1.52 V vs. RHE and lowest Tafel slope of 89.8 mV per decade (Fig. 7(e and f)). It is evident from this that while a high Co/Mo ratio does not ensure a high conductivity, a lower Co/Mo ratio does not allow enough active sites for an enhanced catalytic activity. Hence, it is essential to have an optimum bimetallic ratio for efficient electrocatalysis of OER.

The high catalytic activity of CoO$_x$–MoC/NC-2 than the other CoO$_x$–MoC/NCs was probably due to higher specific surface area, which was confirmed by calculating electrochemical surface areas (ECSAs) from the double-layer capacitance. This was found to be highest for CoO$_x$–MoC/NC-2, which also showed a very high stability.

Li and group effectively utilized a nickel foam to grow MOF and proceeded on to carbonize it. First, they electrodeposited polyaniline (PAni) on nickel foam (NF) (NF@PAni). The layer of PAni promptly traps transition metal ions and upon exposure to H$_2$dodc (2,5-dioxido-1,4-benzenedicarboxylic acid), the MOF-74-M [M$_x$(dobdc)](M = Fe, Co, H$_2$dodc = 2,5-dioxido-1,4-benzenedicarboxylic acid) grows on NF@PAni (NF@PAni@Co/Fe$_2$O$_4$-NRAs). This layered substrate was eventually pyrolyzed at 600 °C (Fig. 8). MOF-74-M was selected for the choice of metal (Fe, Co) and a high O/C ratio of the ligand. SEM images reveal the growth of MOF-74-Co/Fe nanorods (diameter ~ 680 nm) array (NRA) on NF@PAni (Fig. 9(a)). The subsequent heat treatment under N$_2$ atmosphere yielded Co/Fe$_2$O$_4$–C, denoted as NF@NC@Co/Fe$_2$O$_4$–C NRAs. Fig. 9(b) shows the SEM image of Co/Fe$_2$O$_4$–C NRAs with the inset showing a magnified nanorod. The TEM images of Co/Fe$_2$O$_4$–C ascertains the impregnation of metal NPs on a porous carbon matrix advantageous for catalytic activity (Fig. 9(c and d)).

Four types of electrodes were fabricated to show the advantages gained by growing NRAs on NF@PAni, which are as follows: (i) CoFe$_2$O$_4$ on NF@NC (NF@NC-CoFe$_2$O$_4$), (ii) CoFe$_2$O$_4$/C on NF@NC(NF@NC-CoFe$_2$O$_4$/C), (iii) commercial IrO$_2$ on NF@NC (NF@NC-IrO$_2$) on NF@NC, (iv) NF@NC@CoFe$_2$O$_4$/C NRAs. Upon performing the LSV, true to the conjecture of the authors, NF@NC@CoFe$_2$O$_4$/C NRAs showed the lowest onset potential (~1.45 vs. RHE) (Fig. 9(e)) and Tafel slope was found to be ~45 mV dec$^{-1}$ (Fig. 9(f)), indicative of an extremely fast kinetics. The nanorod
morphology of NF@NC@CoFe$_2$O$_4$/C NRAs plays an important role in accelerating the kinetics which in turn is the outcome of lower series resistance ($R_s$) and charge transfer resistance ($R_{CT}$) (Fig. 9(g)). NF@NC@Co/Fe$_2$O$_4$–C NRAs also showed a stable chronopotentiometric response for 30 h (Fig. 9(h)). Polarization curves recorded before and after the durability test show very negligible change in the LSV profile, concluding the longevity of NF@NC-CoFe$_2$O$_4$/C NRAs (Fig. 9(i)).

In a recent work, a bimetallic ZIF-8 comprising both Ni and Zn has been carbonized between 700–1100 °C to obtain Ni induced tubular graphitic nanostructures (ZNDC). ZNDC-1100 shows excellent activity as a catalyst for HER, achieving benchmark 10 mA cm$^{-2}$ current density at a low overpotential of 1.67 V, which is better than IrO$_2$. Such attempts, although few, delineated the potential of MOFs in fabricating successful OER electrocatalysts from MOF derived carbon materials.

### 3.3 MOF derived electrocatalysts for hydrogen evolution reaction (HER)

Hydrogen Evolution Reaction (HER) is an important half-cell reaction in water splitting. Hydrogen being the most important next-generation clean fuel, efficient HER can lead to the sustainable generation of hydrogen that can be used in fuel cells. It is a classic example of a two-electron process and the reaction occurring is as follows:

$$2H^+ + 2e^- \rightarrow H_2$$

The benchmark catalyst in this case happens to be Pt. However in recent times MOF derived carbon materials have also been studied as electrocatalysts for this reaction.

The pyrolyzed derivatives of well-known MOFs, e.g., HKUST-1 (ref. 157) were initially explored for electrocatalysis without further modification (Fig. 10(a and b)). The octahedral shape of HKUST-1 (Fig. 10(c)) upon carbonization changed and adopted a distorted spherical “cauliflower-like” morphology (Fig. 10(d)), with particle sizes of ~20 nm in its carbonized form regarded as Cu/NPC. EDS showed the presence of only C and Cu in Cu/NPC and the BET surface area was calculated to be ~1025 m$^2$ g$^{-1}$. Upon being coated on GCE, Cu/NPC showed a decent activity as an electrocatalyst for HER in acidic medium producing a current density of ~10.66 mA cm$^{-2}$ at a potential of ~1.0 V (vs. RHE). As expected, its activity was much better than that in the
original HKUST-1. This has been attributed to the metallic Cu,
CuO and Cu$_2$O present in Cu/NPC, unlike the strong covalently
bonded Cu in the HKUST framework. Moreover the high BET
surface area as well H$_2$ adsorption capacity provides an added
advantage to Cu/NPC. Although the resistance to Faradaic
process is greater in Cu/NPC than in HKUST-1 owing to the
presence of Cu$_2$O, the overall catalytic activity is much better in
Cu/NPC.$^{158}$

In an extension to the afore-mentioned work, a novel catalyst
containing Cu–Pd bimetallic nanoparticles supported on a
nanoporous carbon was used for HER. Here, HKUST-1 was
prepared in both hydrothermal and electrochemical methods.
Thereby, they were carbonized and some of the Cu replaced by
Pd using galvanic replacement, also known as trans-metallation.
The composite material yielded from the electrochemically
synthesized HKUST-1 (Cu–Pd/NPCC/EC (Fig. 10(e))) showed
a much higher current density and lower overpotential than its
hydrothermally synthesized counterpart (Fig. 10(f)). This has
been attributed to the high specific surface area of Cu–Pd/
NPCC/EC. Moreover, the electrochemical synthesis controls
the oxidation state of Cu by the application of a proper potential
and also enhances the Cu content.$^{151}$

Xu et al., presented a very simple and effective method of
preparing HER electrocatalysts active in both acid and alkaline
medium by carbonizing ZIF-67 at 600 °C under N$_2$ atmosphere
to form Co/N–C.$^{152}$ A vacuum phosphosulfurization process in
the presence of P and S powders led to the formation of CoPS/N–
C (Fig. 11(a)). Similar methods were used to prepare CoP$_2$/N–C
and CoS2/N–C nanocomposites for a comparative study. The ZIF-67 and all its derived nanocomposites were characterized by XRD, Raman spectroscopy, X-ray photoelectron spectroscopy which strongly attest the formation of CoPS/N–C with CoPS supported on N-doped carbon matrix. The polyhedral morphology of ZIF-67 was inherited by the CoPS/N–C with signs of structural collapse explained by the decrease in particle size caused due to the probable reconstruction of carbon framework during high temperature carbonization, phosphorization, sulfurization and phosphosulfurization processes (Fig. 11(b–d)). This shrinkage of carbon framework was also evident from the calculated BET surface areas (ZIF-67 = 1493 m\(^2\) g\(^{-1}\), Co/N–C = 317 m\(^2\) g\(^{-1}\) and CoPS/N–C = 69 m\(^2\) g\(^{-1}\)). Co, P, S, C, N and O in CoPS/N–C were distributed uniformly on the polyhedrons as evident from the microscopic images and elemental mapping (Fig. 11(e)). The electrocatalytic activities of ZIF-67, Co/N–C, CoPS/N–C and CoS2/N–C nanocomposites, towards HER were tested by casting them on a GCE and corresponding LSVs were tested in 0.5 M H\(_2\)SO\(_4\) at a rotating speed of 2000 rpm. Pristine ZIF-67 was inactive whereas the nanocomposites showed a steep rise in the cathodic current, among which CoPS/N–C outperformed the CoP2/N–C, CoS2/N–C and Co/N–C. CoPS/N–C required an overpotential of ~80 mV vs. RHE to achieve a current density of ~10 mA cm\(^{-2}\) (Fig. 11(f)). The Tafel slope analysis proposed a two-electron transfer process following Volmer–Heyrovsky mechanism of bimolecular adsorption and hydrogen evolution (Fig. 11(g)). Also, the low value of the slope (69 mV dec\(^{-1}\)) supports the efficient reaction kinetics. To find out the reason behind this excellent catalytic activity, double-layer capacitance (C\(_{dl}\)) and charge transfer resistance (R\(_{CT}\)) were measured by CV and electrochemical impedance spectroscopy (EIS), respectively. The C\(_{dl}\) for CoPS/N–C was found to be the highest (21.55 mF cm\(^{-2}\)) and R\(_{CT}\) was found to be the lowest (4.8 Ω) among the other nanocomposites, signifying larger surface area, more accessible active sites and faster Faradaic response and hence faster kinetics (Fig. 11(h and i)). LSV recorded before and after 1000 cycles of CV at 50 mV s\(^{-1}\), showed a negligible shift in the overpotential. All the above experiments were also carried out in alkaline medium (1 M KOH) and CoPS/N–C was proven to be active towards HER with an overpotential of ~128 mV, with Tafel slope of 78 mV dec\(^{-1}\).

Wu and co-workers demonstrated a method to preserve the framework by in situ confinement of species inside the pore which prevents the agglomeration of the active sites in the carbon matrix.\(^{160}\) A Cu-based MOF [HKUST-1; Cu\(_3\)(BTC)\(_2\)(H\(_2\)O)\(_3\)] and a Mo-based POM (H\(_3\)PMo\(_{12}\)O\(_{40}\)) were coprecipitated to form octahedral nanocrystallites of NENU-5 where POMs are confined inside the pores of HKUST-1. Upon carbonization of NENU-5 at 800 °C under N\(_2\) atmosphere, Mo reacts with carbonaceous ligands to form MoC\(_x\)–Cu, whereas the presence of Cu\(^{2+}\) assists in the formation of MoC\(_x\) by getting reduced to Cu. Finally, MoC\(_x\)–Cu were converted to MoC\(_x\) by etching the metallic Cu on treating with ferric chloride solution. A pictorial representation of the scheme is shown in Fig. 12(a). The Mo included structure shows a visible colour change, and also changes in the PXRD pattern (Fig. 12(b)).

The FESEM images of NENU-5 and MoC\(_x\) show strikingly similar morphology (Fig. 12(c and d)). The high resolution

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**Fig. 8** Schematic representation of one-step PANI-assisted synthesis of a bimetal–organic framework NRAs and its derived porous CoFe\(_2\)O\(_4\)/C NRAs. Reproduced with permission from ref. 149. Copyright 2017 Wiley.
transmission electron microscopy (HRTEM) image and the elemental mapping confirmed the uniform distribution of Mo nanoparticles with diameter of ~5 nm embedded on the carbon matrix (Fig. 12(e and f)). The electrocatalytic activity of MoC–Cu towards HER was studied in 0.5 M H₂SO₄ and 1 M KOH. An overpotential of 25 mV and 80 mV is required to initiate a cathodic current in acidic and alkaline media, respectively (Fig. 12(g and h)). The corresponding Tafel slopes of 53 mV dec⁻¹ and 59 mV dec⁻¹ signify fast kinetics. The electrocatalyst also showed excellent stability in both the media (Fig. 12(i)). This excellent HER activity is attributed to the Pt like electronic properties of MoC, brought about by high porosity and also by the availability of more active sites. The confinement strategy explored here negates agglomeration of active sites, keeping them exposed and catalytically active. The homogeneity of the morphology and the mesoporosity facilitates the charge and mass transport and hence providing faster reaction kinetics. The in situ confinement strategy for preparing electrocatalyst also opens up opportunities to design and synthesize high performing nanostructured transition metal carbides. These reports establish that indeed it is possible to synthesize highly efficient catalysts for HER from MOF derivatives.

4. Bifunctional electrocatalysts from MOF derived materials

While the superior activity of a material in a particular electrochemical reaction, e.g., OER, ORR or HER, is of utmost importance, practical electrolyzers require efficiency in multiple reactions for real-time use. While bifunctionality in
OER and HER is essential for a material to have practical use in electrochemical water splitting, bifunctionality in ORR and OER is important in reversible fuel cells and metal-air batteries. In the attempt to explore MOF derived electrocatalysts, a few bifunctional catalysts were also developed.

A commendable approach was reported by Qiao et al. who fabricated efficient bifunctional electrocatalysts by growing Co-naphthalenedicarboxylate \( \text{[Co(C}_{12}\text{H}_{6}\text{O}_{4})\cdot\text{H}_{2}\text{O}]_{4} \) framework on a Cu foil by hydrothermal methods at moderate temperature (~80 °C). As perceived from various microscopic techniques, the MOF grows on the Cu foil as arrays of nanowires with a diameter of ~250 nm. These when subjected to carbonization under N\(_2\) atmosphere resulted in porous hybrid carbon nanowires with Co\(_3\)O\(_4\) nanoparticles (Co\(_3\)O\(_4\)-NA) (Fig. 13(a)).

Fig. 10  (a) Asymmetric unit of HKUST-1. (b) Packing of HKUST-1 in a direction. Reproduced with permission from ref. 157. Copyright 2012 Wiley. (c) Pyramidal HKUST nanoparticle before carbonization. (d) HKUST-1 after carbonization. (e) TEM image of Cu–Pd/NPCC/EC. Reproduced with permission from ref. 150. Copyright 2015. Elsevier. (f) LSV scan for HER on (a) bare GCE, (b) Cu/NPCC/HT/GCE, (c) Cu/NPCC/EC/GCE, (d) Pd/GCE, (e) Cu–Pd/NPCC/HT/GCE and (f) Cu–Pd/NPCC/EC/GCE in 0.5 M H\(_2\)SO\(_4\) solution at 5 mV s\(^{-1}\). Reproduced with permission from ref. 151. Copyright 2018. Elsevier.
microscopic images of the formed nanowires are shown in Fig. 13(b and c). Domains of crystalline Co$_3$O$_4$ demarcated by lattice fringes surrounded by amorphous phases were visible in HRTEM (Fig. 13(d and e)). Co$_3$O$_4$-NA grown on Cu-foil can be directly used as working electrode for studying the electrocatalytic activity towards OER without any further processing. The authors systematically studied the catalytic contribution from Cu-foil, MOF and Co$_3$O$_4$-NA. The polarization curve
recorded in 0.1 M KOH shows both MOF and Co$_3$O$_4$-NA to be active for OER (Fig. 13(f)). The corresponding Tafel plots are shown in Fig. 13(g). However, the pristine MOF showed a much lesser activity with the onset at $\sim$1.55 V vs. RHE, whereas Co$_3$O$_4$-NA showed an onset at $\sim$1.47 V vs. RHE. Co$_3$O$_4$-NA is not only active towards OER but are also active towards ORR and selectively produces water as the product, following a four electron pathway (Fig. 13(h)). This exceptional catalytic behavior arises due to a complex synergistic effect between Co$_3$O$_4$ and C which causes the active Co sites to be more electrophilic, thereby facilitating the adsorption and oxidation of OH$^-$ groups, showing enhanced OER activity in alkaline solutions. Also, the nanowire morphology of the catalysts assists in fast electron transfer. The nanowire morphology is also particularly advantageous in providing swift electrolyte penetration, whereas the mesopores assist the access of reactants in the electrolyte to the active sites. The nanowire morphology also helps in the faster release of the evolved O$_2$. The most noteworthy advantage is growing the catalyst on conducting supports which ensures better adhesion between nanowire arrays and substrates, greatly enhancing the electron transport, structural stability, without the use of binders and conductive fillers. Owing to this fast kinetics, Co$_3$O$_4$-NA shows a Tafel slope much smaller than benchmark IrO$_2$/C for OER.

Recently, we have reported synthesis and detailed characterization of an efficient and stable bi-functional electrocatalyst derived from a 3D metal–organic framework $\{[\text{Co(bpe)}_2(NCN)]_2\text{[CN]$_2$:5H$_2$O})\}$ (Co-MOF), (bpe = 1,2-bis(4-pyridyl)ethane and N(CN)$_2$ = dicyanamide) (Fig. 14(a)). The 3D framework of Co-MOF is composed of bridging bpe linkers which forms a 2D $[\text{Co(bpe)}_2]$ layer and this is further pillared by N(CN)$_2^-$ anions (Fig. 14(b)). The presence of large no of Co$^{II}$-
Nitrogen moieties in the structure could lead to the formation of electrochemical active M–N–C sites. This Co-MOF was chosen as a carbonization template due to two reasons, (a) the presence of electroactive CoII metal ion, which can also catalyze organic precursors to form carbon nanotubes under pyrolyzing condition at H2 atmosphere; (b) secondly, the presence of N(CN)2, a well-known chemical precursor for g-C3N4 that can lead to the formation of N-doped graphitic nanostructures. The two different catalysts were prepared by pyrolyzing Co-MOF at different conditions. Pyrolyzing at 800 °C under Ar atmosphere and at 800 °C under H2/Ar atmosphere resulted in Co-NCNT-Ar and Co-NCNT-H2 (NCNT = nitrogen doped carbon nanotube), respectively. Furthermore, Co-NCNT-H2 was calcined in air at 250 °C to obtain Co3O4@Co/NCNT composite material. Both the carbonized materials were characterized by XRD, Raman spectroscopy, XPS, FESEM and TEM techniques. XRD pattern of Co-NCNT-Ar suggests the presence of metallic cubic Co phase on the regular graphitic carbon (Fig. 14(c)). Raman spectra showed D and G bands and their ratio suggests a high degree of graphitization in Co-NCNT-Ar and Co3O4@Co/NCNT (Fig. 14(d)). Raman spectra of Co3O4@Co/NCNT also showed the bands characteristics of typical Co3O4 spinel phase. XPS survey spectra for both the composite materials showed the presence of Co, C, N and O. Along with metallic Co, the presence of spinel like Co–N4, Co(CO)4 and Co–N–C was also realized from XPS spectra (Fig. 14(e)). The detailed morphological feature of the carbonized product was realized from FESEM and TEM imaging. The microscopic study showed that in Co/NCNT-Ar, the Co nanoparticles are embedded in a continuous self-coiled N-doped carbon nanotube matrix, where the average diameter of Co nanoparticles and nanotubes are in the range of 50–110 nm and 50–200 nm, respectively (Fig. 14(f)). HRTEM showed lattice fringes with spacing of about 0.209 nm which corresponds to the {111} of the cubic metal structure. Pyrolysis under H2 atmosphere also led to the formation of self-coiled dense N-doped carbon nanotube where metallic Co nanoparticles (40–50 nm) are embedded. After calcination in air, TEM images revealed the formation of core–shell nanoparticles Co3O4@Co encapsulated in a CNT matrix. The length and diameter of the nanotubes are in the range of 100–200 nm and 10–20 nm, respectively. The lattice fringe of the outer shell of Co3O4@Co nanoparticles is 0.46 nm which is characteristic cubic [111] structure of Co3O4 and higher than that of Co (0.209 nm) (Fig. 14(g)). The permanent porosity was measured by the N2 adsorption study at 77 K to elucidate the textural parameters (Fig. 14(h)). Both Co-NCNT-Ar and Co3O4@Co/NCNT showed typical type IV isotherm with BET surface area of 58 and 118 m2 g−1, respectively. Both the samples showed wide pore size distribution (2–30 nm). Electrochemical ORR and OER activities for Co-NCNT-Ar and Co3O4@Co/NCNT were tested. Cyclic...
voltammetry (CV) in O₂ saturated 0.1 M KOH solution for Co-NCNT-Ar and Co₃O₄@Co/NCNT was measured and showed good catalytic activity towards ORR at 0.87 and 0.9 V (vs. RHE), respectively (Fig. 15(a and b)). LSV of Co₃O₄@Co/NCNT using a RDE electrode at different rotation speeds revealed a remarkable catalytic performance for ORR according a current density of 1 mA cm⁻² at low overpotential of 0.88 V (onset 0.93 V vs. RHE), and this value is 0.83 for Co-NCNT-Ar (onset at 0.9 V vs. RHE). Co₃O₄@Co/NCNT reveals a very close ORR activity to that of commercial Pt/C (Fig. 15(c and d)). Koutecky-Levich (K-L) plots suggested first order reaction kinetics with respect to the dissolved O₂ concentration and number of electrons transferred (n) per O₂ molecule during the ORR to be in the range of 3.92 to 3.96 suggesting the reduction of O₂ predominantly through the 4 electron transfer pathway. Interestingly Co-NCNT-Ar showed 2 electron transfer pathway. RRDE measurement further confirmed it based on the percentage of H₂O₂ production (Fig. 15(e)). Co-NCNT-Ar showed about 54% H₂O₂ production in the entire potential range (0.8 to 0.3 V vs. RHE) and Co₃O₄@Co-NCNT showed only 1% H₂O₂ production. The relatively lower Tafel slope of 61 mV dec⁻¹ for Co₃O₄@Co-NCNT compared to Co-NCNT-Ar (74 mV dec⁻¹) indicating faster reaction kinetics for the former (Fig. 15(f)).

OER activity is also observed to be superior for Co₃O₄@Co-NCNT compared to Co-NCNT-Ar as the former attained a current density of 10 mA cm⁻² at a potential of 1.61 V vs. RHE, whereas later reached the same current density at 1.76 V at 1600 rpm (Fig. 16(a)). In a similar condition RuO₂ and IrO₂
Co\textsubscript{3}O\textsubscript{4}@Co-NCNT surpasses the benchmark catalysts (Fig. 16(a)). The Tafel slopes for the OER are 58.7 mV per decade and 80 mV per decade for Co\textsubscript{3}O\textsubscript{4}@Co-NCNT and Co-NCNT-Ar, respectively, indicating reaction kinetics substantially faster for the former (Fig. 16(b)). Interestingly the overpotential difference between ORR (at current density of \( 1 \text{ mA cm}^{-2} \)) and OER (at current density of \( 10 \text{ mA cm}^{-2} \)) is about 0.73 V, indicating high efficiency of Co\textsubscript{3}O\textsubscript{4}@Co-NCNT as ORR and OER bifunctional catalyst (Fig. 16(c)). This is an important parameter for evaluating the bifunctionality for a reversible oxygen electrode. This value is lowest among all the state of art catalysts. Furthermore chronoamperometric stability measurement suggested no significant decline in the current in the course of 30–40 h of continuous polarization, indicating high stability of both the catalyst (Fig. 16(d)). The high catalytic activity for Co\textsubscript{3}O\textsubscript{4}@Co-NCNT was attributed to the high surface area, presence of highly active sites and the synergistic interaction between conductive metallic Co cores and the semi-conductive Co\textsubscript{3}O\textsubscript{4} shell. All these parameters facilitate an easy mass diffusion and faster charge transfer in the NCNT network.

In a similar work by Muhler et al., bifunctionality in OER and ORR was demonstrated based on core–shell Co@Co\textsubscript{3}O\textsubscript{4} nanoparticles embedded in CNT grafted N-doped polyhedral obtained by the pyrolysis of ZIF-67.\textsuperscript{120} In this case, in the carbonized form, polyhedrons containing Co nanoparticles are wrapped in few layers of CNT. The Co@Co\textsubscript{3}O\textsubscript{4} particles are wrapped in graphene like highly graphitized carbon sheets. The width of the Co\textsubscript{3}O\textsubscript{4} shell can be controlled by the time of calcination of the sample in O\textsubscript{2} environment and two such samples were prepared with calcination times of 2 and 6 h respectively (Co@Co\textsubscript{3}O\textsubscript{4}/NC-1 and Co@Co\textsubscript{3}O\textsubscript{4}/NC-2). While Co@Co\textsubscript{3}O\textsubscript{4}/NC-1 shows a better activity in ORR, Co@Co\textsubscript{3}O\textsubscript{4}/NC-2 has a better performance in OER, with a current density of \( -10 \text{ mA cm}^{-2} \) achieved merely at 1.64 V. However, upon considering bifunctionality, it is observed that overall Co@Co\textsubscript{3}O\textsubscript{4}/NC-1 is a more efficient electrocatalyst, with comparable activity in ORR with Pt/C and in OER with IrO\textsubscript{2} and Pt/C. As suggested in this work, the small overvoltage in bifunctionality between ORR and OER translates to energy saving of at least 150 mV and 100 mV for Co@Co\textsubscript{3}O\textsubscript{4}/NC-1 and Co@Co\textsubscript{3}O\textsubscript{4}/NC-2, respectively.\textsuperscript{120}

An attempt towards fabricating bifunctional catalysts was also made by carbonizing ZIF-67 at various temperatures from 600–2000 °C under Ar atmosphere.\textsuperscript{154} While at lower temperature, the morphology of the ZIF-67 nanoparticles is maintained in the carbonized form, at higher temperatures, carbon nanotubes are observed at the surface, whose formation has been catalyzed by the presence of cobalt. At 1200 °C, cobalt form large aggregates, wrapped by sheets of graphitic carbon. Beyond 1600 °C, the morphology was primarily that of hollow carbon onions. The sample that had been pyrolyzed at 800 °C, i.e., Co@C-800, shows an excellent ORR activity with an onset potential of 0.92 V (vs. RHE), along with a 4 electron transfer pathway and excellent stability. This stability can be attributed...
to the protection of the Co nanoparticles by the layers of carbon matrix. Moreover, the excellent bifunctionality of Co@C-800 is established upon investigating its role in OER. It showed an onset potential of 1.43 V and a potential of merely 1.61 V is required to achieve a current density of 10 mA cm$^{-2}$. In this work, the direct effect of the carbonization temperature on the electrochemical activity has been shown. In case of Co@C-800, Co and N dopants could induce a positive charge on the adjacent carbon atoms, thereby assisting in the adsorption of OH$^{-}$ and electron transfer.

MOF derived dual catalysts have been also attempted for HER and OER, thereby showing activity as a catalyst for overall water splitting. A recent work involves the pyrolysis of the zeolitic imidazolate framework ZIF-8 (Zn[MIM])$_2$ to generate a N containing porous carbon. Using cathodic polarization treatment (CPT) up to 8 h, the N and O containing surface groups are modulated without compromising its large surface area or pore size distribution (Fig. 17(a)). A sample having undergone 6 h CPT shows an excellent HER activity in 0.5 M H$_2$SO$_4$ electrolyte with an overpotential of 155 mV, a Tafel slope of 54.7 mV dec$^{-1}$ and an exchange current density of 0.063 mA cm$^{-2}$ (Fig. 17(b)). CPT of 4 h results in excellent OER activity in 0.1 M KOH electrolyte with an overpotential of 476 mV and a Tafel slope of 78.5 mV dec$^{-1}$ (Fig. 17(c)). These carbon electrolyzers successfully run a two-electrode water splitting reaction at a current density of 10 mA cm$^{-2}$ over 8 h under a potential of 1.82 V with a Faradaic efficiency of $\sim$99% in 0.1 M KOH (Fig. 17(d)). By careful investigation, it has been found that in this case the presence of oxidized N and carboxylic O sites on the surface is specifically effective in improving the HER kinetics as these are the active sites. Although in this case, it has been proven that surface area and pore size are not determining factors, larger ECSA definitely ensures better activity. Moreover, investigation of Raman spectra also showed that higher carbonization time inflicts higher number of defects and severe structural damage to the graphitic carbon matrix, thereby lowering activity.

ZIF-67 because of its large cobalt content is an automatic choice for carbonization. Porous Co-P/NC nanopolyhedrons comprising CoP$_x$ nanoparticles embedded in N containing C matrix was prepared by the direct carbonization of ZIF-67, followed by phosphidation. This shows excellent performance in both HER and OER in 0.1 M KOH, with low overpotentials and high current density, showing an activity even better than Pt or IrO$_2$. This superior activity has been attributed to their high specific surface area along with interconnected mesoporosity, good conductivity and the dual effect of doped CoP$_x$ and the N-dopants. In a similar approach, Sun et al. also grew ZIF-67 on a carbon cloth, followed by subsequent carbonization and phosphidation to yield a Co, N and P containing carbon matrix named Co-P/NC/CC. This material shows an excellent HER activity in 1.0 M KOH with an onset potential of $-0.1$ V vs. RHE. It also showed an onset potential of 1.5 V vs. RHE for OER in...
1.0 M KOH and achieves a current density of 10 mA cm\(^{-2}\) at an overpotential of merely 330 mV. The superior activity of Co–P/NC/CC is not only because of a larger ECSA, but also because the large content of cobalt phosphides. A two-electrode electrolyzer fabricated using this system at cathode and anode in 1.0 M KOH for overall water-splitting, where it shows an onset of 1.7 V and achieves current densities of 10 and 100 mA cm\(^{-2}\) at cell voltages of 1.77 and 1.93 V, respectively.\(^{162}\)

Bifunctionality achieved in mesoporous nitrogen doped graphene (MNG) obtained from ZIF-67 has also been exploited in a multi-functional electrode for Zn–air battery and water splitting device. Here, ZIF-67 has been first grown as fibers on
Table 1 Concise summary of the MOF derived carbon nanocomposites described in this review

<table>
<thead>
<tr>
<th>Type of electrocatalyst</th>
<th>Material</th>
<th>Derived from MOF</th>
<th>Morphology</th>
<th>Heteroatoms/transition metals present</th>
<th>Activity</th>
<th>Ref. no.</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>ORR catalyst</strong></td>
<td>P-CN-Co-20</td>
<td>BMZIF (bi-metallic ZIF-67)</td>
<td>Co nanoparticles on graphitized carbon</td>
<td>N, P, Co</td>
<td>Onset: -0.04 V vs. Ag/AgCl</td>
<td>141</td>
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<tr>
<td></td>
<td>NCD-1</td>
<td>${Mg,(ndc)<em>2(HCO_2)(H_2O)}</em>{\cdot}$N$_2$H$<em>4$Co$</em>{20}$</td>
<td>Spherical dots</td>
<td>N</td>
<td>1 mA cm$^{-2}$ at 0.36 V vs. RHE</td>
<td>129</td>
</tr>
<tr>
<td></td>
<td>Co@CN</td>
<td>${[NH_2Me_2]_22H_2O\cdot$2H$<em>2$O-DMF$}</em>{\cdot}$</td>
<td>Co nanoparticles on carbon nitride</td>
<td>Co, N</td>
<td>1 mA cm$^{-2}$ at 0.8 V vs. RHE</td>
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</tr>
<tr>
<td></td>
<td>NSMC-900</td>
<td>SCUT-12 $[\text{Zn}_3(\text{TDC})<em>2(\text{DABCO})]</em>{\cdot}$4DMF</td>
<td>Porous planar</td>
<td>N, S</td>
<td>Onset: 0.98 V vs. RHE; comparable to 40 wt% Pt/C</td>
<td>144</td>
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<td></td>
<td>MDC300</td>
<td>ZIF-67</td>
<td>Polyhedral</td>
<td>Co, N</td>
<td>Onset: 0.86 V vs. RHE</td>
<td>145</td>
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<td></td>
<td>P-Z8-Te-1000</td>
<td>ZIF-8 grown on Te nanowires</td>
<td>Fibrous</td>
<td>N, P</td>
<td>Onset: -0.22 V vs. Ag/AgCl; better than 20 wt% Pt/C</td>
<td>146</td>
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<td></td>
<td>BHPC-950</td>
<td>ZIF-8 supported on silica matrix</td>
<td>Interconnected macroporous carbon texture</td>
<td>N</td>
<td>Onset: 0.93 V vs. RHE; Zn-air &amp; Li-S batteries</td>
<td>147</td>
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<td><strong>OER catalyst</strong></td>
<td>Co$_2$-MoC/NC-1</td>
<td>Co-Mo-MOF</td>
<td>3D nanowire assembly</td>
<td>Co, Mo, N</td>
<td>Onset: 1.52 V vs. RHE; Tafel slope 89.8 mV dec$^{-1}$.</td>
<td>148</td>
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<td>NF@NC@Co/Fe$_2$O$_4$-C NRAs</td>
<td>MOF-74-M ($M =$ Fe, Co)</td>
<td>Nanorod array</td>
<td>Co, Fe, N</td>
<td>Onset: 1.45 V vs. RHE; Tafel slope 45 mV dec$^{-1}$.</td>
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<td>ZNDC-1100</td>
<td>Bimetallic ZIF-8 (Ni, Zn)</td>
<td>Tubular graphitic nanostructure</td>
<td>Ni, N</td>
<td>10 mA cm$^{-2}$ at 1.67 V</td>
<td>163</td>
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<td><strong>HER catalyst</strong></td>
<td>Cu/NPC</td>
<td>HKUST-1</td>
<td>“Cauliflower-like” morphology</td>
<td>Cu</td>
<td>10.66 mA cm$^{-2}$ at -1.0 V</td>
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<td>Cu-Pd/NPCC/EC</td>
<td>Electrochemically prepared HKUST-1</td>
<td>Metal NPs on porous network</td>
<td>Cu, Pd</td>
<td>Onset: -0.13 V vs. RHE</td>
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<td>CoPS/N-C</td>
<td>ZIF-67</td>
<td>Polyhedra with CoPS NPs</td>
<td>Co, P, S, N</td>
<td>Overpotential: 80 mV vs. RHE; Tafel slope 68 mV dec$^{-1}$.</td>
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<td>MoC$_x$</td>
<td>NENU-5</td>
<td>Nano-octahedrons</td>
<td>Mo</td>
<td>Overpotential: 80 mV vs. RHE in alkaline and 25 mV in acidic media</td>
<td>160</td>
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<tr>
<td><strong>Bifunctional electrocatalyst</strong></td>
<td>Co$_3$O$_4$C-NA</td>
<td>${\text{Co}((\text{C}_6\text{H}_6\text{O}_4)\cdot(\text{H}_2\text{O})<em>4)}</em>{\cdot}$ framework grown on Cu foil</td>
<td>Hybrid carbon nanowires</td>
<td>Co</td>
<td>OER onset: 1.61 V vs. RHE; ORR half-wave potential: 0.78 V vs. RHE; Tafel slope: 89 mV dec$^{-1}$.</td>
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<td>Co$_3$O$_4$@Co/NCNT</td>
<td>Co-MOF</td>
<td>N-doped carbon nanotubes with metal NPs</td>
<td>Co, N</td>
<td>OER onset: 0.9 V vs. RHE. Comparable to RuO$_2$ and IrO$_2$.</td>
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<td></td>
<td>Co-P/NC</td>
<td>ZIF-8</td>
<td>Nanopolyhedrons</td>
<td>Co, P, N</td>
<td>HER: -154 mV overpotential to reach 10 mA cm$^{-2}$. OER onset: 1.5 V vs. RHE. Water-splitting</td>
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<td>MNG-CoFe</td>
<td>ZIF-67 fibres</td>
<td>Fibrous</td>
<td>Co, Fe, N</td>
<td>ORR, OER, HER better than benchmark catalysts. Zn-air battery and water-splitting device</td>
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</table>
the surface of NaCl crystals and post carbonization, NaCl has been washed away with acid, to yield MNG, which has a much higher ECSA than the porous carbon derived from typical polyhedral ZIF-67. MNG–CoFe with encapsulated CoFe alloy nanoparticles showed excellent ORR, OER and HER performances, comparable to benchmark noble metal catalysts. MNG–CoFe when assembled as a multifunctional electrode in the rechargeable Zn–air battery showed high power density and cycling durability. Moreover in a water-splitting device, it showed very low overpotential, outperforming the benchmark catalysts. \(^{164}\)

Although in most cases, Co and Ni nanoparticles have been exploited for catalysis, Cu has also been used for bifunctional catalysis. For this purpose, a MOF Cu-NP MOF (containing Cu\(^{2+}\), pyrazine and HEDP = 1-hydroxyethylidene-1,1-diphosphonic acid) has been used where Cu\(^{2+}\) is connected by P containing 1-hydroxyethylidene-1,1-diphosphonic acid (HEDP) and N containing pyrazine. Upon carbonization, Cu\(_n\)P nanoparticles coated by N and P-doped carbon shell are formed, which extends to a hierarchical porous carbon matrix (Cu\(_n\)P@NPPC). Cu\(_n\)P@NPPC is an excellent catalyst for both HER and OER. \(^{165}\)

Thus several unique approaches have been adopted to synthesize MOF-derived porous carbon with distinct morphologies and composition showing superior activity. While we have mentioned some over here, many more attempts have been undertaken in this global endeavor of finding cheap alternatives of noble metals for renewable energy technology. The various MOF-derived electrocatalysts that have been discussed in this review has been concisely summarized in Table 1 with their properties and activities.

5. Conclusions and future perspectives

This review documents some of the recent research conducted to develop low cost, noble metal-free electrocatalysts derived from MOF. MOFs act as perfect template for generating such carbon materials owing to their innate porosity, stable structure, high carbon content and ease of heteroatom doping. However, the synthesis of efficient carbon based materials requires careful selection of the MOF precursor and the annealing temperature, along with morphology and size control. Till date, quite a few materials have been generated using MOF which are excellent noble metal-free electrocatalysts and have helped in taking a big stride towards cheap alternative catalysts. New and innovative approaches were brought into light which could be utilized to design a plethora of such functional materials. The methodically developed MOF derived materials having favourable electronic structure based on heteroatom doped carbon matrix along with metal or metal compound NPs, can show excellent electrochemical activity. Moreover, suitable morphologies and pore structure that facilitate electron and mass transport are instrumental in developing highly potent electrocatalyst for OER, ORR and HER.

Highly potent MOF-derived carbon-based electrocatalysts can be synthesized by carefully considering the following points:

1. Careful selection of the MOF is supremely important since ligand composition provides the carbon template with heteroatoms like N, P and S. Similarly, redox-active metal ion from the MOF provides the metal or metal oxide nanoparticles.

2. Morphology, size and shape of the as synthesized MOF is also crucial as it determines the morphology and corresponding textural parameter of the resultant carbonaceous material.

3. Pyrolysis temperature and condition are also important parameters which plays a crucial role in determining the morphology and composition of the resultant carbon nanocomposite.

Targeted design and synthesis of tailored MOFs has the potential to develop next-generation materials with superior activity for the commercialization of energy devices like fuel cells and metal–air batteries. Until now, MOF derived functional materials have been exploited as bifunctional catalysts but trifunctional catalysts which can catalyze all the three half reactions—OER, ORR and HER efficiently are yet to be developed from such materials. Such trifunctional materials are ideal for practical use in regenerative fuel cells, electrolyzer, overall water-splitting and metal–air batteries. This missing link can be filled by careful considerations of the design strategies discussed in this review. Thus, this review not only paves way for better understanding of the foundation of designing materials but also throws a challenge to develop an all-in-one trifunctional catalyst from MOF precursor. Another obstacle that needs to be overcome is the poor conductivity of pristine MOFs in order to develop electrochemically active MOFs. Absence of studies that provide a deep insight to the mechanism of the electrocatalytic activities of these materials is also another hurdle in their otherwise steady advancement and this void can be filled by suitable proper theoretical approaches. Additionally, merely the development of new materials is not adequate to fight the global energy demand, sincere efforts to support their application in clean energy technology are also necessary, which would require more rigorous and meticulous performance and stability test. Thus, the challenge is still open to develop electrocatalysts with supreme efficiency and stability along with proper mechanistic insights for successful commercialization of sustainable clean energy techniques.

Conflicts of interest

There are no conflicts to declare.

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