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Recent advances in heterostructured cathodic electrocatalysts for non-aqueous Li–O₂ batteries†

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Developing efficient energy storage and conversion applications is vital to address fossil energy depletion and global warming. Li–O₂ batteries are one of the most promising devices because of their ultra-high energy density. To overcome their practical difficulties including low specific capacities, high overpotentials, limited rate capability and poor cycle stability, an intensive search for highly efficient electrocatalysts has been performed. Recently, it has been reported that heterostructured catalysts exhibit significantly enhanced activities toward the oxygen reduction reaction and oxygen evolution reaction, and their excellent performance is not only related to the catalyst materials themselves but also the special hetero-interfaces. Herein, an overview focused on the electrocatalytic functions of heterostructured catalysts for non-aqueous Li–O₂ batteries is presented by summarizing recent research progress. Reduction mechanisms of Li–O₂ batteries are first introduced, followed by a detailed discussion on the typical performance enhancement mechanisms of the heterostructured catalysts with different phases and heterointerfaces, and the various heterostructured catalysts applied in Li–O₂ batteries are also intensively discussed. Finally, the existing problems and development perspectives on the heterostructure applications are presented.

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1. Introduction

Since the 21st century, the fossil energy crisis and environmental problems have continuously forced people to search for clean and effective energy storage and conversion technologies.^{1–4} Currently, the rapid developments of fuel cells,^{5–8} supercapacitors,^{9–11} and batteries^{12–15} have attracted more and more research attention. A comparison of different available and currently under-development energy storage devices is provided in Fig. 1 in terms of their theoretical energy densities. Metals (such as Al, Mg, Fe, and Zn) are appropriate for metal–air batteries in the aqueous environment.¹⁶ They are thermodynamically unstable in aqueous environments, but under specific conditions, their surfaces may be passivated by corresponding oxides or hydroxides, rendering them compatible with aqueous electrolytes to some extent. Zn–air batteries have been intensively assessed among these aqueous metal–air batteries. Zn–air batteries with a low cost of fabrication could deliver a theoretical energy density of 1086 W h kg^{−1} and

theoretical working voltage of 1.65 V. However, the practically attainable energy density of Zn–air batteries is usually around 350 W h kg^{−1},¹⁷ making them more suitable for low-power applications, such as small hearing aids rather than future energy vehicles and large-scale grids. Fe–air batteries show a theoretical energy density of 1229 W h kg^{−1}, comparable to that of Zn–air batteries, but their practical energy density is much lower (60–80 W h kg^{−1}),¹⁸ and is only one fifth of Zn–air batteries. As for Al–air and Mg–air batteries, they exhibit a significant theoretical energy density of 8100 and 6815 W h kg^{−1}, respectively. Nevertheless, due to the severe parasitic corrosion of their metallic anodes in contact with the aqueous electrolyte, only a very small fraction of the electrodes can be

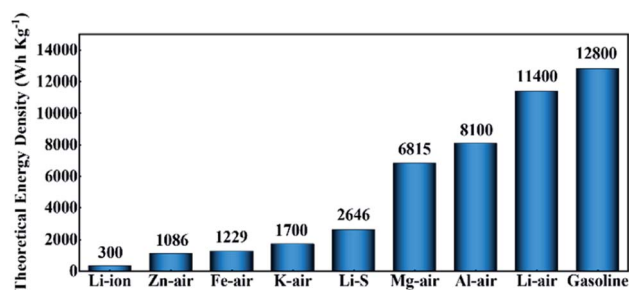


Fig. 1 Gravimetric energy densities (W h kg^{−1}) of typical rechargeable batteries and gasoline.^{16,21,22}

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actually utilized.^{19,20} Among all metal–air batteries, nonaqueous Li–O₂ batteries display the largest theoretical energy density of 11 400 W h kg⁻¹ (even close to that of gasoline). In addition, the anticipated realistic energy density of nonaqueous Li–O₂ batteries matches the useable energy density of gasoline for automotive applications (about 1700 W h kg⁻¹), and this means that the nonaqueous Li–O₂ batteries hold great potential to be one of the most likely energy storage devices for future energy vehicles, which is considered as an important solution to deal with the problems of global warming and energy crisis.^{23–25} However, they still face many problems before practical use, such as high charging overpotentials, poor cycle stability, low coulombic efficiency, *etc.*^{26–28} Researchers found that the main reason for these problems is the sluggish dynamics of the oxygen reduction reaction (ORR) and oxygen evolution reaction (OER) during discharging and charging. In order to solve these obstacles, various ways have been used to accelerate the ORR/OER kinetics, including the addition of redox mediators, applying photo-assisted systems, and the development of various catalysts. The concept of using a photocatalyst-assisted charging strategy to alleviate the serious problem of high overpotentials in Li–O₂ systems was introduced by Yu *et al.*,³³ and it was found that the photogenerated holes in the photocathode might aid in the Li₂O₂ oxidation processes, thereby greatly reducing the overpotentials. Yu's group³⁴ used cobalt-tetramino-benzoquinone (Co-TABQ) as a bifunctional catalyst for Li–O₂ batteries under visible light conditions, exhibiting an overpotential gap of 0.2 V with a round-trip efficiency of 94.0%. Wang *et al.*³⁵ established a novel photo-assisted Li–O₂ system by applying a hierarchical TiO₂/Fe₂O₃ heterojunction, which delivered an ultralow overpotential gap of 0.19 V. However, the key to develop these photo-assisted Li–O₂ systems lies in the light itself, which would undoubtedly bring about difficulties to the practical application of Li–O₂ batteries. So efficient cathode catalysts without photo-assistance are urgently needed. It is evident that noble metals and their oxides with high catalytic performance are undoubtedly the ideal choices.^{36–38} However, the scarcity in the Earth's crust and high price of precious metals hinder their large-scale usage in Li–O₂ batteries. Therefore, the rational design of cathode catalysts in Li–O₂ batteries in recent years was mainly carried out through two pathways: (1) reducing the loading of precious metals *via* rationally constructing nanosized or single-atom catalysts.^{38–41} (2) Employing non-precious metal catalysts, including non-precious alloys, metal oxides,^{42,43} metal sulfides,^{44,45} metal nitrides,^{46,47} metal carbides,^{48,49} carbon materials, *etc.*^{50–52}

However, most of these non-precious catalysts only show single catalytic activity for the OER or ORR, and the development of high-efficiency bifunctional catalysts is actually challenging and essential for the practical application of Li–O₂ batteries.^{53–55} At present, various strategies including surface modification,^{56–58} strain engineering,^{59,60} phase modulation^{61–63} and heterostructure construction have been explored to realize the bifunctional ability for Li–O₂ catalysis.^{64–66}

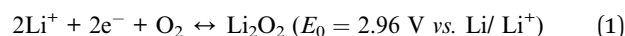
Among them, heterostructures have been intensively investigated, and are usually defined as composite structures composed of interfaces formed by different solid materials in

a broad sense, including conductors, insulators and semiconductors. The interface effect derived from the inherent electric fields among multiple components is beneficial for enhancing the electrocatalytic properties, which can remarkably promote surface reaction dynamics with fast mass and charge transfer.^{67–69}

In this review, the construction and mechanisms of Li–O₂ batteries were briefly introduced, and various functions of heterostructures in them were summarized and listed. Then, recent research reports on the application of heterostructures in Li–O₂ batteries were provided. We focused on the summary of structure–function relationships between heterostructures and their catalytic properties in these reports, aiming to reveal the underlying mechanisms of their excellent ORR and OER performance. Finally, a personal view of the challenges and opportunities faced by various heterostructures in Li–O₂ batteries was included.

2. Fundamentals of nonaqueous Li–O₂ batteries

As shown in Fig. 2a, a typical non-aqueous Li–O₂ cell consists of a Li metal anode, a porous air cathode and a non-aqueous electrolyte.^{70,71} The ideal operation of Li–O₂ batteries is based on the electrochemical formation during discharging and decomposition during charging of lithium peroxide (Li₂O₂). The typical electrochemical reaction equation is as follows:



In 2014, Bruce's group elaborated on the formation mechanisms of discharge products in Li–O₂ batteries,³² and they found that the formation pathways of Li₂O₂ are mainly controlled by the solubility of the LiO₂ intermediate, which depends primarily on the solvation of the cations by the solvent molecules. In a strong Li⁺ solvating solvent, Li₂O₂ is usually formed through surface mechanisms, and in a weak Li⁺ solvating solvent, Li₂O₂ is normally generated through solution mechanisms (Fig. 2b).^{31,72,73}

2.1. Surface mechanism and solution mechanism

In recent years, the Li₂O₂ mechanism has been intensively studied,^{31,32,74} and it is widely accepted that the ORR in Li⁺ containing non-aqueous solvents occurs starting from the one-electron reduction of O₂ to generate superoxide (O₂⁻), which combines with Li⁺ to form LiO₂. In strong Li⁺ solvating solvents,^{75–77} such as dimethylsulfoxide and 1-methylimidazole, LiO₂ is dissolved mainly in the electrolyte solution, and then it would get reduced or disproportionate to large-sized Li₂O₂ on the cathode surface.^{78–80} In weak Li⁺ solvating solvents including tetraethylene glycol dimethyl ether and acetonitrile, due to the low solubility of LiO₂ in the electrolyte solvent, LiO₂* is formed and deposits on the cathode surfaces, which is then reduced *via* an electrochemical process or disproportionation to produce a Li₂O₂ film.^{81–83}



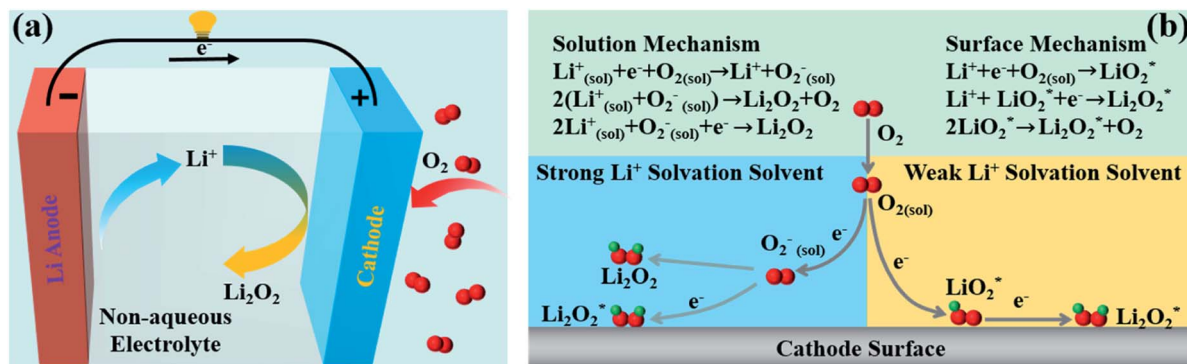


Fig. 2 (a) Schematic of a nonaqueous Li–O₂ cell. (b) Reduction mechanisms of Li–O₂ batteries through different reaction routes.^{29–32}

2.2. Current status for large-scale commercial applications

In fact, the practical application of Li–O₂ batteries is not only limited to the problem of cathode catalysts. For example, the electrolytes in Li–O₂ batteries are not only decomposed at high voltage during charging, but are also always easily reduced by superoxide. Another requirement characteristic of the electrolyte in Li–O₂ cells is low volatility,⁷³ which is very important in open systems through oxygen flows. Moreover, it is difficult to find corresponding substitutes for Li anodes, and as a result, safety issues such as explosions caused by Li dendritic short-circuits and blocking resulting from the production of thick insulating layers on the Li surface are also hard to avoid during cycling.⁸⁴ Furthermore, Li–O₂ cells currently studied in the laboratory are usually coin-cells,^{85–87} and few research studies were conducted on the implementation and evaluation of large-sized batteries for practical applications. In recent years, some groups have focused on the investigation of pouch Li–O₂ cells for future commercial applications.^{88–90} Zhao's group⁹⁰ has taken a crucial step to successfully construct a Li–O₂ pouch cell, which shows a high energy density of 768.5 W h kg^{–1} (based on the weight of the entire battery). Actually, the key points before the large-scale commercialization of Li–O₂ batteries are supposed to be the following aspects: (1) side reactions in the high-loading air electrodes, (2) pulverization of the Li-metal anodes and (3) large-scale inhomogeneity of the pouch cell. In general, although practical Li–O₂ batteries still have many problems to be solved, it is expected that through unremitting efforts, high performance Li–O₂ secondary batteries would eventually be fabricated in mass production. Once being successfully developed, Li–O₂ batteries could broaden application prospects in many fields, such as digital electronics, medical and health, electric vehicles, aerospace, etc.

3. Characteristics of heterostructures

3.1. Catalyst structure modulation

As shown in Fig. 3a, through heteroepitaxial growth, the growth orientation of active materials can be adjusted and the surfaces with high reactivity can be exposed, thereby affecting the diffusion of Li⁺ and the adsorption of O₂ species to improve the ORR and OER kinetics.^{91,92} Moreover, the defect-free interfaces

and structural connectivity between the epitaxial layer and the substrate are favorable for relaxing structural changes in the cathode caused by the generation and decomposition of Li₂O₂ during repeated cycling, preventing the active material from detaching.^{91,92} In fact, because perfect lattice matches are normally not realized, lattice distortions are found at the interfaces of the heterostructures, which significantly increase the number of active sites to enhance their electrocatalytic activities.^{93,94} Additionally, the disordered atomic arrangements can reduce the surface energies of the heterostructures and effectively change the adsorption capacity of the intermediate products, affecting the formation processes of Li₂O₂.⁹⁵

3.2. Boosting electron/ion transport

The heterostructures we are talking about actually function at special interfaces formed by the contact of two different materials. We take the PN heterojunction as an example to illustrate the formation mechanism of the built-in electric field. As shown in Fig. 3b, because the difference in Fermi energy levels between the two materials will cause electron transfer across the hetero-interfaces, an equilibrium state of equal Fermi energy levels is thus established. In order to keep their original electron affinity (ϕ) and workfunction (χ) unchanged, a space charge region will be formed. Built-in field plants will appear on both sides of the hetero-interfaces.^{96,97} Due to the action of the built-in electric field, the transportation of Li⁺ and electrons will be accelerated, thereby promoting the ORR and OER kinetics.

3.3. Synergetic effect

Different components of heterostructure composites play various roles in the electrocatalytic processes, notably exhibiting superior performance, compared with any one-component counterparts (Fig. 3c).^{98,99} For example, it is reported that graphene and CuGeO₃ could respectively act as ORR and OER electrocatalytically active components, and their heterogeneous structures will exhibit bifunctional electrocatalytic properties with stronger ORR and OER activities.¹⁰⁰ In addition, the combination of NiCo₂O₄ and NiO could deliver high electrocatalytic activity with effective decomposition of by-products during cycling, which is mainly due to the synergy effect from



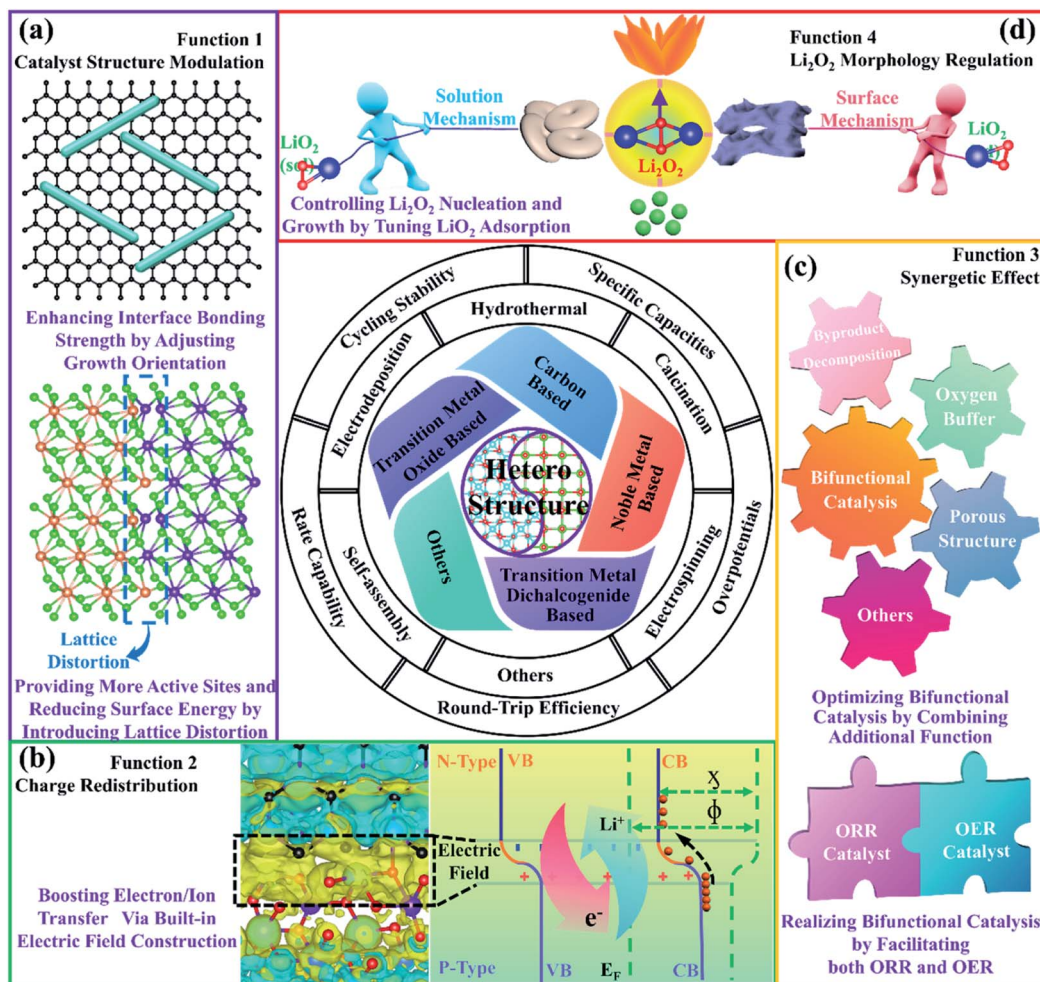


Fig. 3 Summary of the typical heterostructures and their functions in Li–O₂ batteries.

heterogeneous structures.¹⁰¹ Furthermore, it is evident that CeO₂ as an O₂ buffer combined with materials with high OER catalytic activity can exhibit larger specific capacity and lower overpotentials than those of single components.^{102–104} More importantly, porous nanostructured heterostructures also aid in accelerating the transportation of electrolyte and O₂, as well as providing a large number of active sites to facilitate the electrocatalytic reactions.

3.4. Facilitating O₂ transfer and regulating the morphology of discharge products

In fact, the strong interactions of the heterostructures can effectively change the charge distribution of the active sites, thereby tuning the adsorption energy of the catalysts to O₂ species (Fig. 3d).^{105,106} If the adsorption energy of O₂ is perfected, O₂ transport is largely promoted. Generally speaking, the change in the adsorption energy of O₂ species will be linked to the morphologies of the discharge products, especially LiO₂; as the adsorption energy of O₂ increases, Li₂O₂ tends to form a film and small particles by the surface growth model.¹⁰⁷ If it is reduced, Li₂O₂ usually becomes toroid *via* the solution growth model.¹⁰⁸ For heterostructured composite materials, Li₂O₂

deposition mechanisms based on two components are usually combined, and special shapes are thus constructed *via* the competition of the two mechanisms, such as flower, pea, sheet, etc.^{99,107,109}

4. Developments and applications of heterostructures for nonaqueous Li–O₂ batteries

4.1. Carbon supported heterostructures

Because of their low cost, environmental friendliness, high surface area, superior electrical conductivity, easy preparation, robust mechanical strength and good physicochemical stability,^{110–112} carbon materials have been widely studied and used in Li–O₂ batteries. More importantly, they can also show excellent catalytic properties for boosting the ORR.^{113,114} Their OER catalytic activity, however, is evidently insufficient, largely limiting their practical Li–O₂ battery performance.^{116,117} Therefore, it is necessary to employ bifunctional cathode catalysts with excellent performance towards both the ORR and OER by constructing heterogeneous structures containing carbon



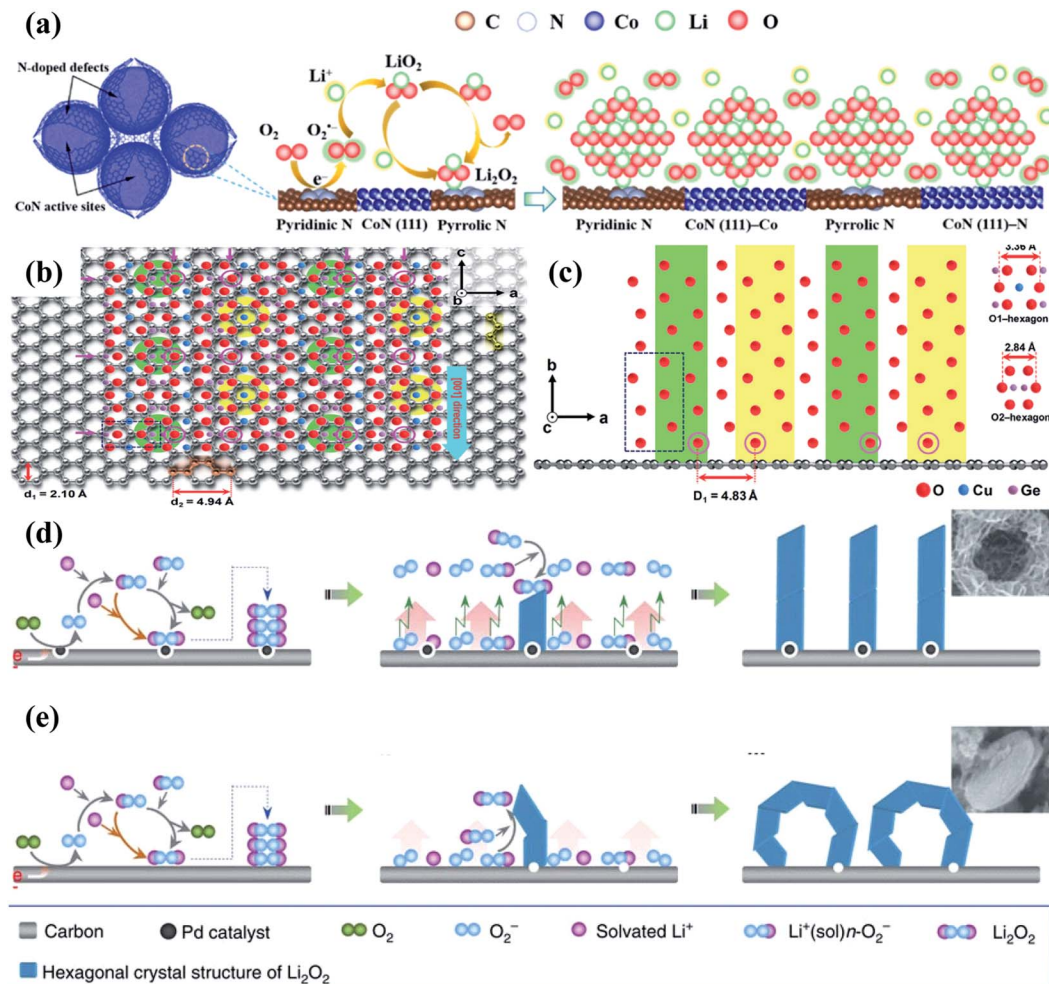


Fig. 4 (a) Schematic of the biphasic N-doped cobalt@graphene multiple capsule heterostructure and the proposed reaction mechanism during the discharge process. Reproduced with permission: copyright 2017, American Chemical Society.¹¹⁵ (b) Top and (c) side views from the nanowire-axis direction. Reproduced with permission: copyright 2018, Wiley.¹⁰⁰ Different formation mechanisms of Li_2O_2 aggregates on (d) P-HSC and (e) HSC cathodes. Reproduced with permission: copyright 2013, Springer Nature.⁵²

materials and active OER catalytic materials. Tan *et al.*¹¹⁵ prepared a dual-phase nitrogen-doped cobalt@graphene multi-capsule heterostructure *via* high-temperature synthesis and nitrogen and ammonia treatment, and Fig. 4a shows the electrocatalytic reaction processes of this heterostructure during cycling. N-doped graphene with a high surface area and excellent electrical conductivity functions as the porous and conductive support, on which the exposed Co and CoN nanoparticles and the N-doped defects together act as the active sites to enable strong binding interactions with Li_2O_2 . The cathode catalyst thus delivered superior electrochemical catalytic activities, showing a low overpotential of 0.9 V at 0.1 mA cm^{-2} , and there was almost no change in the charge and discharge overpotentials after 30 cycles with a fixed capacity of 1.0 mA h cm^{-2} . Many studies focusing on carbon supported heterostructures also demonstrate a similar conclusion, such as Co_4N nanorods anchored on CNF papers,⁸⁹ Co_9S_8 nanorods on the porous carbon foil,¹¹⁸ $(\text{Mn}_{1/3}\text{Co}_{2/3})\text{O}$ on CNT microspheres,¹¹⁹ *etc.*

Notably, carbon based materials are proven to tightly anchor the active catalyst on their surfaces, beneficial for improving the

specific capacities and cycling stability of Li– O_2 batteries.^{124,125} For example, Lee *et al.*¹⁰⁰ successfully controlled the growth direction of 1D CuGeO_3 nanowires with the specific crystal orientation on 2D conductive graphene by a facile hydrothermal route followed by thermal reduction. It is found that the lattice mismatch between CuGeO_3 and graphene is only 2.2% (Fig. 4b and c, $d_2 = 4.94 \text{ \AA}$, $D_1 = 4.83 \text{ \AA}$), and the angles between them are 60 and 120°. This is because two hexagons (green and yellow circles), as shown in Fig. 4b, composed of oxygen atoms intersected in a zigzag pattern along the *a*-axis and *c*-axis directions of graphene, forming a zigzag line of oxygen atoms along the *c*-axis direction. Compared with individual graphene and CuGeO_3 cathodes, the CuGeO_3 -graphene cathode shows better cycle stability of 100 cycles at 200 mA g^{-1} with a specific capacity limit of 1000 mA h g^{-1} . The excellent catalytic performance of the CuGeO_3 -graphene heterostructured catalyst can be attributed to the bifunctional phase boundary synergy effect.

Besides, applying a heterogeneous structure in a cathode catalyst for the Li– O_2 batteries can also effectively modulate the growth and morphology evolution of the discharge products,



thereby largely reducing the OER overpotentials and enhancing the cycling performance. Xu *et al.*⁵² successfully synthesized a free-standing honeycomb-like palladium-modified hollow spherical carbon (P-HSC) deposited onto carbon paper through electrophoresis with template replication. The discharge products exhibit a unique architecture composed of Li_2O_2 nanosheets, and their relative formation mechanisms are depicted in Fig. 4d and e. Compared with the carbon surfaces, those of the Pd nanocrystals are less sticky to superoxide, which would weaken the binding interaction of superoxide and the carbon substrate. In response, superoxide molecules could easily diffuse away from the cathode. Benefiting from the mutual effect of spherical carbon and Pd nanoparticles, the nucleation and crystallization of Li_2O_2 preferentially occurs on prismatic crystal faces, resulting in nanosheet shaped Li_2O_2 growth. Kwak and co-workers¹²⁷ dispersed molybdenum carbide particles on carbon nanotubes by ball milling and thermal treatment, and their catalytic properties were investigated in Li–O₂ cells. It is worth noting that toroidal Li_2O_2 was formed on the carbon nanotube cathode in the discharge processes, while Li_2O_2 in the form of a well-dispersed layer covered the catalyst surfaces, which lead to a larger battery efficiency (88%) and better reversibility (over 100 cycles) with lower overpotentials.

In general, owing to multiple merits, it is a good choice to select carbon-based substrates for forming heterostructures. These highly conductive carbonaceous substrates with a large surface area can not only easily enable rapid electron transport and mass transfer, which aid in reducing the ORR and OER overpotentials, but also regulate the morphology of the discharge products, further improving Li–O₂ battery performance.

4.2. Noble metal based heterostructures

Among various catalysts, noble metals have been intensively investigated and widely testified to feature superior catalytically active properties Li–O₂ batteries. Due to their limited amount on earth and high cost, however, researchers have to look for alternatives or conduct structural modification of noble metals. Recent research studies show that constructing noble metal based heterostructures can not only greatly improve the catalytic performance of the cathode catalysts, but also make them more affordable for boosting practical applications of Li–O₂ batteries. Liang *et al.*¹²⁰ prepared a surface sulfur-vacancy modified Ru/ZnIn₂S₄ (RuZIS-Vs) Mott–Schottky heterojunction by hydrothermal and subsequent chemical reactions and used it as a cathode catalyst material for Li–O₂ batteries, showing low ORR and OER overpotentials (0.39/0.38 V) and excellent cycling stability (321 cycles). They found that sulfur vacancies induced the interface defects of ZnIn₂S₄, which matched well with doped Ru, thereby establishing typical metal/semiconductor Mott–Schottky interfaces. By comparing the electronic structures of ZIS, ZIS-Vs and Ru-ZIS-Vs, it is clear from Fig. 5a and b that the Mott–Schottky interfaces could effectively reduce the bandgap of ZIS by varying its valence and conduction band values, which would greatly promote charge transfer on the heterostructure and advance the ORR and OER catalytic activity. Similarly,

Agyeman *et al.*¹²¹ decorated Pd nanoparticles on NiCo₂O₄ nanosheets (Pd@NiCo₂O₄) *via* a simple hydrothermal and subsequent reduction processing, and their discharge products show a flower-like morphology. It is demonstrated that Pd@NiCo₂O₄ and NiCo₂O₄ exhibit different electronic structures caused by Pd doping and increased the oxygen vacancies in the heterostructure. As evidenced in Fig. 5c, it is mainly due to the fact that the electronegativity of O (3.44) is much higher than those of Pd (2.2), Co (1.88) and Ni (1.91), thereby attracting electrons from these metals into the O sites; notably, noble metal-based heterostructures could also contribute to accelerating the decomposition of Li_2O_2 . For instance, Huang and co-workers¹²⁸ prepared a core–shell Pd@Pd₄S heterostructure decorated on a porous carbon matrix. It is evident that the charging voltage of the Pd@Pd₄S–C cathode remains below 3.7 V with low overpotentials, which is much lower than those of other counterparts. In fact, during the charging process, the decomposition of Li_2O_2 was mainly divided into two parts, including decomposition of small Li_2O_2 particles to transform into $\text{Li}_{2-x}\text{O}_2$ through delithiation and subsequent decomposition of $\text{Li}_{2-x}\text{O}_2$ with the release of O₂ at higher oxidation voltages. The stable charging voltage of about 3.7 V for the charge profile of the Pd@Pd₄S–C cathode indicates that the Pd@Pd₄S heterostructure can effectively promote Li_2O_2 delithiation and largely reduce OER overpotentials to achieve high battery efficiency. In fact, instead of directly accelerating the decomposition process of Li_2O_2 , it is more common that noble metal-based heterostructures could induce the formation of a more easily decomposed Li_2O_2 structure during the ORR process. Cong's group¹²² applied a simple one-step electrospinning method to obtain a silver-modified perovskite La_{0.9}FeO_{3- δ} (Ag@LFO) as an effective cathode catalyst for Li–O₂ batteries. As shown in Fig. 5d, the binding energies of O₂ species on the surfaces of LFO (O₂ is –2.84 eV and LiO₂ is –6.01 eV) are much lower than those on Ag@LFO, indicating that they are more easily adsorbed on the Ag@LFO surfaces. It is also illustrated that the discharge product generation would prefer to follow the surface model on the Ag@LFO cathode, presenting the reason for the formation of thin-film Li_2O_2 on the surfaces of Ag@LFO and toroid Li_2O_2 on the surfaces of LFO. Actually, film-like Li_2O_2 is considered to be easier to decompose during charging, resulting in lower overpotentials and longer cycle life.

The activity of precious metal-based heterostructured catalytic materials is usually not as high as that of the precious metal itself. In recent years, to reduce costs, precious metals are often coupled with other materials in the form of nanosized structures, such as nanowires, nanoparticles, and nanoclusters. To fully exploit the catalytic properties of precious metal atoms, the application of precious metal single-atom catalysts has emerged as one of the most promising directions. Recently, Lu *et al.*¹²⁹ successfully synthesized ultra-high-density single-atom catalysts containing metal contents up to 23 wt% for 15 metals (including Pd, Ru, Ir, and Pt) on chemically distinct carriers *via* a versatile approach combining impregnation and two-step annealing, and this approach could enable kilogram-scale material output, boosting the commercialization of precious metal-based single-atom catalysts.



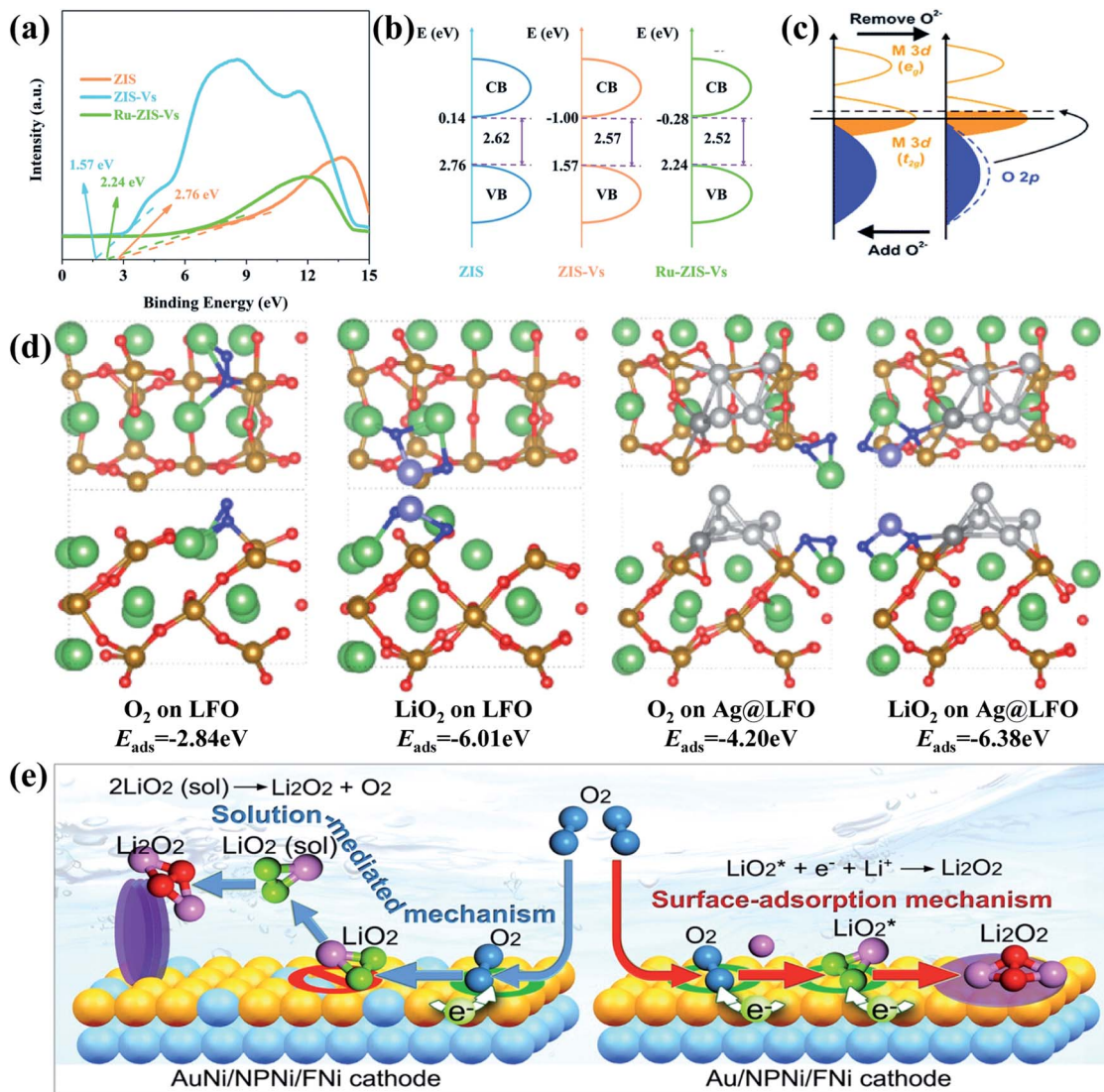


Fig. 5 (a) XPS valence band spectra and (b) band diagram of ZIS, ZIS-Vs and RuZIS-Vs. Reproduced with permission: copyright 2020, Royal Society of Chemistry.¹²⁰ (c) Molecular orbital structure change of Pd@NiCo₂O₄ as a result of Pd impregnation. Reproduced with permission: copyright 2017, Royal Society of Chemistry.¹²¹ (d) Optimized structures and the corresponding binding energies of O₂ and LiO₂ on LFO and Ag@LFO. Color codes: Li (violet), O (red), Fe (golden), Ag (light gray for Ag clusters and gray for replacing lanthanum in perovskite), lanthanum (green) and oxygen atom of O₂ and LiO₂ (blue). Reproduced with permission: copyright 2019, American Chemical Society.¹²² (e) The mechanisms of electrochemical growth of the film-like and nanosheet-like Li₂O₂. Reproduced with permission: copyright 2017, American Chemical Society.¹²³

4.3. Transition metal oxide based heterostructures

Owing to affordable cost, wide structural diversity, multiple valence states and relatively high electrocatalytic activity, transition metal oxides are considered as one of the alternative materials to noble metals as cathode catalysts for Li-O₂ batteries. Their electrical conductivity, however, is normally too limited to enable fast reaction kinetics,¹³⁰⁻¹³² and it is highly necessary to boost charge transfer and O₂ adsorption on the surfaces for perfecting battery performance. Applying transition metal oxide based heterostructures as catalysts is proven to be one of the most effective strategies. Wu's group¹²⁶ successfully prepared Mo₂C/MoO₂@ reduced graphene oxide heterostructures by *in situ* decoration of Mo₂C between MoO₂ and

reduced graphene oxide through a hydrothermal-calcination method. The atomic MoO₂ (-111), Mo₂C (101) and MoO₂/Mo₂C interface models are listed in Fig. 6a for revealing the O₂ adsorption, and the DFT calculation data demonstrate that the Mo₂C/MoO₂ interfaces show greater O₂ adsorption energy. This is due to the charge distribution imbalance around the interfaces caused by the accumulation of valence electrons, which could offer stable O₂ chemical adsorption on the Mo atoms. Nucleation instead of crystal growth was thus preferred during cycling, and film-like amorphous discharge products tended to form on the composite catalyst, resulting in an ultra-low potential gap of 0.57 V, and a stable cycling performance with a charge potential of less than 4.0 V after 100 cycles at 200 mA



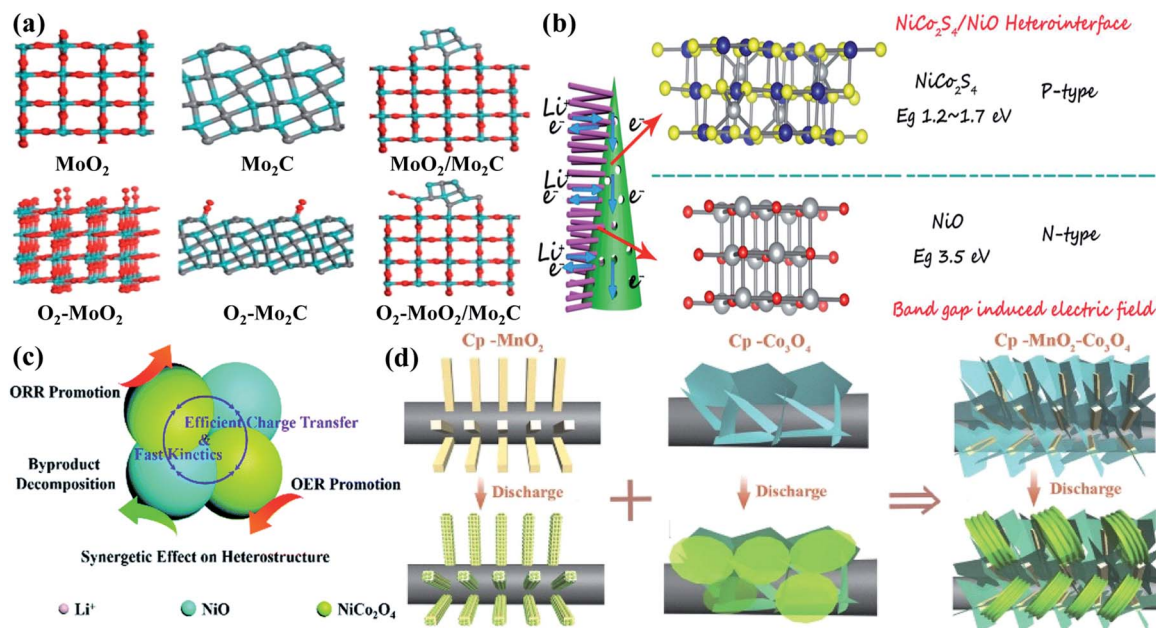


Fig. 6 (a) Views of MoO_2 (-111), Mo_2C (101) and $\text{MoO}_2/\text{Mo}_2\text{C}$ interfaces with corresponding O_2 adsorption models (red, cyan and grey balls are O, Mo, and C atoms). Reproduced with permission: copyright 2020, Elsevier.¹²⁶ (b) Summary of the enhanced high rate reaction mechanism with unique local electric fields from the $\text{NiCo}_2\text{S}_4/\text{NiO}$ heterostructure. For the spheres in the models (gray, blue, yellow and red balls represent Ni, Co, S and O atoms). Reproduced with permission: copyright 2019, Wiley.¹⁰⁷ (c) Specific roles of the $\text{NiO}-\text{NiCo}_2\text{O}_4$ heterostructure for $\text{Li}-\text{O}_2$ catalysis during discharge and recharge processes. Reproduced with permission: copyright 2019, Royal Society of Chemistry.¹⁰¹ (d) Schematic illustration of different discharge product morphologies on carbon paper supported MnO_2 and Co_3O_4 cathodes. Reproduced with permission: copyright 2017, Wiley.⁹⁹

g^{-1} . Wang *et al.*¹⁰⁷ reported, as shown in Fig. 6b, that the p-n heterojunction could be constructed from NiCo_2S_4 nanotubes and NiO nanoneedles with distinctly different bandgaps. This unique structure could introduce a built-in electric field at the hetero-interfaces, effectively facilitating interfacial charge transfer kinetics and fast ORR and OER kinetics. Boosted charge transfer could also be traced in $\text{PdO}-\text{Co}_3\text{O}_4$ heterostructures,¹³⁴ which delivered a stable cycling performance of more than 90 cycles. Moreover, the synergetic effect was realized in urchin-like $\text{NiO}-\text{NiCo}_2\text{O}_4$ microspheres¹⁰¹ for perfecting their $\text{Li}-\text{O}_2$ battery performance. As depicted in Fig. 6c, NiCo_2O_4 was believed to facilitate both the ORR and OER for $\text{Li}-\text{O}_2$ batteries, and NiO was likely to boost the by-product decomposition. More importantly, fast transport of Li^+ and electrons could be enabled to further promote the reaction kinetics, and superior specific capacities with low overpotentials and enhanced cycling stability were thus obtained. It is worth noting that the electrochemical performance is usually closely related to the formation and evolution of discharge products,^{135,136} which could be governed by the heterostructures. Zhang and co-workers⁹⁹ deposited Co_3O_4 nanosheets and $\alpha\text{-MnO}_2$ nanorods on the carbon paper through a simple electro-deposition and calcination route. Compared with the granular morphology on the MnO_2 cathode and nanosheet/film structure on the Co_3O_4 cathode, mooncake/large-sheet Li_2O_2 was homogeneously (Fig. 6d) grown on the composite cathode with an embedded architecture. This is mainly because $\alpha\text{-MnO}_2$ provided the growing sites, and Co_3O_4 nanosheets with evidently increased

height further facilitated the formation of dissolved LiO_2 . Unique embedded growth of Li_2O_2 was thus realized, bringing about a large reversible specific capacity of 4850 mA h g^{-1} with almost 100% coulombic efficiency, and a low potential gap of 0.95 V at 103 mA g^{-1} . 1-D $\text{RuO}_2/\text{Mn}_2\text{O}_3$ hollow heterostructures were also fabricated through single-nozzle electrospinning followed by heating speed controlled annealing.¹⁴⁰ Interestingly, their double-walled fiber structure could effectively trigger Li_2O_2 generation and decomposition inside and outside on the catalysts with close contact, displaying improved overpotential characteristics and stable cycle life of over 100 cycles in the $\text{Li}-\text{O}_2$ cells.

Although the TMO based heterostructures exhibit high redox activity in charge-discharge reactions, the overpotential depends strongly on the composite phase, and the heterostructures obtained from the combination of two transition metal oxides normally show high overpotentials, which are caused by the low electrical conductivity limiting the charge transport. The composite of transition metal oxides and high electrical conductivity materials can therefore effectively minimize the obstacles and maximize the activity of the catalytic sites.

4.4. Transition metal dichalcogenide based heterostructures

In recent years, transition metal dichalcogenides (TMDs), the chemical formula of which appears as MX_2 (M and X represent the transition metal and chalcogen elements, respectively),¹⁴¹ have received broad research interest. Most of them with



a unique X-M-X structure show typical 2D material properties and could expose more catalytically active sites.^{142,143} Besides, the weak van der Waals interaction and the missing of surface dangling bonds could effectively alleviate the strict thermal and lattice matching demand, contributing to easy heterostructure construction.¹⁴⁴ Furthermore, TMDs could display different electrical characteristics, which are mainly affected by the polype and the number of transition metal d electrons, and this could result in semiconducting or metallic behaviors,¹⁴⁵ favorable for forming efficient heterostructures with various other catalysts for Li-O₂ batteries. Hu *et al.*¹⁰⁸ developed heterostructured NiS₂/ZnIn₂S₄ nanosheets through a one-step hydrothermal method, whose density of electronic states close to the Fermi level at the heterogeneous interface increases significantly (Fig. 7a). On the one hand, the established built-in fields on the heterostructures are beneficial for fast interfacial charge transfer, providing numerous active interfaces for redox reaction kinetics. On the other hand, the heterostructures could induce strong Li⁺ and O₂ adsorption and weak LiO₂ adsorption (Fig. 7b), which boost LiO₂ desorption on their surface and

solvation in the electrolyte. This effectively promoted toroid-like Li₂O₂ deposition and decomposition in low Gutmann donor number (DN) solvents through the solution pathway, which guaranteed outstanding electrocatalytic activities. Similar findings also appeared in Ni₃Se₂/NiSe₂ heterostructures,¹³³ and it is also evident to realize advanced charge transfer by enabling distinct electron redistribution along the interfaces (Fig. 7c). Moreover, their 3D porous structure aided in barrier-free mass transport and urchin-like Li₂O₂ generation with intimate contact with the catalyst (Fig. 7d), leading to a large discharge specific capacity of 23 092 mA h g⁻¹ at 500 mA g⁻¹ and reduced charging overpotential of 0.38 V, and a long cycling life of more than 500 cycles can also be achieved with a fixed capacity of 1000 mA h g⁻¹ at 100 mA g⁻¹. What is more, Liang's group⁹⁵ conducted deliberate CoSe₂@NiSe₂ heterostructure engineering *via* the hydrothermal method combining heat treatment. As shown in Fig. 7e and f, the formation of a heterogeneous interface caused the distortion of local fine atom arrays. These distortions can not only serve as active sites, but also reduce the surface energies of the catalyst, thereby promoting the LiO₂ and

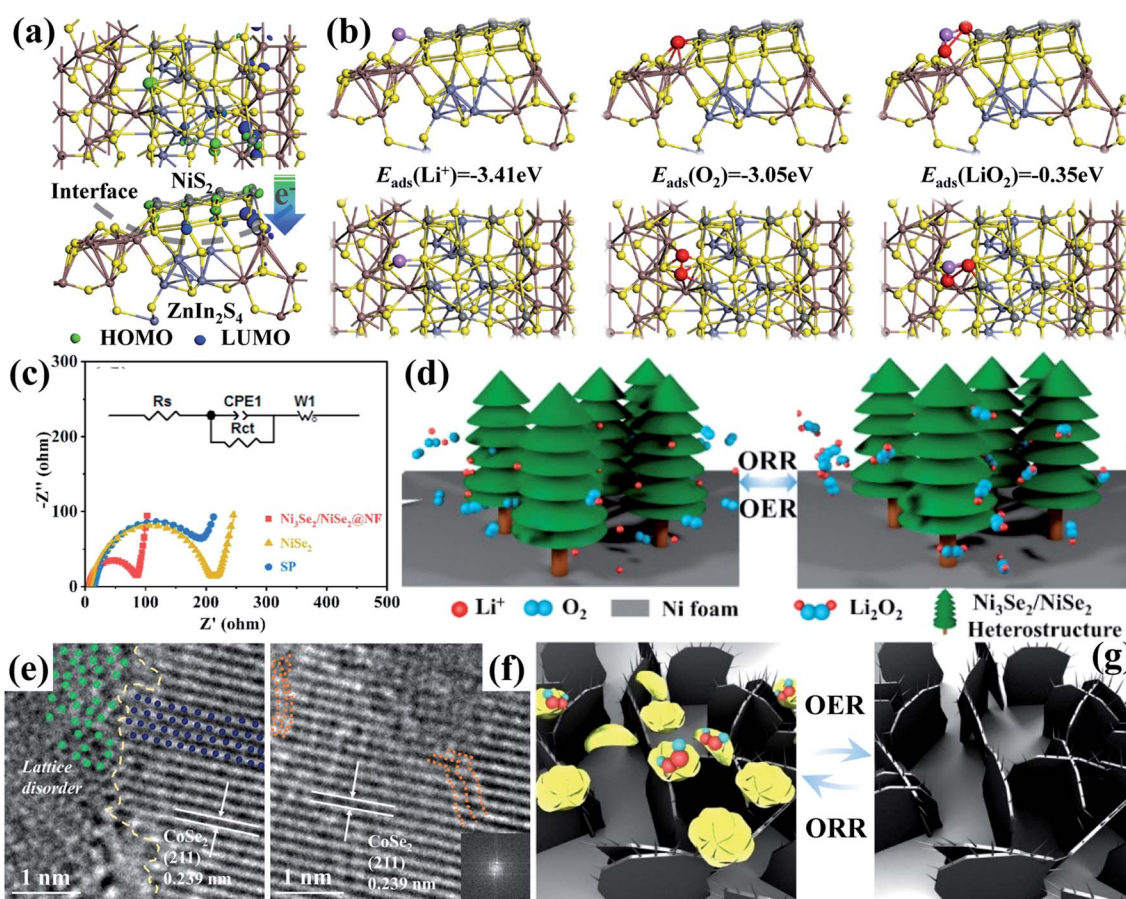


Fig. 7 (a) Charge distribution at the interface of NiS₂ and ZnIn₂S₄ with (b) adsorption energies (ΔE_{ads}) of Li⁺, O₂ and LiO₂ adsorbed on NiS₂@ZnIn₂S₄ with optimized structures. Reproduced with permission: copyright 2020, American Chemical Society.¹⁰⁸ (c) Nyquist plots of different cathodes with the inset showing the fitting equivalent circuits. (d) Schematic diagram of the reaction mechanisms of the Ni₃Se₂/NiSe₂@NF cathode during discharge and charge processes. Reproduced with permission: copyright 2020, Elsevier.¹³³ (e) HRTEM image of CoSe₂@NiSe₂ at the interface with the disordered edges. (f) HRTEM image of CoSe₂@NiSe₂ and inset shows the corresponding FFT pattern from the distortion region. (g) Schematic illustration of the oxygen electrode reactions on the CoSe₂@NiSe₂ cathode during cycling. Reproduced with permission: copyright 2020, Elsevier.⁹⁵



adsorption and reversible flower-shaped Li_2O_2 formation (Fig. 7g) with a low potential gap (1.2 V) during cycling.

TMD based heterostructures show a high initial capacity and excellent cycling stability. To achieve high ion and electron transfer rates, the compositing of metallic phases for TMDs seems to be a favorable option. However, the problem of the conversion of pure metallic phases to semiconductor phases during charging and discharging has not yet been solved, and the combination of the two phases to obtain a heterogeneous structure has been conducted as a possible strategy in some studies,^{44,146} displaying the advantages of both high electrical conductivity and high stability.

4.5. Other heterostructures

In addition to the above heterostructures, there are also some other heterostructures applied in Li–O₂ batteries. For example, Jin *et al.*¹³⁷ synthesized Co and Fe layered double hydroxide (LDH)–RuO₂ nanosheet heterostructures by the self-assembly technique (Fig. 8a). The strong interface electronic coupling between LDH and RuO₂ nanosheets with oxygen vacancies greatly improved the diffusion of Li⁺ and reactants, which are responsible for the enhanced transfer kinetics towards the OER and ORR. Lee's group¹³⁸ simulated the ORR pathway of the heterostructure of h-BN and Ni (111) as a cathode catalyst through DFT calculations; the structure of the heterostructure is shown in Fig. 8b, and they found that the heterostructure

thermodynamically preferred to undergo the $2e^-$ pathway (Fig. 8c), which is obvious different from most of the previous results with the $4e^-$ pathway. The main reason is that the $(\text{Li}_2\text{O}_2)_2$ adsorption is remarkably strong on the h-BN/Ni (111) surface, contributing to a maximum discharge potential of 1.93 V and a minimum charge potential of 3.83 V, comparable to those of noble metal based catalysts. In recent years, photocatalysis has also been developed rapidly and applied to the field of Li–O₂ batteries. Veeramani and coworkers¹³⁹ developed a photocathode nanocomposite of cadmium selenide/zinc sulfide quantum dots grown on carbon nanotubes (CdSe/ZnS QD@CNT), and it shows a low charge voltage within 4.0 V; this is because during charging, the photoexcited holes created by the photoelectrode absorbing photons from the light source on the heterostructure interfaces can assist to oxidize Li_2O_2 to oxygen gas and Li⁺ in the valence band (VB) position (Fig. 8d).

The rapid development of heterogeneous structures based on the stacking of two-dimensional materials (including graphene, hBN, MoS₂, hydroxides, and MXenes) in recent years has also led to new directions in the field of energy storage and conversion,^{147–149} which offers the opportunity to build electrodes that will combine the advantages of each building block and eliminate the associated disadvantages. For example, hBN crystals have proved to be a similar substrate to graphene, allowing a tenfold increase in their electronic quality.

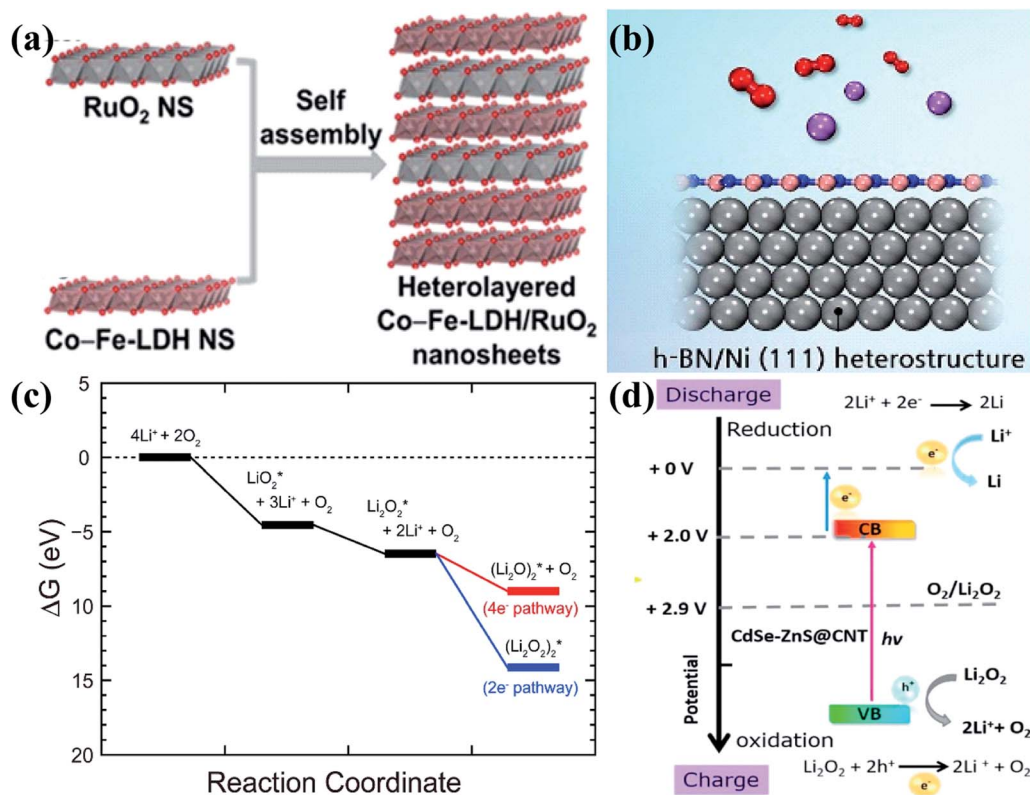


Fig. 8 (a) Schematic illustration of the synthesis of Co and Fe layered double hydroxide (LDH)–RuO₂ nanosheets. Reproduced with permission: copyright 2019, Elsevier.¹³⁷ (b) Schematic illustration of the h-BN/(111) heterostructure. (c) Free energy diagram of the ORR on the h-BN/Ni (111) surface. Reproduced with permission: copyright 2016, Elsevier.¹³⁸ (d) The theoretical energy band diagram for the improved performance of rechargeable photoelectrochemical Li–O₂ batteries. Reproduced with permission: copyright 2018, Elsevier.¹³⁹



5. Summary and perspectives

In order to cope with the energy crisis, the development of Li–O₂ batteries has attracted widespread attention from researchers, and it is essential to develop high-performance catalysts with high activity and long-term durability. In this review, we summarized the latest research progress of heterostructured catalysts used in Li–O₂ batteries, and various strategies for battery performance improvement were proposed. Moreover, analysis of the mechanisms behind performance enhancement and the functions of hetero-interfaces for Li–O₂ catalysis were emphasized, and the existing problems of the current heterostructure research studies and possible directions for future studies were finally presented, as depicted in Fig. 9.

1. The critical roles of heterostructures for Li–O₂ catalysis are mainly played in the following aspects: (1) the transportation of electrons and ions could be accelerated. Since the difference

between the Fermi levels of the heterostructure components normally induces a built-in electric field, the migration of ions and electrons would be effectively promoted under the action of coulombic force to advance the reaction kinetics. To this end, it is believed that gradient hetero-interfaces with an ideal built-in interface potential should be induced for fast charge and mass transfer, such as multi-layer heterostructures, and even the construction of the built-in electric field inside the components through precise tuning of the composition of each component needs to be considered. It is worth noting that these strategies can also bring about new insights of designing high-current energy storage devices; (2) synergy effects are proposed to contribute to the excellent bifunctional catalytic activities of heterostructures, which refers to the outcomes of $1 + 1 > 2$ under the actions of different components. In fact, the deeper intrinsic mechanisms for synergy effects have not been fully explored, and the as-presented synergy effects were comprehensive

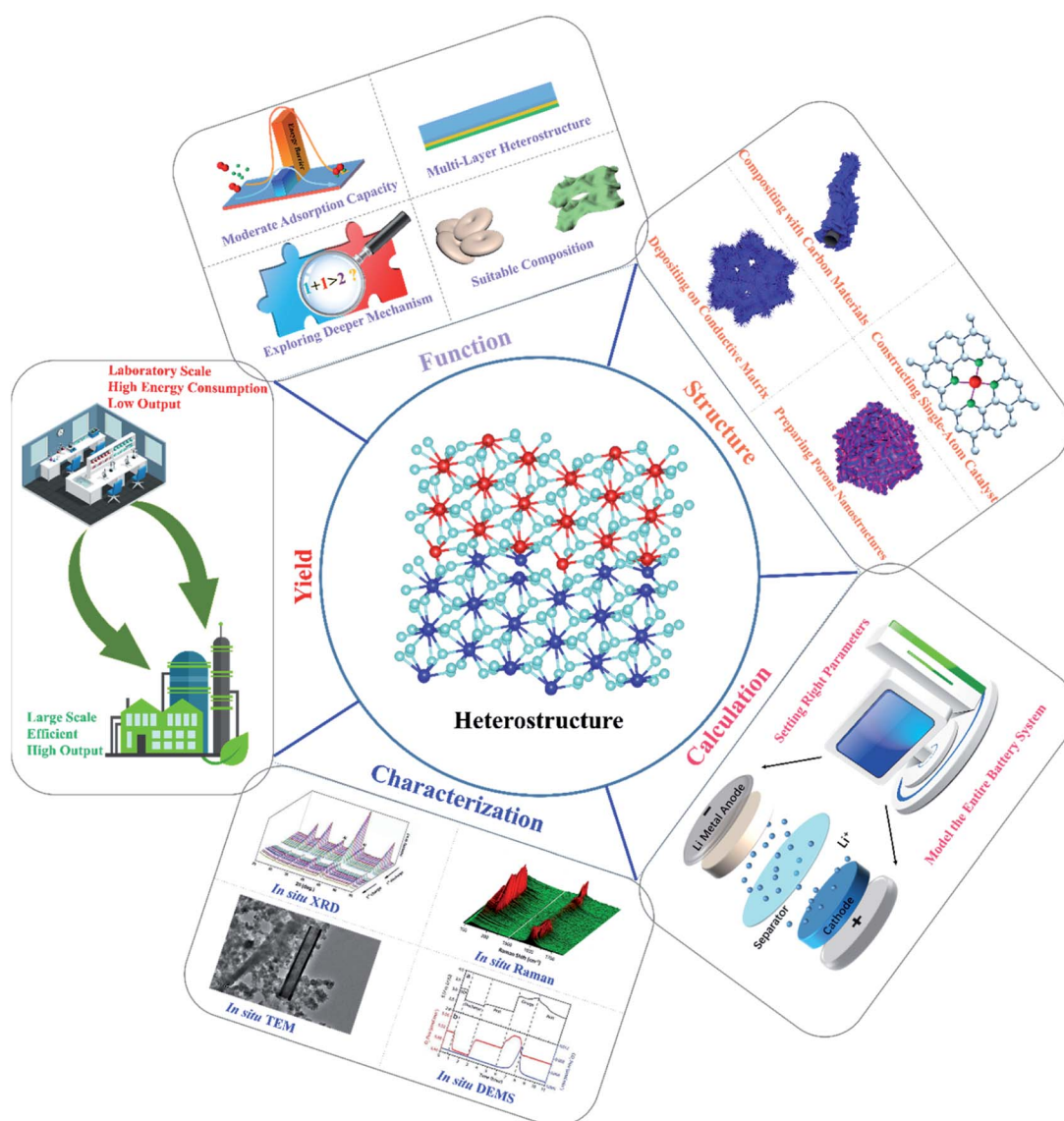


Fig. 9 Directions and perspectives for future studies on heterostructured cathode catalysts for Li–O₂ batteries.



macroscopic properties largely affected by different components. Consequently, clearer evidence should be provided to illustrate the insightful effect of synergy in future research studies; (3) the adsorption behaviors of the catalysts are basically controlled by the electronic properties of the active sites, and regulating the adsorption energies of O_2 species on heterostructures is thus essential for improving the Li- O_2 battery performance. It is found that the electronic structures of the surrounding active sites on heterostructures could be effectively tuned, thereby changing the adsorption energies of O_2 species and resulting in the alterations of ORR and OER pathways. Actually, too low or too high adsorption energies of O_2 species tend to pose difficult adsorption or desorption during cycling, which demonstrates that the ideal heterostructured catalysts should show a moderate adsorption capacity to deliver the smallest reaction barrier in the adsorption and desorption processes. (4) Each component in the heterostructures affects the nucleation and growth of the discharge products *via* different formation mechanisms, and there is definitely a competition between them. There are currently more than 100 candidates for Li- O_2 battery cathode catalysts (including graphene, molybdenum sulfide, Pt, *etc.*), and more than 10 000 hetero structures composing them. How to select and use them to build heterostructures still remains challenging, and employing computational material pre-selection to build a genome for heterostructures for Li- O_2 batteries is a good way to solve this problem. As a result of these research efforts, controlled access to small and readily decomposable Li_2O_2 or even highly conductive amorphous Li_2O_2 is possible, thus greatly extending the limits of current Li- O_2 batteries.

2. Fig. 9 also shows the strategies for constructing high-performance carbon, noble-metal, and TMO and TMD-based heterostructures. Carbon-based heterostructures are favored by researchers as catalysts, mainly because of their low price and high electrical conductivity. However, in view of the fact that carbon materials themselves are vulnerable to superoxide attack and enable irreversible decomposition, the rational designs of carbon-based heterostructures are focused on composing heterostructures on the supported carbon substances, which can not only utilize the high electronic conductivity and large specific surface area of the carbon matrices, but also avoid the formation and accumulation of by-products to prevent the rapid death of Li- O_2 batteries. In contrast, noble metal-based heterostructures usually deliver low overpotentials and stable cycle performance, and the purpose of preparing such heterostructures is to reduce the amount of noble metals in the catalysts with high catalytic performance close to those of noble metal catalysts. In this respect, guaranteeing uniform dispersion of noble metals throughout another component is of great significance, which is still a problem remained to be solved. Moreover, the rapid development of single-atom catalysts in recent years has paved the road for the embedding of noble metal atoms into the carbon matrix or other compounds. These heterostructures tend to feature superior electrochemical catalytic behaviors to those of traditional heterostructures containing nanoparticles with limited utilization of noble metals. Furthermore, directly depositing catalysts onto the conductive

current collectors or gas diffusion layers is an effective way to enhance the electrical conductivities and restrain the side reactions of heterostructure cathodes, including carbon paper, nickel, titanium foams or meshes. TMD-based heterostructures are still less researched in Li- O_2 batteries, but there has been a lot of progress in the field of HER catalysis and supercapacitors, which can bring some experience to the development of Li- O_2 batteries. The main disadvantage of TMD materials is low conductivity as well as low specific surface area; the problem of low specific surface area can be solved by preparing porous nanostructures (nanoflower and hollow nanotube), and the low electrical conductivity can be enhanced by constructing heterostructures. The other component of the heterostructure should be carefully selected, such as high conductive composite (metal/carbon material/superconducting TMD), or phase modulation is recommended to obtain a mixed phase heterostructure (1T/2H) with improved electrical conductivity. In addition to the design of materials in terms of the function and category of the heterostructures, the dimensionality of the heterostructured material also has a significant impact on the electrochemical performance, and most heterostructured materials in Li- O_2 batteries are currently of multiple dimensions; most of them are combinations of 3D + nD ($n = 0, 1$ and 3) materials. The large specific surface area of 1D or 2D materials increases the contact area with the electrolyte and improves the kinetics of ion transfer, in addition to the fact that they typically have a large number of exposed metal active sites and thus exhibit greater specific capacities. By combining the high electrical conductivity of single-walled carbon nanotubes, graphene, or MXene with TMOs or TMDs holding favorable redox activity, high catalytic performance can be achieved when individual shells or layers are assembled into stacked 1D or 2D heterogeneous structures.

3. At present, most research studies on the reaction mechanisms were carried out by means of theoretical calculations to obtain the reaction energy barrier, electronic state density distribution, charge distribution, with simulating the conversion kinetics of the corresponding device combined with the experimental results. It is essential to comprehend the mechanism of the catalytic process by theoretical calculations, while the parameter settings in the calculation processes determine the accuracy of the theoretical calculation results. The differences between the theoretical research models and the actual catalyst statements may lead to discrepancies, leading to inaccurate conclusions. Notably, the theoretical calculations are mainly based on the adsorption conditions and electrocatalytic reactions on the catalyst surfaces, but there are a lot of other factors that should also be involved, such as the dissolution of lithium in the anode, the heat transfer between the electrolyte and the electrodes, and even the changes in the electrolyte flow. Therefore, the entire Li- O_2 battery system with relative influencing factors should be taken into consideration, and theoretical models and parameters need to be set as close as possible to actual conditions, reasonably offering the guidance for catalyst design and performance improvement.

4. In-depth and systematic characterization studies are highly demanded to intensively investigate Li- O_2 battery reactions based on heterostructured catalysts, and typical advanced



characterization techniques are listed in Fig. 9. *In situ* Raman could detect the changes in the types and contents of the deposited substances on the electrode surface during charge and discharge. *In situ* infrared is used to constantly detect the intermediate products of the catalytic cycle and the adsorption state of the real reaction. *In situ* XRD can qualitatively and quantitatively analyze the crystallization reaction product Li_2O_2 and other crystalline by-products. *In situ* TEM can also detect the change of the product structure to reveal the detailed nano-level mechanism at the atomic resolution and visualize the ORR and OER processes. *In situ* DEMS is used to analyze the gas consumption and emissions during battery cycling. In particular, these *in situ* technologies can help us determine the reaction pathway and the growth process of Li_2O_2 in the ORR and OER processes of Li–O₂ batteries based on heterostructured catalysts, which can also be better compared with theoretical calculations.

5. Although heterostructures show great potential in Li–O₂ batteries, some of the preparation procedures for the heterostructured catalysts consist of multiple steps, which largely affect the controllable modulation of the morphology of the materials. As shown in Table S1,† the synthesis routes reported in this review are based on the lab conditions rather than laboratory-scale production, and some of these methods require high energy consumption and cumbersome fabrication processes with low output (Fig. 9). Therefore, the development of green, mild and efficient preparation strategies is highly desired, and plays a vital role in realizing large-scale commercial applications of bi-functional electrocatalysts for Li–O₂ batteries. In addition, the active centers of the homogeneous structural material with a single phase are assumed to be uniformly dispersed on the entire catalyst phases, while the heterostructure composed of multiple phases normally offers heterogeneous interfaces, making it more difficult to clarify the relative catalytic mechanism.

Author contributions

Jun Wang, Yao Liu, and Shulei Chou contributed to the conception of the study; Qing Xia contributed to the analysis and manuscript preparation; Deyuan Li, Lanling Zhao, Yuxin Long, Xue Han, Zhaorui Zhou, Yiming Zhang, Yebing Li and Abulgasim Ahmed Abbaker Adam performed the analysis with discussions.

Conflicts of interest

There are no conflicts to declare.

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