

## REVIEW

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3, 7384Ion-irradiation of catalyst and electrode materials  
for water electrolysis/photoelectrolysis cells,  
rechargeable batteries, and supercapacitorsYahui Wang, <sup>a</sup> Xing Cheng, <sup>a</sup> Kai Zhang, <sup>b</sup> Ge Chen, <sup>\*a</sup> Ruzhi Wang <sup>\*c</sup>  
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Innovatively designed, modified, and synthesized catalyst/electrode materials are of great importance in improving the performance of electrochemical energy devices for energy storage and conversion. Among various material modification technologies, the ion-irradiation technique has been recognized as a promising technology, which can effectively modify and design materials through controlling the key irradiation parameters, such as irradiation ion species, energy and fluence. The advantages of the ion-irradiation technique are high precision, favorable controllability, favorable repeatability, etc.; meanwhile, it can realize arbitrary element doping and defect control for many catalyst/electrode materials. Notably, it does not induce any impurities into the target materials during the ion-irradiation process, which is superior to many chemical modifications. The effects of ion-irradiation on catalyst/electrode materials in recent years are described in this review, including defect introduction, regulation of the electronic structure, morphology control, synthesis, and element doping. Then, the applications of ion irradiated catalyst/electrode materials in water electrolysis/photoelectrolysis cells, lithium-ion batteries, and supercapacitors are systematically summarized with an emphasis on the advantages of boosting the properties of materials. Finally, the challenges and strategies are proposed for developing more practical ion-irradiation techniques towards high-performance catalyst/electrode materials.

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## Introduction

In an effort to decrease the depletion rate of fossil fuels and reduce the associated environmental impact, exploring clean and sustainable energy sources such as solar, wind, waterfall, *etc.* for generating clean electrical energy has become imperative in recent years. However, such electricity is intermittent, and highly efficient energy storage and conversion technologies are needed to make them suitable for many applications.<sup>1,2</sup> In this regard, electrochemical energy devices such as rechargeable batteries, supercapacitors, and water electrolysis cells have been recognized as the reliable, efficient, and practical means to meet the renewable energy storage and conversion requirement. Achieving higher energy/power densities, lifetime, and safety of these devices are always the relentless pursuit,<sup>3,4</sup> and the development of highly efficient catalyst/electrode materials has been the major focus, which can be approached through innovative design and fabrication as well as modification for the device performance optimization.<sup>5–7</sup> Particularly, elaborately modified catalyst/electrode materials are believed to be one of the most efficient ways of obtaining high performance.<sup>8,9</sup>

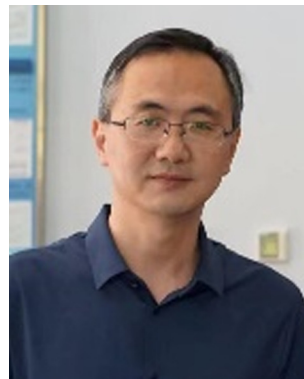
Among various modification strategies, ion-irradiation has been identified as an environmentally friendly technique that can modify the material without using any chemical reagents. When the energy of ion irradiation is large enough, it is able to transfer considerable energy and momentum to a single atom or an electron. This is also the main reason why ion-irradiation can adjust the properties of various materials to a considerable extent. During the irradiation process, energetic ions hit the target material and make the impacted target nuclei and electrons generate defects (*e.g.* vacancy dislocations), which result in changes in the physical and chemical properties of the materials.<sup>10,11</sup> Ion irradiation can introduce nanocrystallization, ordered surface nanostructures, and latent tracks in the target materials. To date, ion-irradiation has been widely applied in a variety of fields such as solar cells,<sup>12</sup> biomedicine,<sup>13</sup> microelectronics<sup>14–16</sup> and renewable energy.<sup>17</sup>

In this review, we have mainly reviewed the application of ion-irradiation for catalyst/electrode materials in electrochemical energy conversion and storage devices, including water electrolysis/photoelectrolysis cells, rechargeable batteries, and supercapacitors. The ion-irradiation effects on the catalyst/electrode materials are mainly discussed in terms of defect introduction,



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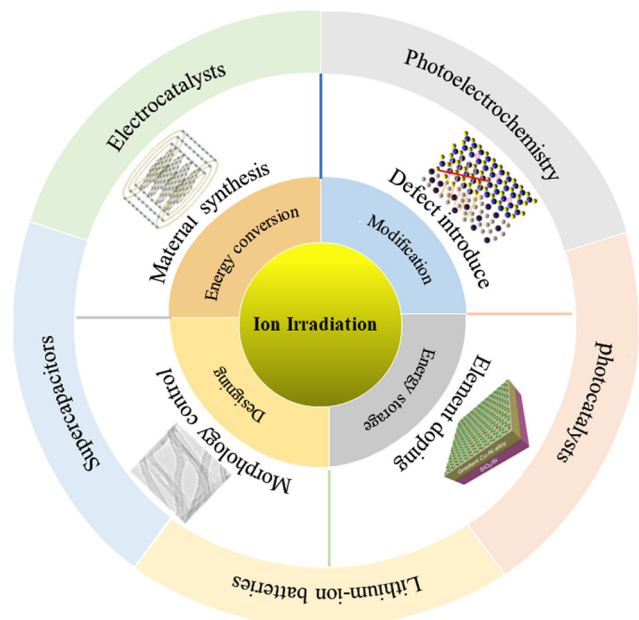


Fig. 1 Design considerations of the application of the ion-irradiation technique for modifying and designing catalyst/electrode materials for energy storage and conversion devices.

and element doping, morphology control and material synthesis. Particularly, we have emphasized ion-irradiation advantages for boosting the performances of the device. Fig. 1 shows the ion-irradiation of catalyst/electrode materials for various applications. Finally, we proposed the challenges and directions in the future which may provide some guidance for researchers in this field.

## 2. Fundamentals of the ion-irradiation technique

### 2.1 Concepts and principles of the ion-irradiation technique

Ion-irradiation is a technique that refers to ionizing the atoms of an element and then accelerating to a higher energy to make them possess a high kinetic energy. Then, they are injected into the surface of the target materials, resulting in a change in the physical or chemical properties of the material.

The principles of ion-irradiation are shown in Fig. 2.<sup>17</sup> First, the irradiated hot electrons from the ion source bombard the injected gas molecules for generating ions. Then, a magnetic analyzer is used to select specific elemental ions with mass-charge ratios. Subsequently, the ions are accelerated by a step through the accelerator to obtain a higher energy, and then focused through quadrupole lenses to obtain uniform beam spots. Finally, the ion beam is uniformly injected into the surface of the solid sample using a deflection scanner. The ions with high energy will collide with the atoms and electrons of the solid target surface many times, thus weakening the energy of the implanted ions. Eventually, ions penetrate into the solid material to achieve the purpose of irradiation. During irradiation, a series of physical and chemical interactions can occur between the implanted ions and the solid sample.

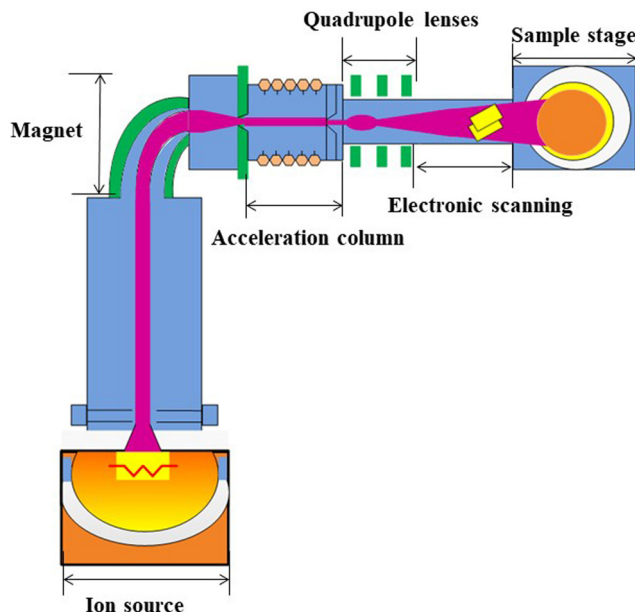


Fig. 2 Basic structure of an ion implanter with magnetic analysis. Reproduced with permission from ref. 17, Copyright 2020, AIP Publishing.

### 2.2 Composition of an ion implanter

The process of ion-irradiation on the sample is carried out using an ion implanter composed of an ion source, a magnetic analyzer, an accelerator, quadrupole lenses, electronic scanning, and a target chamber (Fig. 3).<sup>18</sup> The ion source is the main part of the ion-implantation implanter to produce ion beams with various strengths. Electrons produced by a direct current discharge or high-frequency discharge of an ion source act as the bombardment particles. When the energy of the foreign electron is higher than the ionization potential of the atom of targeted materials, the element is ionized by collision. A magnetic analyzer is used to select an elemental ion with a specific mass-charge ratio, and an accelerator is used to obtain higher energy of the ion-beam through a quadrupole lens.

### 2.3 Key parameters in the ion-irradiation process

For ion-irradiation, some irradiation parameters play a crucial role in modifying the properties of the targeted materials,<sup>19</sup> including irradiated energy ( $E$ ), ion fluence ( $N_I$ ), ion flux ( $j$ ) and implantation temperature ( $T$ ). The detailed explanations are as follows.

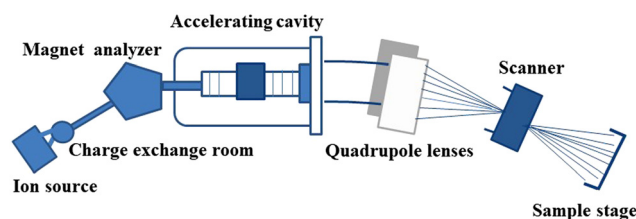


Fig. 3 Schematic diagram of the ion-irradiation principle. Reproduced with permission from ref. 18, Copyright 2017, Wiley-VCH.



(1) **Irradiated energy ( $E$ ).** The  $E$  determines the penetration depth of the implantation into the matrix. Typical values range from a few keV to several MeV, corresponding to implantation depths ranging from a few dozens of nanometers to a few microns. The higher the ion energy, the deeper the implantation depth. With the increasing energy, the average implantation depth will become deeper, and the depth distribution of the implanted species will become broader.

(2) **Ion fluence ( $N_I$ ).** The  $N_I$  is related to the number of solute monomers that are implanted into the matrix. The ion fluence ranges from a few  $10^{14}$  to several  $10^{17}$   $\text{cm}^{-2}$  in the implantation process.

(3) **Ion flux ( $j$ ).** The  $j$  is defined as the number of ions introduced into the unit surface area of the matrix per unit time. The formula is expressed as  $j = N_I/t$ , for electrode materials, which is measured in  $\mu\text{A cm}^{-2}$  or in ions per  $\text{cm}^2$  per s. The larger the ion flux that is provided, the shorter the implantation time needed.

(4) **Implantation temperature ( $T$ ).** Temperature affects the atomic mobility of the implanted monomers and the stability of the irradiation induced defects. Thus, it controls the evolution of the solid solution toward the nucleation and growth of the precipitates.

## 2.4 Type of ion-irradiation and characteristics

Different irradiated energies correspond to various applications. The following types of ion-irradiation are often used for the modification of catalyst/electrode materials: low energy ion-irradiation, medium energy ion-irradiation, swift heavy ion-irradiation (SHII) and high charge ion-irradiation.

The  $E$  of low energy ion-irradiation is usually less than 1 keV, which is widely used in materials as a common irradiation.<sup>20,21</sup> Low-energy ion irradiation can modify the surface morphology and achieve the selective doping of two-dimensional materials. The ion-implantation with an energy up to tens of keV can effectively realize the defect engineering of two-dimensional materials.<sup>22,23</sup>

The  $E$  of medium energy ion-irradiation ranges from a few hundred keV to a few MeV and is suitable for the micro/nanostructure modification and elemental doping of various thin film electrode materials.<sup>17,19</sup>

The  $E$  of SHII is usually more than 1 MeV. At higher energy, inelastic collisions occur between incident ions and target electrons. For inelastic collisions, energy loss can be transformed into defects through indirect complex processes. Since the transferred energy is very large, a large number of defects can be generated at a very low flux.<sup>24</sup> Therefore, a large number of defects can be produced at a very low fluence to engineer the defect structure and even the nano-morphology of the material.

High charge ion-irradiation is usually used in the modification of semiconductor materials. The high charge means that it possesses high potential energy, which is stored by removing many from the neutral atom. Potential energy is the sum of all binding energies of the lost electrons. It can reach ten to hundreds of keV.<sup>25,26</sup> Highly charged ions are slower, and the

target ion interactions are concentrated on the near-surface region.

Compared with other techniques, ion-irradiation possesses the following unique advantages:<sup>19</sup>

(1) **Favorable controllability.** In the irradiation process, there is a wide range of options and controllable implanted parameters (such as energy, fluency, and temperature). Therefore, ion-irradiation can precisely control the doping concentration and depth as well as longitudinal and transverse diffusion.

(2) **High precision.** Since the energy and fluence of implanted ions can be controlled, consequently, the depth and concentration of implanted ions in the host matrix can be accurately controlled during the implanted process.<sup>17</sup>

(3) **High purity.** Since the ion-irradiation is carried out in a vacuum system, it does not introduce other impurities. Meanwhile, a high resolution mass analyzer is used to ensure the high purity of the implanted ions.

(4) **Favorable repeatability.** Because the ion-irradiation technology is accurate, controllable, and accurately controls the parameters of the ion beam (species, energy, fluency, beam current, temperature, and so on),<sup>27</sup> consequently, it has favorable repeatability.

(5) **High flexibility.** Since every type of ion-implantation has its unique characteristics, therefore, it can be selected to satisfy the different purposes of specific modifications of materials which include a variety of properties of thin films, solids and nanostructured samples.

## 3. Ion-irradiation effects on the catalyst/electrode materials

In this section, we will focus on the ion-irradiation/implantation effects on the catalyst/electrode materials, including the defect introduction, element doping, morphology control, and material synthesis, and these effects are helpful in material modification and design.

### 3.1 Defect introduction

The electrochemical properties of the material are closely associated with defect structures. Introducing defects into materials is one of the promising approaches for improving the performance of materials. A great number of methods have been developed to introduce defects into materials, including plasma treatment,<sup>28</sup> thermal annealing,<sup>29</sup> chemical functionalization<sup>30</sup> and ion irradiation.<sup>31</sup> Among these methods, ion-irradiation can offer a wide range of controllability in the defect type by controlling the irradiation parameters such as the energy, type, and fluence of the injected ions.

Sun *et al.*<sup>32</sup> proposed using the C ion-irradiation to introduce sulfur vacancies to improve the hydrogen evolution reaction (HER) performance of  $\text{MoS}_2$ . They found that the irradiation dose plays a major role in tuning the number of sulphur vacancies and the amorphous phase on the basal plane of  $\text{MoS}_2$  which improves the HER performance of  $\text{MoS}_2$ . The superior activity for the HER is due to the vacancies generated by ion-irradiation, which



exposes more active sites and thus improves the catalytic performance.

Also, He *et al.*<sup>33</sup> fabricated a single-layer MoS<sub>2</sub> and followed this by using 500 keV Au irradiation to generate sulfur defects (Fig. 4a). They further investigated the impacts of defects on the properties of MoS<sub>2</sub>. As the defect density increases, the characteristic photoluminescence (PL) peak of MoS<sub>2</sub> was found to first shift to higher energy and then shift back to lower energy; after ion-irradiation, the electron transfers from MoS<sub>2</sub>

to the absorbed O<sub>2</sub> at the defect sites, thus leading to a depletion of the electrons and an enrichment of the holes.

Huang *et al.*<sup>34</sup> used a chemical vapor deposition (CVD) method to directly grow ReS<sub>2</sub> nanosheets on a p-Si substrate, which was further irradiated with Ar ion-irradiation (Fig. 4b). Re defects were successfully introduced onto the ReS<sub>2</sub>/Si photocathode by ion-irradiation. The existence of Re vacancies not only produces adequate unsaturated S atoms but also influences intrinsic charge compensation from S to Re-Re bonds,

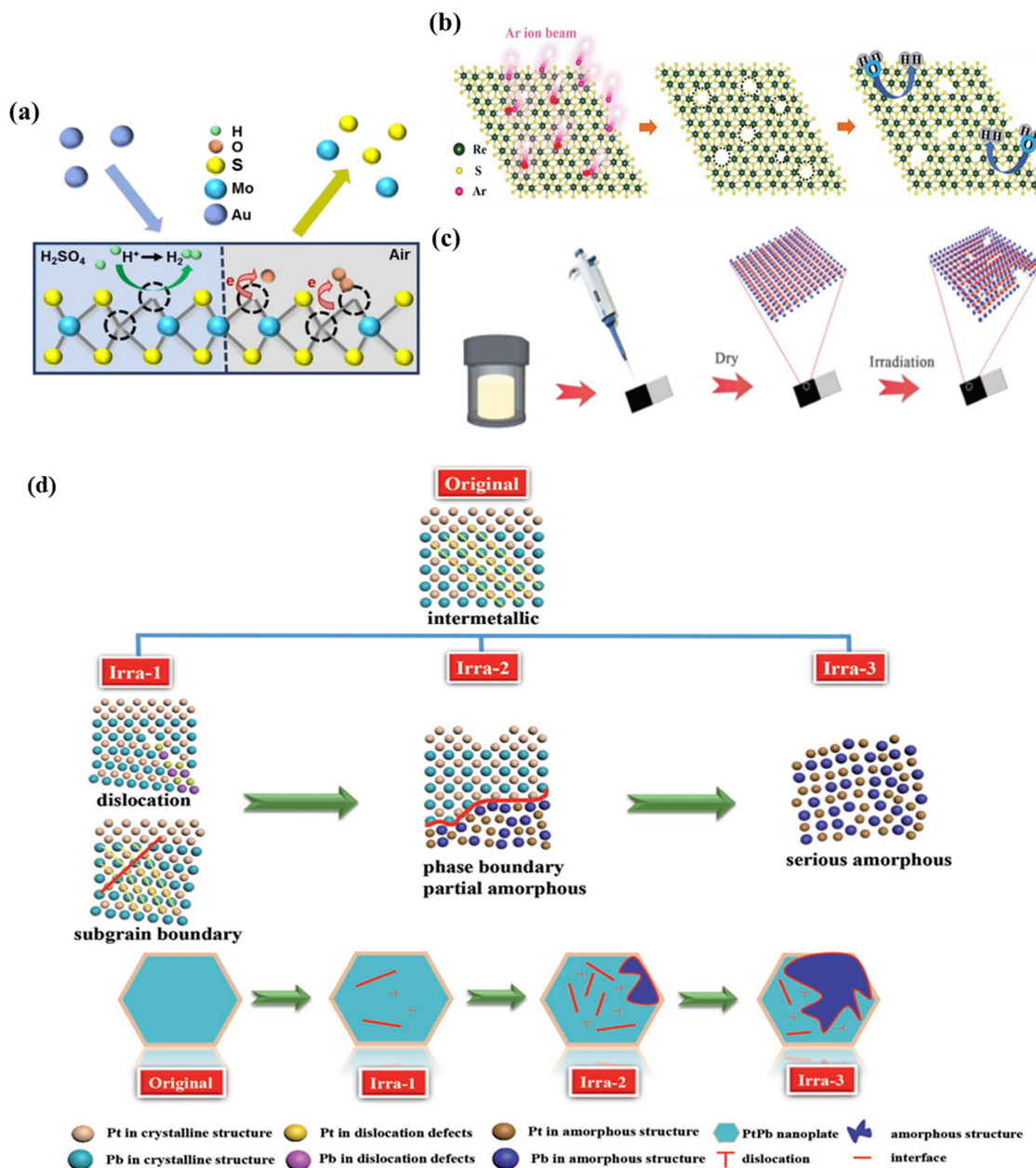


Fig. 4 (a) Scheme illustration of the defects in MoS<sub>2</sub> generated by Au ion-irradiation. Reproduced with permission from ref. 33. Copyright 2018, American Chemical Society. (b) Schematic diagram of Ar ion beam bombardment to 2D ReS<sub>2</sub> nanosheets and the formation of atomic Re vacancy defects for enhancing catalysis toward the solar-driven PEC-HER. Reproduced with permission from ref. 34. Copyright 2018, Wiley-VCH. (c) Scheme illustration of the preparation procedure of Fe-Bi<sub>2</sub>Te<sub>3</sub>/Ti. Reproduced with permission from ref. 35. Copyright 2020, Royal Society of Chemistry. (d) Simulation models of PtPb nanoplate structure evolution by controlling the C<sup>+</sup> ion fluences. Reproduced with permission from ref. 18. Copyright 2017, Wiley-VCH.



allowing  $H^+$  to be absorbed on the active sites neither strongly nor weakly. In addition, the distinctive stacked 3D structure has the obvious benefit of exposing more active sites and larger superficial areas to contact with electrolytes. These two reasons jointly enhanced the solar-driven hydrogen evolution reaction (PEC-HER) performance of  $ReS_2/Si$  photocathodes.

In a work by Wang,<sup>35</sup> high-energy Fe ions (a dose of  $4 \times 10^{14}$  ions per  $cm^2$ ) were irradiated on the surface of  $Bi_2Te_3$  nanosheets, which resulted in the generation of a variety of defects (Fig. 4c). The existence of defects resulted in an increase in the active sites and improvement in the conductivity of  $Bi_2Te_3$ , ensuring the rapid transfer and transport of charge during the catalytic process. Guo *et al.*<sup>18</sup> reported a new strategy by controlling the defects of PtPb nanoplates using  $C^+$  ion-irradiation. By controlling the ion fluence, PtPb nanoplates could be well transformed from single crystal to polycrystal with various degrees of dislocations, subgrain boundaries, and partial amorphization. After ion-irradiation, PtPb nanoplates can largely maximize the Pt catalytic efficiency and utilization enhancing electrocatalytic performance (Fig. 4d). The same group<sup>36</sup> synthesized PtPb NPs under nonaqueous conditions and followed this by 1 MeV  $Kr^{3+}$  ion-irradiation. Upon ion-irradiation, novel structure PtPd nanoplates are formed, in which the crystalline phase and amorphous phase co-existed; particularly, the crystalline phase is surrounded by the amorphous phase annularly to form a crystalline/amorphous interface. As a result,  $Kr^{3+}$  ion-irradiation makes the Pt electronic structure regulated due to the interface fusion of the crystalline/amorphous interface. Meanwhile, the crystalline/amorphous interface provides plenty of active sites, and the interface can

activate the C–H and O–H bonds to enhance the catalytic performance.

Chen *et al.*<sup>37</sup> prepared a single-layer  $MoS_2$  by the CVD method and transferred it onto various substrates such as Au, single-layer graphene (SLG), hexagonal boron nitride (BN) and  $CeO_2$ , and created defects by  $Ar^+$  ion-irradiation (Fig. 5a). They explored the substrate effect on the electronic structure of  $MoS_2$  by a series of measurement. It was found that these substrates can tune the electronic energy levels in  $MoS_2$  due to the charge transfer that occurred on the interface. Furthermore, when the reduced  $CeO_2$  as a substrate upon annealing significantly affects the energy band alignment of  $MoS_2$  and  $CeO_2/Au$  substrates, the reduction introduces electronic conductivity into the  $CeO_2$  substrate. The introduction of defects by ion-irradiation enhances the HER activity of single layer  $MoS_2$ . The tunability of defect types and concentrations in  $MoS_2$  by ion-irradiation at different energies provides a possibility to engineer the properties of  $MoS_2$  and other two-dimensional (2D) materials.

The introduced vacancies by ion-irradiation could also regulate the electron density of the material, which can help in better understanding of the relationship between the structure and performance. For example, He *et al.*<sup>38</sup> fabricated a series of Co-based catalysts (oxygen defect-containing  $Co_3O_4$  ( $Co_3O_4-O_v$ ) and multi phased  $CoO/Co_3O_4$ ) (a schematic illustration of  $Co_3O_4$ ,  $Co_3O_4-O_v$ , and  $CoO/Co_3O_4$  samples is shown in Fig. 5b) through  $Ar^+$  ion-radiation on pristine  $Co_3O_4$ .  $Co_3O_4$  was subjected to argon ion-irradiation at 25 keV ( $Co_3O_4-O_v$ ) and 50 keV ( $CoO/Co_3O_4$ ) with a dosage of  $5 \times 10^{15}$  ions per  $cm^2$ . The theoretical calculation and ultraviolet photoelectron

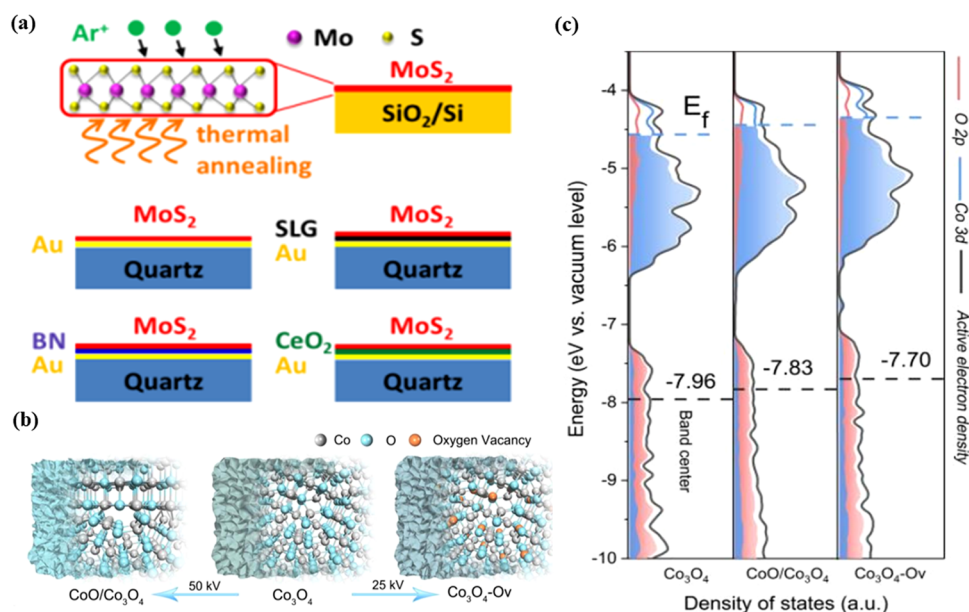


**Fig. 5** (a)  $MoS_2/Au$ ,  $MoS_2/SLG/Au$ ,  $MoS_2/BN/Au$  and  $MoS_2/CeO_2/Au$  studied in the work. All the structures are supported by quartz substrates. The  $MoS_2$  single layers in all these structures are transferred from CVD grown  $MoS_2$  on the  $Si/SiO_2$  substrate. Vacuum thermal annealing or  $Ar^+$  irradiation is used to treat the samples to either enhance the contact with substrates or produce defects. Reproduced with permission from ref. 37. Copyright 2018, American Chemical Society. (b) Schematic illustration and (c) PDOS plots of  $Co_3O_4$ ,  $Co_3O_4-O_v$  and  $CoO/Co_3O_4$  samples. Reproduced with permission from ref. 38. Copyright 2020, Wiley-VCH.

spectrometer characterization together indicated that the band center of  $\text{Co}_3\text{O}_4$  upshifted, thereby improving the adsorption capacity of the oxo group and greatly reducing the reaction barrier. This consequence further proved that the surface active electron density of  $\text{Co}_3\text{O}_4$  can be effectively regulated by ion-irradiation (Fig. 5c).

### 3.2 Element doping

Ion-irradiation is known to be a novel and effective element doping means commonly used in metal oxides, heterojunctions, and graphene which usually involves two steps in fabricating materials including ion-implantation and annealing.

Liu *et al.*<sup>39</sup> successfully prepared noble metal nanoparticles and metallic (Fe, V) and non-metallic (N) doped  $\text{TiO}_2$  by ion implantation (the energy of 20–50 kV, with different ion sources Au, Fe, V, and N, at a fluence from  $0.1$  to  $20 \times 10^{16}$  ions per  $\text{cm}^2$ ) and subsequent annealing. After doping of Au, the photocatalytic activity of the Au- $\text{TiO}_2$  sample was enhanced because of Au NPs in response to the incident light, which enhance the absorption, the local electric field and the excitation of active electrons and holes. For Fe doped  $\text{TiO}_2$ , the hybridization of the  $\text{Ti}^{4+}$  3d orbital with  $\text{Fe}^{3+}$  3d orbitals occurs, which introduces a new energy in the band structure of  $\text{TiO}_2$  and therefore improves the response to visible light. For V-doped  $\text{TiO}_2$ , the increased V doping promotes the transformation from the anatase to the rutile phase. Besides, the implanted V-doped  $\text{TiO}_2$  exhibits a red-shifted absorption edge which leads to a significant enhancement of light absorption in the visible light region. For N-doped  $\text{TiO}_2$ , the experimental results and theoretical calculations demonstrate that the substitution of nitrogen for oxygen atoms in  $\text{TiO}_2$  induces an isolated localized state of N 2p above the O 2p dominant valence band. The  $\text{TiO}_2$ -based composite films are fabricated by ion implantation (Au, Fe, V, and N element doping in  $\text{TiO}_2$ ) and show high crystallinity and stability, and the enhanced photocatalytic activities towards visible light were observed. The photocatalytic activity is greatly improved, suggesting that the N doping level had an obvious influence on the photocatalytic activity of the  $\text{TiO}_2$  samples. This report suggested that different irradiation sources can affect the properties of materials in different ways.

The same group reported the formation and thermal evolution of  $\text{Al}_2\text{O}_3$  substrates by high fluence  $\text{Zn}^+$  ion implantation at different fluences ( $1 \times 10^{17}$  and  $2 \times 10^{17}$  ions per  $\text{cm}^2$ ).<sup>40</sup> They first implanted  $\text{Zn}^+$  onto the surface of the  $\alpha\text{-Al}_2\text{O}_3$  sample. Then, the implanted sample was annealed in an oxygen atmosphere at  $700^\circ\text{C}$  for 2 h. During the annealing process,  $\text{Zn}^+$  diffused to the surface and was oxidized into ZnO which indicates that the implanted  $\text{Zn}^+$  was successfully doped onto the surface of  $\alpha\text{-Al}_2\text{O}_3$ . Particularly, the higher quality ZnO film formed on the surface of  $\text{Al}_2\text{O}_3$  single crystal at a fluence of  $1 \times 10^{17}$  ions per  $\text{cm}^2$  by  $\text{Zn}^+$  implantation combined with sequential annealing at  $700^\circ\text{C}$  for 2 h in the oxygen atmosphere. Zhao *et al.*<sup>41</sup> reported ion-implantation for the direct synthesis of nitrogen-doped graphene (NG). Firstly, they fabricated a newly multilayered substrate (Ni/Cu/a-SiC/ $\text{SiO}_2$ /Si). Then, nitrogen ion-implantation (with an ion energy of 40 keV

at  $1 \times 10^{16}$  ions per  $\text{cm}^2$ ) was performed on the multilayered substrate of Ni/Cu/a-SiC/ $\text{SiO}_2$ /Si. Finally, the N ion-implanted substrate was annealed by using rapid thermal processing to obtain NG (Fig. 6a). Through nitrogen ion-implantation, they proved the successful synthesis of nitrogen-doped graphene using transmission electron microscopy (TEM), Raman spectroscopy, X-ray photoelectron spectroscopy (XPS), and the measurement of electronic properties, demonstrating that the N ion-implantation could effectively dope graphene. The ion-implantation technique not only can be expected to produce NG but also opens up a new way to extend more element doping of other materials.

### 3.3 Morphology control

During the ion-irradiation process, the sputtering effect is a common phenomenon which is assumed to result from cascades of atomic collisions by energetic ions or recoil atoms.<sup>42–44</sup> When energetic ions shoot into a target substrate, the incident ions impact the atoms of the target substrate, and these atoms may obtain enough energy to be ejected out of the substrate; thus, the surface sputtering effect can modulate the surface morphology of materials.<sup>45,46</sup> Ishaq *et al.*<sup>47</sup> used different ion fluences to investigate the effects of  $\text{H}^+$ ,  $\text{N}^+$ , and  $\text{Ar}^+$  ion-irradiation on the structural and morphological changes of multi-walled carbon nanotubes (MWCNTs) and amorphous carbon nanowires (a-CNWs). They found that, at room temperature, defects were introduced in MWCNTs under 70 keV  $\text{N}^+$  ion beam irradiation at a low dose of  $1 \times 10^{16}$  ions per  $\text{cm}^2$ . Meanwhile, under high fluence  $\text{N}^+$  irradiation ( $7 \times 10^{16}$  ions per  $\text{cm}^2$ ), MWCNTs could transform into a-CNWs (Fig. 6b and c). At 1000 K, the 70 keV  $\text{N}^+$  ion beam irradiation of defective MWCNTs is transformed from curling of carbon clusters to an onion-like structure at a dose of  $4 \times 10^{16}$  ions per  $\text{cm}^2$  and the formation of an onion-like structure is at  $7 \times 10^{16}$  ions per  $\text{cm}^2$  (Fig. 6d and e). The type of the ion source also affects the morphology of MWCNTs. At 1000 K, under  $\text{Ar}^+$  ion-irradiation ( $7 \times 10^{16}$  ions per  $\text{cm}^2$ ), the defective MWCNTs transform into a high-density onion-like structure (Fig. 6f and g). When the ion source is changed to an  $\text{H}^+$  ion source, the ion fluence is changed from  $1 \times 10^{16}$  to  $1 \times 10^{17}$  ions per  $\text{cm}^2$ , and the a-CNWs gradually increase the fraction of the material with a diamond structure (Fig. 6h). These results showed that the change of ion-irradiation parameters (including ion irradiation energy, dose and annealing temperature) has been seen for crystalline–amorphous and amorphous–crystallization changes in the carbon nanostructure. Additionally, Jiang *et al.*<sup>48</sup> utilized Ag ion-irradiation and annealing to fabricate hollow microsphere arrays. After Ag ion-irradiation, the surface of the polystyrene sphere (PS) microsphere was carbonized. Through annealing or chloroform immersion, the non-carbonated zone can be easily removed, and then the hollow microsphere. They have investigated the influence of the ion-irradiation energy and fluence on the morphology and structure of the PS microsphere array. They found that, at 15 kV, with the increase of Ag ion fluences ( $1 \times 10^{16}$ ,  $3 \times 10^{16}$ , and  $5 \times 10^{16}$  ions per  $\text{cm}^2$ ), the morphology and size of PS microspheres were almost



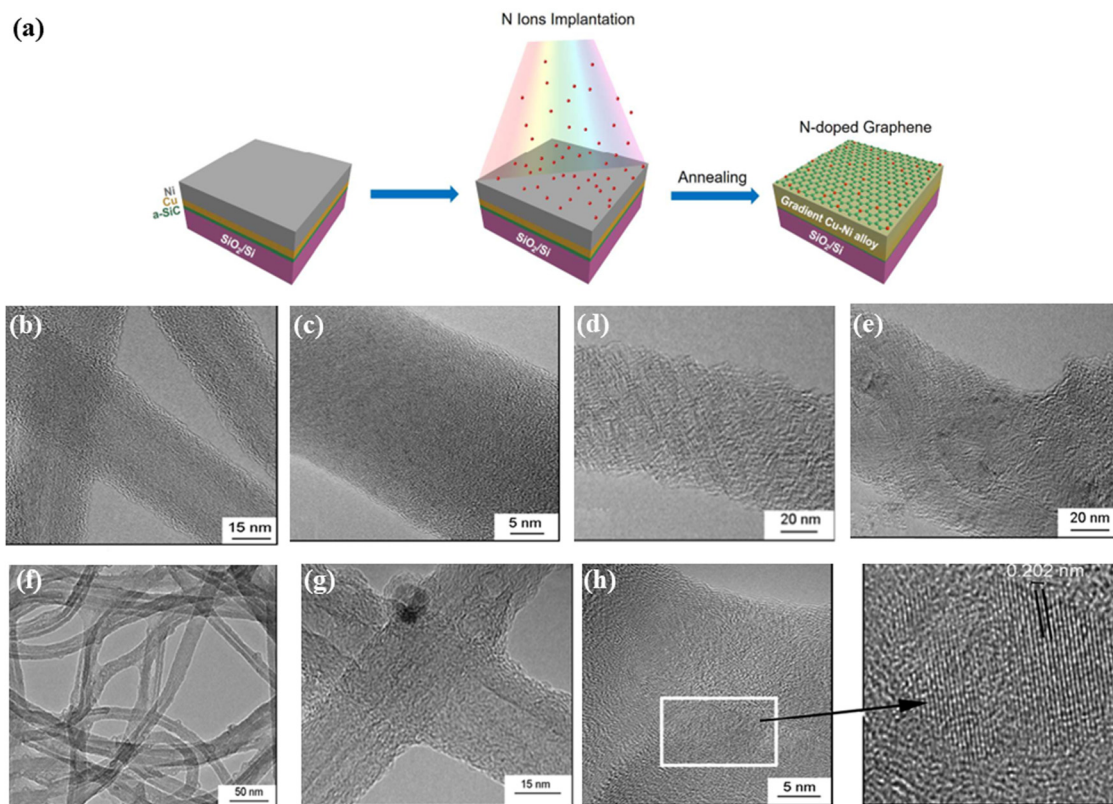


**Fig. 6** (a) Schematic illustration of the co-segregation method combined with ion implantation for one-step synthesis of NG. Transitions of MWCNTs to amorphous carbon nanowires under 70 keV  $N^+$  ion beam irradiation at room temperature. Reproduced with permission from ref. 41. Copyright 2018, Elsevier. (b) A MWCNT irradiated at a dose of  $1 \times 10^{16}$  ions per  $cm^2$  and (c) an MWCNT irradiated at a dose of  $7 \times 10^{16}$  ions per  $cm^2$ . HR-TEM images of  $N^+$  ion irradiated defective MWCNTs (1000 K) at a dose of (d)  $4 \times 10^{16}$  ions per  $cm^2$  and (e)  $7 \times 10^{16}$  ions per  $cm^2$ . (f and g) HR-TEM images of  $Ar^+$  ion-irradiated defective MWCNTs at a dose of  $7 \times 10^{16}$  ions per  $cm^2$  at 1000 K. (h) A diamond particle nucleates on a-CNWs under  $H^+$  ion beam irradiation at 1000 K, and the crystalline diamond structure with a spacing of about 0.202 nm was formed into a-CNWs at a dose of  $1 \times 10^{17}$  ions per  $cm^2$ . Reproduced with permission from ref. 47. Copyright 2013, Elsevier.

unchanged. However, when the irradiated fluence reached  $7 \times 10^{16}$  ions per  $cm^2$ , the coalescence phenomenon of neighboring PS microspheres occurred. Then, they chose this irradiated fluence ( $7 \times 10^{16}$  ions per  $cm^2$ ) in subsequent experiments and attempted to change the energy to investigate the influence of ion-irradiated energy on the morphology of PS spheres. With the increasing irradiated energies (10, 15, 20, 40, and 60 kV), not only the morphology and size of the PS sphere but also the structure of the close-packed 2D hexagonal array has undergone dramatic changes. Under the same irradiated parameters, after thermal annealing at 350 °C for 30 min, the structure of samples has transformed into spherical shell arrays. The main reason for the formation of spherical shell arrays is due to the carbonization effect produced by calcination and the sputtering effect generated through ion-irradiation (regulating different irradiated parameters). The Ag ion beam modification diversifies the structure and morphology of the PS sphere array and greatly improves its thermostability.

In a work by Giulian,<sup>49</sup> Pt nanoparticles (NPs) in amorphous  $SiO_2$  were irradiated with 185 MeV Au ions in the fluences varying from  $2 \times 10^{14}$  to  $2 \times 10^{13}$  ions per  $cm^2$ . From the TEM images, with the increasing irradiation fluence, it was observed

that Pt NPs transform from spherical to rod-like shapes. The deformation process is studied as a function of the NP size, varying irradiation fluence and energy give a broad prospective of the shape transformation. Based on the above reports, the morphology of materials can be further regulated by changing ion-irradiation parameters (including ion irradiation energy, dose and annealing temperature), thus providing a broader application prospect for ion irradiation technology.

### 3.4 Material synthesis

Besides the modifications, as in an earlier study, ion-implantation has been used to synthesize nanoparticles. Taguchi *et al.*<sup>50</sup> synthesized C-SiC coaxial nanotubes by ion-irradiation as shown in Fig. 7. Firstly, C-SiC coaxial nanotubes were synthesized by heat treatment of MWCNTs and Si powder without direct contact with each other at 1200 °C. Then, coaxial nanotubes were deposited on a molybdenum grid TEM sample holder with a holey-carbon layer. The deposited coaxial nanotubes were irradiated with both 200 keV Si ions with a dose of  $1.1 \times 10^{21}$  ions per  $cm^2$  and 400 keV electrons simultaneously using the TEM instrument. The hybrid consisted of one-dimensionally stacked graphene nanodisks with diameters less than 50 nm and cylindrical



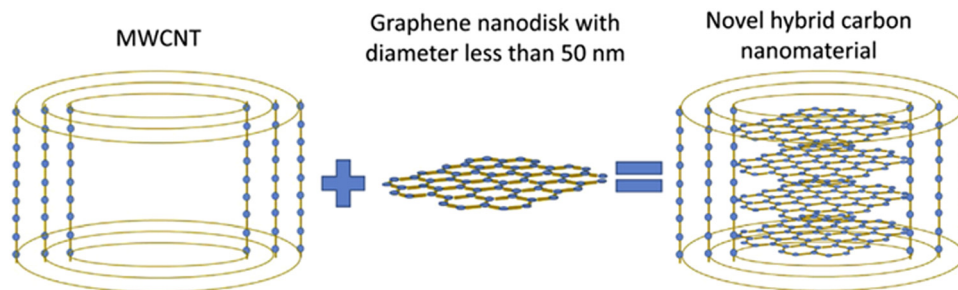


Fig. 7 Schematics of the microstructure of novel hybrid carbon nanomaterials inside an amorphous silicon carbide nanotube synthesized by ion-irradiation. Reproduced with permission from ref. 50. Copyright 2019, Elsevier.

multiwalled carbon nanotubes inside an amorphous SiC tubular layer. A sudden emergence of new continuous graphitic layers in the microstructure was observed by *in situ* transmission electron microscopy following ion-irradiation, where these layers were perpendicular to the length direction of the nanotube. Also, Tsai *et al.*<sup>51</sup> synthesized multilayer silicene by B ion-implantation. The boron ions were implanted into the target materials and reacted with carbon at the surface of SiC to form B<sub>4</sub>C during annealing. Simultaneously, silicon atoms were apparently squeezed out of the surface and then rearranged into the multilayer silicon.

Zheng *et al.*<sup>52</sup> fabricated tungsten oxide (WO<sub>3-x</sub>) nanowires by Ar<sup>+</sup> ion beam irradiation and subsequently annealed them in a vacuum. In the first step, the WO<sub>3</sub> thin film (~70 nm thickness) was deposited on silica slides at 400 °C using an ultra high vacuum magnetron sputtering system. Then, the size of as-deposited WO<sub>3</sub> thin films was controlled by tuning irradiation parameters (ion energies of 80, 130, 190 keV and with different

fluences of  $1 \times 10^{17}$  and  $1.5 \times 10^{17}$  ions per cm<sup>2</sup>) and annealing conditions. The results demonstrated that the nanowire length increases with the increasing irradiation fluence and with the decreasing ion energy. The electrical conductivity of the nanowire film was higher than that of the pristine film, which is attributed to the high-density irradiation-induced vacancies on the oxygen sublattice greatly improving the conductivity. Furthermore, Wang *et al.*<sup>53</sup> reported that the number of graphene layers can be controlled by modulating the C ion-implantation (with ion dosages of  $4 \times 10^{15}$  and  $8 \times 10^{15}$  ions per cm<sup>2</sup>). Fig. 8 shows the Raman spectra and scanning tunneling microscopy (STM) topographical images of the monolayer and bilayer graphene synthesized by ion-implantation. The Raman mapping result indicates that ion-implantation can fabricate highquality monolayer and bilayer graphene.

In general, this review involves the influence of ion-irradiation on materials, including defect introduction, element doping, morphology control and material synthesis. These effects are

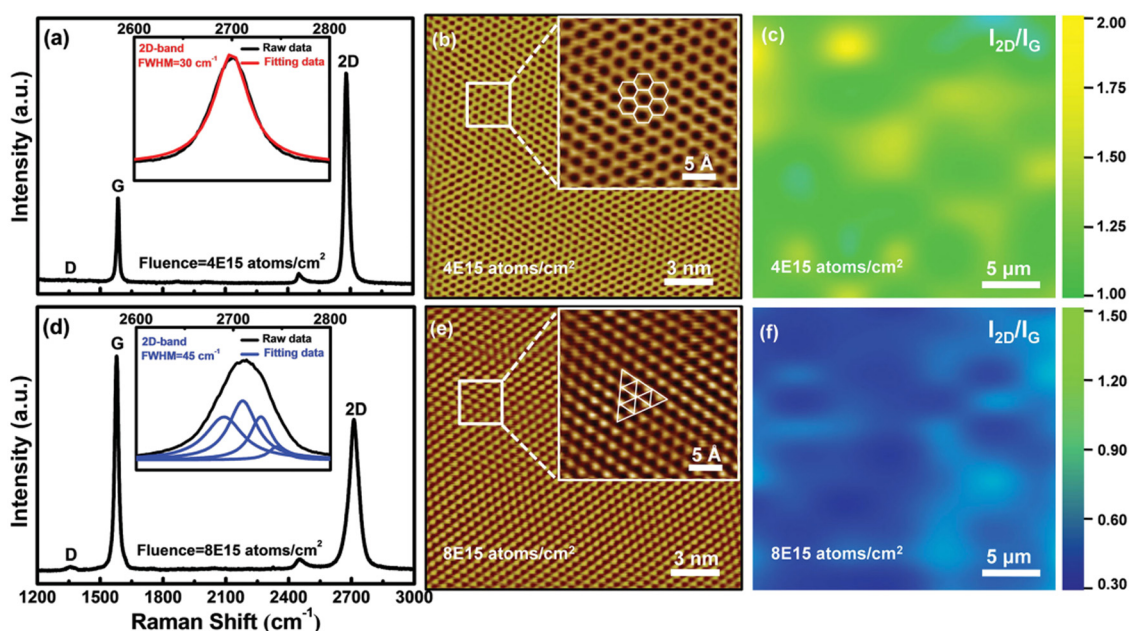


Fig. 8 Raman spectra of the (a) monolayer and (d) bilayer graphene. STM topographical images of the (b) monolayer and (e) bilayer graphene. The  $I_{2D}/I_G$  peak ratio derived from Raman mapping conducted on the (c) monolayer and (f) bilayer graphene. Reproduced with permission from ref. 53. Copyright 2015, Wiley-VCH.



the means of modification/designing of catalyst and electrode materials, which is a positive effect. However, ion-irradiation is not suitable for the modification/designing of all materials. For example, the ultrathin nanosheet/nanoplate is easy to be penetrated.<sup>54–57</sup> Consequently, materials may be unstable under ion-irradiation. In other words, ion-irradiation technology is not suitable for all catalyst/electrode materials.

## 4. Application of ion-irradiation technology in catalytic/electrode materials for energy conversion and storage

### 4.1 Application of ion-irradiation technology for electrocatalysts

**4.1.1 HER electrocatalysts.** Water electrolysis to produce hydrogen is one of the most promising technologies for energy storage and conversion, which involves two half reactions. One is the HER and the other is the oxygen evolution reaction (OER). Both reactions need electrocatalysts to speed up the reaction kinetics.<sup>3,58</sup> Therefore, developing efficient catalysts is essential for accelerating the reaction kinetics and greatly reducing electrode overpotentials.

Recently, the 2D MoS<sub>2</sub> nanosheet as a promising alternative to Pt for the HER has drawn persistent interest. Several studies have revealed that the electrocatalytic HER performance of nanoscale MoS<sub>2</sub> mainly arises from its edge states, while the basal plane is inert. If the inert basal plane sites can be optimized, meanwhile the edge activity can be maintained, the catalytic activity of MoS<sub>2</sub> will be enormously improved.<sup>59–62</sup> It was found that the inert basal plane of MoS<sub>2</sub> could be activated by introducing sulphur defects into the surface. However, ion-irradiation is a novel strategy which can be used to introduce sulphur defects into MoS<sub>2</sub> and thus improve catalytic activity.<sup>33,63,64</sup> Sun *et al.*<sup>32</sup> use C ions to irradiate MoS<sub>2</sub> by regulating different C ion fluences ( $5 \times 10^{12}$ ,  $2 \times 10^{13}$ , and  $5 \times 10^{13}$  ions per cm<sup>2</sup>). They found that ion-irradiation generates sulphur vacancies which were active sites on MoS<sub>2</sub> basal planes, which could improve HER performance. It was shown that different ion fluences could tune the amount of sulphur vacancies and the amorphous region on the basal plane of MoS<sub>2</sub>. When MoS<sub>2</sub> was ion-irradiated by a fluence of  $2 \times 10^{13}$  ions per cm<sup>2</sup>, the best HER performance could be achieved with an HER onset potential of 77 mV at 10 mA cm<sup>-2</sup> and a small Tafel slope of 66 mV dec<sup>-1</sup>.

Different from the above low energy ion-irradiation, Madauss *et al.*<sup>65</sup> used Xe ions (irradiated energy 91 MeV) to appropriately modify the MoS<sub>2</sub> surface and thus achieved high catalytic activity for the hydrogen evolution reaction. At a potential of -0.6 V vs. RHE, the irradiated MoS<sub>2</sub> exhibits a higher current density (35.3 mA cm<sup>-2</sup>) than non-irradiated MoS<sub>2</sub> (-13.3 mA cm<sup>-2</sup>) (Fig. 9a). The irradiated sample has a smaller Tafel slope (104 mV dec<sup>-1</sup>) than the non-irradiated MoS<sub>2</sub> (106 mV dec<sup>-1</sup>) (Fig. 9b). The results are due to a large

increase in the number of low-coordinated Mo atoms after ion-irradiation, which can form bonds with adsorbing species and thus improving catalytic activity.

Additionally, Xia *et al.*<sup>66</sup> fabricated MoSe<sub>2</sub> nanosheets grown on carbon cloth (CC) by the CVD method, which were subsequently irradiated by Ar<sup>2+</sup> ions for providing additional active sites on the inert basal planes of nanosheets. MoSe<sub>2</sub> was irradiated by using different doses of Ar<sup>2+</sup> ions. Compared with pristine MoSe<sub>2</sub>/CC, the higher dose ion-irradiated MoSe<sub>2</sub> nanosheets ( $5 \times 10^{15}$  ions per cm<sup>2</sup>) showed the lowest overpotential of -171 mV (Fig. 9c) at a current density of -100 mA cm<sup>-2</sup> and the smallest Tafel slope of 35 mV dec<sup>-1</sup> (Fig. 9d). The first-principle calculation was employed to investigate the electron states of four different vacancy (single Mo vacancy (V<sub>Mo</sub>), single Se vacancy (V<sub>Se</sub>), double Se vacancy (V<sub>Se2</sub>) and the absence of one MoSe<sub>2</sub> molecular (V<sub>MoSe2</sub>)) cases observed in the MoSe<sub>2</sub> basal plane. The calculated results showed that the pristine MoSe<sub>2</sub> exhibits a higher bandgap (1.23 eV) than V<sub>Mo</sub> and V<sub>Se2</sub> band gaps (0.04 eV and 0.81 eV, respectively), which suggests the improved electronic conductivity, thereby promoting the electronic transferring during the HER process. Upon ion-irradiation, the existence of vacancies in the MoSe<sub>2</sub> basal plane can not only lead to better conductivity but also produce more electrocatalytic active sites in MoSe<sub>2</sub>, ensuring the better HER catalytic ability of MoSe<sub>2</sub>.

Wang *et al.*<sup>35</sup> prepared Bi<sub>2</sub>Te<sub>3</sub> nanosheets on a Ti plate and followed this by Fe<sup>+</sup> irradiation, which can regulate the balance of hydrophilicity and hydrophobicity for the material. The contact angle result indicates that the Fe ion irradiated Bi<sub>2</sub>Te<sub>3</sub>/Ti exhibits hydrophilic properties, whereas the non-irradiated Bi<sub>2</sub>Te<sub>3</sub>/Ti exhibits hydrophobic properties (Fig. 10a). The result suggests that the Fe<sup>+</sup> irradiation can manipulate the hydrophilicity/hydrophobicity balance of the Bi<sub>2</sub>Te<sub>3</sub> surface, facilitating the release of hydrogen bubbles from the catalyst surface and exposing the active sites in time. While the damage caused by large bubbles on the electrode material was avoided, the stability of the material was improved (Fig. 10b and c). Silicon (Si) is rarely used as the HER catalyst due to its inert electrochemical activity. However, Wu *et al.*<sup>67</sup> fabricated nanoporous Si as the HER catalyst by using Ar<sup>+</sup> ion-irradiation followed by an annealing method. When the Ar<sup>+</sup> ion-irradiation fluence reached  $1 \times 10^{17}$  ions per cm<sup>2</sup>, the nanoporous Si had more positive HER onset potentials and a smaller value of Tafel slope than the pristine Si (Fig. 10d and e). The higher catalytic HER activity of the irradiated Si could be attributed to the formation of nanopores during Ar<sup>+</sup> irradiation, which promotes H<sub>2</sub> bubbles escaping from the catalyst surface and thus exposes more active sites. They also investigated the influence of fluences ( $5 \times 10^{16}$ ,  $1 \times 10^{17}$ , and  $2 \times 10^{17}$  ions per cm<sup>2</sup>, respectively; the irradiated samples were named NPS-1, 2, and 3) on the HER performance of the samples. It was found that NPS-2 (at  $1 \times 10^{17}$  Ar<sup>+</sup> ions per cm<sup>2</sup> irradiated) showed the highest HER performance (Fig. 10f). The largely enhanced HER activity comes from the unique morphology and large specific surface area, which results from Ar<sup>+</sup> ion irradiation and post-irradiation annealing. This method provides a novel research strategy to



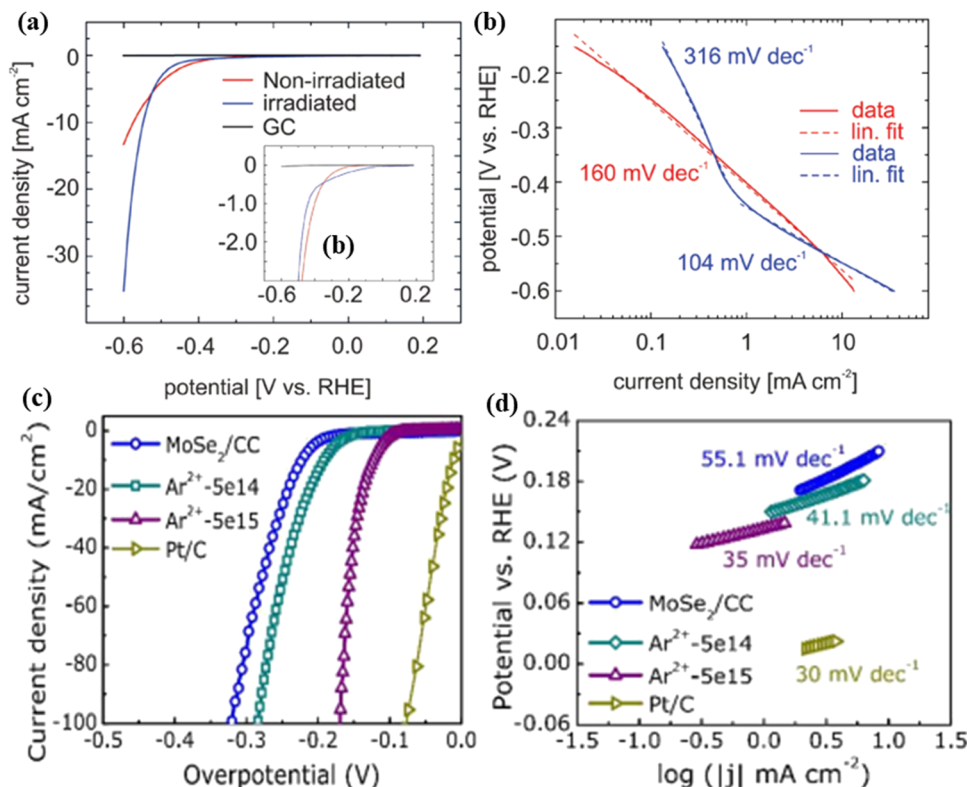


**Fig. 9** (a) Current density as a function of the potential (V vs. RHE) showing strong activity enhancement of the irradiated MoS<sub>2</sub> compared to that of the non-irradiated MoS<sub>2</sub>. The inset of (a) shows the less negative onset potential for the irradiated MoS<sub>2</sub>. (b) *V*–*I* curves with Tafel-slopes of the irradiated (blue) and non-irradiated (red) MoS<sub>2</sub>. Reproduced with permission from ref. 65. Copyright 2018, Royal Society of Chemistry. The HER performance of the as-fabricated catalysts. (c) LSV curves and (d) Tafel plots of MoS<sub>2</sub>/CC, Ar<sup>2+</sup>-5e14, Ar<sup>2+</sup>-5e15 and Pt/C electrodes, respectively. Reproduced with permission from ref. 66. Copyright 2018, American Chemical Society.

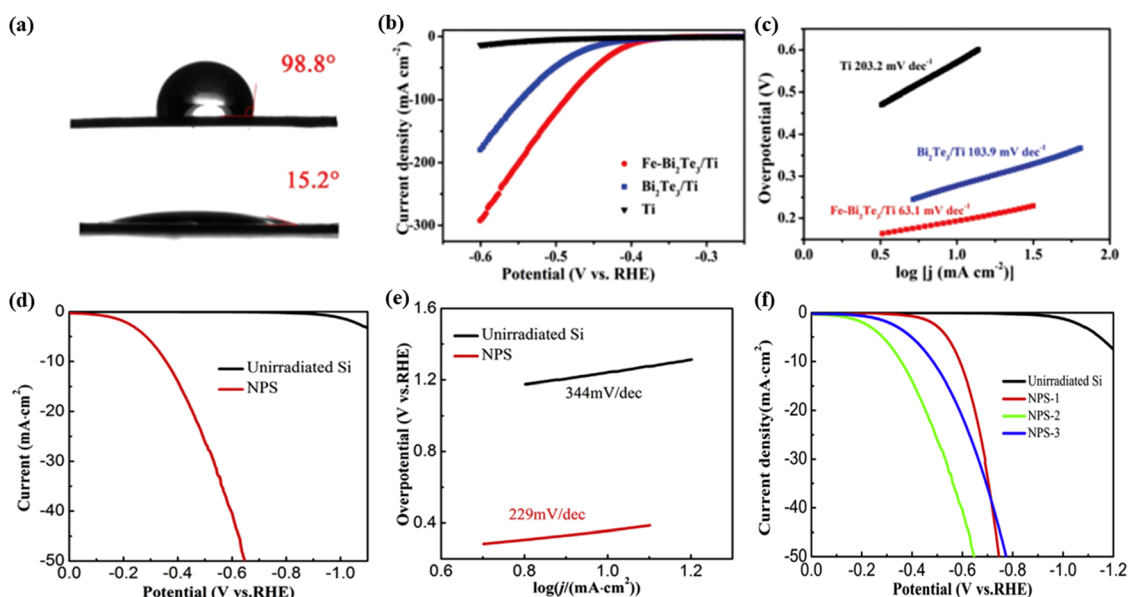


**Fig. 10** (a) Contact angles of Bi<sub>2</sub>Te<sub>3</sub>/Ti (top) and Fe-Bi<sub>2</sub>Te<sub>3</sub>/Ti (bottom), respectively. (b) LSV curves of the Ti plate, Bi<sub>2</sub>Te<sub>3</sub>/Ti and Fe-Bi<sub>2</sub>Te<sub>3</sub>/Ti at a scan rate of 5 mV s<sup>-1</sup>. (c) The corresponding Tafel plots. Reproduced with permission from ref. 35. Copyright 2020, Royal Society of Chemistry. (d) Polarization curves of the nanoporous and non-irradiated Si. (e) Tafel slopes derived from the polarization curves. (f) The polarization curves for the fluences of NPS-1, NPS-2 and NPS-3. Reproduced with permission from ref. 67. Copyright 2018, Elsevier.

further improve the catalytic performance for energy conversion and catalysis.

**4.1.2 OER electrocatalysts.** The OER is an important half-reaction of water splitting for hydrogen generation.<sup>11</sup> To obtain affordable, efficient and environmentally friendly OER electrode materials, the compounds of VIII 3d-transition metals (such as Fe, Co, and Ni) including metal oxides,<sup>68–73</sup> sulfides<sup>74–78</sup> and phosphides<sup>79–83</sup> have been extensively investigated for their low cost and abundance as well as similar outermost electron configurations to Ru, Ir and Pt.

Xia *et al.*<sup>11</sup> used  $N^+$  ion-irradiation (a dose of  $5 \times 10^{15}$  ions per  $cm^2$ ) to introduce N dopants and oxygen vacancies in NiO nanosheets grown on carbon cloth (Fig. 11a). In the experiment,  $N^+$  ion-irradiated NiO nanosheets exhibited a high OER performance with a potential of 1.98 V at a current density of  $100 \text{ mA cm}^{-2}$  (Fig. 11b). The double layer capacitance ( $C_{dl}$ ) values of pristine and irradiated NiO/CC are estimated to be 4.15 and  $6.49 \text{ mF cm}^{-2}$ , respectively (Fig. 11c), which demonstrate that the irradiated sample possesses larger effective electrochemical surfaces. The charge transfer resistance ( $R_{ct}$ ) value of the irradiated electrode is evaluated as  $13.8 \Omega$ , which is

smaller than that of the pristine sample ( $16.9 \Omega$ ) (Fig. 11d), indicating the better electrical integration. The high turnover frequency (TOF) value ( $92.1 \text{ s}^{-1}$ ) (Fig. 11e) demonstrates the high intrinsic activity of irradiated samples. Density functional theory (DFT) calculations further reveal that the introduced oxygen vacancies and opportune N dopants could narrow the NiO band gap which vested a better conductivity for charge transfer during the OER process. This work provides a novel path to enhance the OER electrocatalytic performance by ion irradiation.

He *et al.*<sup>38</sup> reported a novel strategy using an  $Ar^+$  ion-irradiation technology to regulate the electronic properties of Co-based catalysts (including oxygen vacancy-containing  $Co_3O_4$  ( $Co_3O_4-O_v$ ) and multi phased  $CoO/Co_3O_4$ ) as OER catalysts.  $Co_3O_4$  was subjected to argon ion-irradiation at 25 keV ( $Co_3O_4-O_v$ ) and 50 keV ( $CoO/Co_3O_4$ ) with a dosage of  $5 \times 10^{15}$  ions per  $cm^2$ . Notably,  $CoO/Co_3O_4$  demonstrates a much higher current density and lower onset potential than  $Co_3O_4-O_v$  (Fig. 11f). The overpotential of  $CoO/Co_3O_4$  (270 mV) is lower than that of the pristine  $Co_3O_4$  (410 mV) at  $10 \text{ mA cm}^{-2}$ . The Tafel slope of  $CoO/Co_3O_4$  ( $55 \text{ mV dec}^{-1}$ ) is smaller than those of the

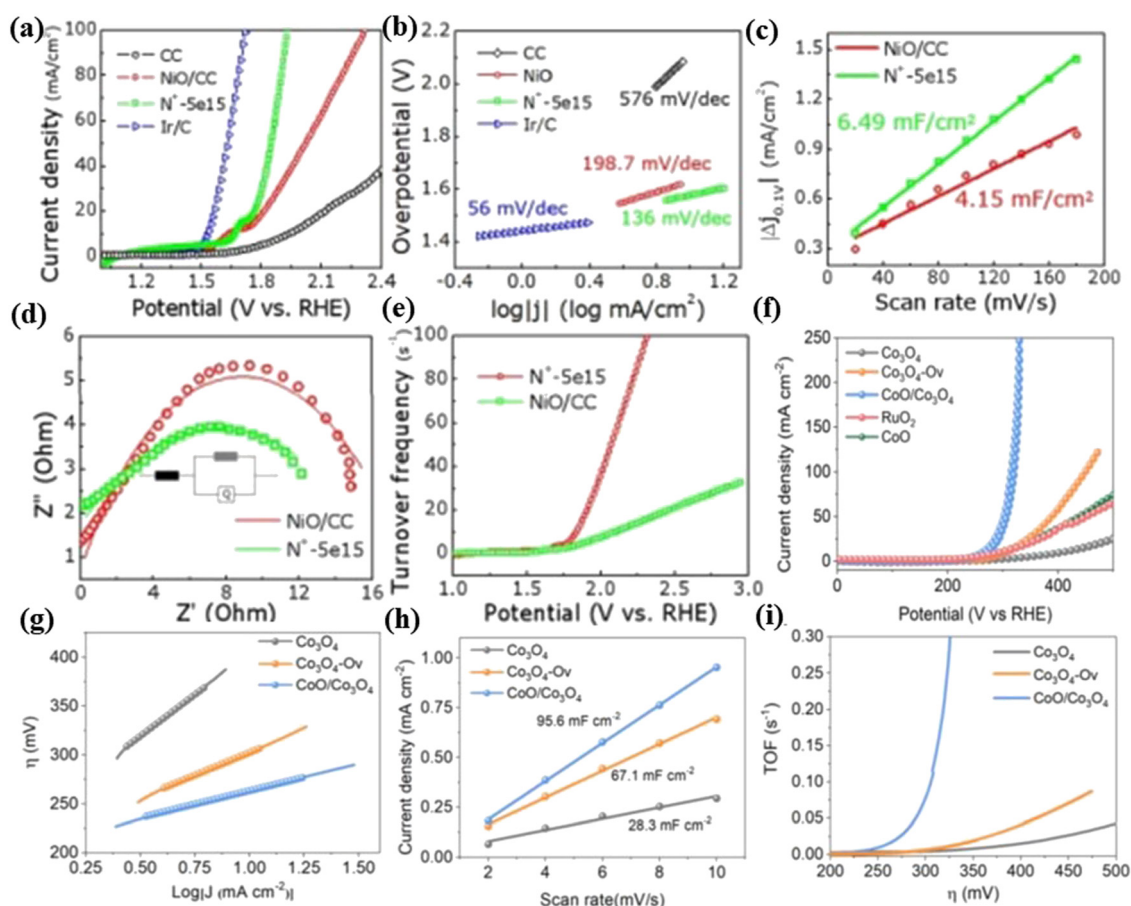
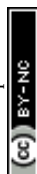


**Fig. 11** (a) LSV curves and (b) Tafel plots of carbon cloth, NiO/CC,  $N^+$ -5e15 and Ir/C electrodes. (c) Capacitive measurement results. (d) Nyquist plots of NiO/CC and  $N^+$ -5e15. (e) TOF of NiO/CC and  $N^+$ -5e15. Reproduced with permission from ref. 11. Copyright 2019, Royal Society of Chemistry. (f) Polarization curves of the OER for  $Co_3O_4$ ,  $Co_3O_4-O_v$ ,  $CoO/Co_3O_4$ ,  $CoO$  and  $RuO_2$ . (g) Tafel plots of the OER for  $Co_3O_4$ ,  $Co_3O_4-O_v$  and  $CoO/Co_3O_4$ . (h) Capacitive current measured at 1.16 V vs. RHE is plotted as a function of the scan rates of pristine  $Co_3O_4$ ,  $Co_3O_4-O_v$  and  $CoO/Co_3O_4$ . (i) Potential-dependent TOF curves of  $Co_3O_4$ ,  $CoO/Co_3O_4$  and  $Co_3O_4-O_v$ . Reproduced with permission from ref. 38. Copyright 2020, Wiley-VCH.



pristine  $\text{Co}_3\text{O}_4$  ( $169 \text{ mV dec}^{-1}$ ) and  $\text{Co}_3\text{O}_4\text{-O}_v$  ( $92 \text{ mV dec}^{-1}$ ), indicating the favorable OER kinetics for the multi-phased  $\text{CoO}/\text{Co}_3\text{O}_4$  sample (Fig. 11g). In addition, the  $C_{dl}$  of  $\text{CoO}/\text{Co}_3\text{O}_4$  is  $95.6 \text{ mF cm}^{-2}$ , which is higher than that of other samples (Fig. 11h). These results indicated that more active sites were generated and exposed to the irradiation of  $\text{Ar}^+$  ions. Besides, the calculated TOF value of  $\text{CoO}/\text{Co}_3\text{O}_4$  (Fig. 11i) is higher than those of  $\text{Co}_3\text{O}_4\text{-O}_v$  and  $\text{CoO}$ , suggesting its high intrinsic activity. Introducing  $\text{Ar}^+$  ion radiation can effectively modulate the surface active electron density of  $\text{Co}_3\text{O}_4$  by forming the  $\text{CoO}$  phase and oxygen vacancies, which can precisely and appropriately enhance the adsorption capacity of the oxo group and greatly reduce the reaction barrier. Using the ion-irradiation technology has great potential to be developed as a general strategy to regulate the surface electronic properties of metal oxides.

#### 4.2 Application of the ion-irradiation technique for photoelectrochemistry

Photoelectrochemical (PEC) water splitting has been considered to be a promising clean technique for generating hydrogen gas,<sup>84</sup> which has attracted much attention due to the increased demand for renewable energy.<sup>85,86</sup> Semiconductors are eminently attractive electrode materials for solar-driven PEC water splitting.<sup>34,87,88</sup> To better improve photo-redox reactions, these semiconductors must have a rational bandgap for being activated (*i.e.*, photoexcitation) by light absorption.<sup>89,90</sup> Many types of semiconductors such as  $\text{TiO}_2$ ,  $\alpha\text{-Fe}_2\text{O}_3$ , and  $\text{BiVO}_4$ , *etc.* have been intensively researched as potential photoelectrodes.<sup>91–95</sup>

Ion-irradiation has been playing an important role in modifying the properties of semiconductor materials. In general, the ion-irradiation effects on semiconductor materials are manifested as the changes in optical, physical, and electrical properties.

Singh *et al.*<sup>96</sup> reported the effect of 120 MeV  $\text{Ag}^{9+}$  ion irradiation on the PEC properties of  $\text{TiO}_2$  thin films deposited on a conducting glass substrate ( $\text{SnO}_2\text{:In}$ ). The PEC properties of the films were found to be greatly affected by the fluence of irradiation. The optimal observed fluence of radiation for the best PEC response was  $5 \times 10^{11}$  ion per  $\text{cm}^2$ , at which the photocurrent density reaches a maximum of  $0.76 \text{ mA cm}^{-2}$  (Fig. 12a). The irradiated samples also exhibited a maximum flat-band potential at the junction and a maximum charge carrier density. The satisfying PEC properties by ion-irradiation may be due to the formation of the desired structure of vertically aligned grains caused by ion-irradiation, which can efficiently separate the photogenerated electron-hole pairs before recombination. Dass *et al.*<sup>97</sup> modified the nanostructured 2%Cr- $\text{Fe}_2\text{O}_3$  and  $\text{CuO}$  thin films by using  $\text{Au}^{13+}$  SHI for PEC. The photocatalytic activity of the 2%Cr- $\text{Fe}_2\text{O}_3$  thin film irradiated increases at a fluence of  $10^{12}$  ions per  $\text{cm}^2$ , which might be due to more efficient absorption of photons resulting from the introduction of defects and the occurrence of structural rearrangements. Furthermore, it was observed that the decrease in photoresponse of irradiated 2%Cr- $\text{Fe}_2\text{O}_3$  and irradiated  $\text{CuO}$  thin films at higher fluences of  $10^{13}$  and  $10^{12}$  ions per  $\text{cm}^2$  (Fig. 12b and c) is attributed to the formation of a large number of discontinuities or dislocations in films. These defects act as recombination

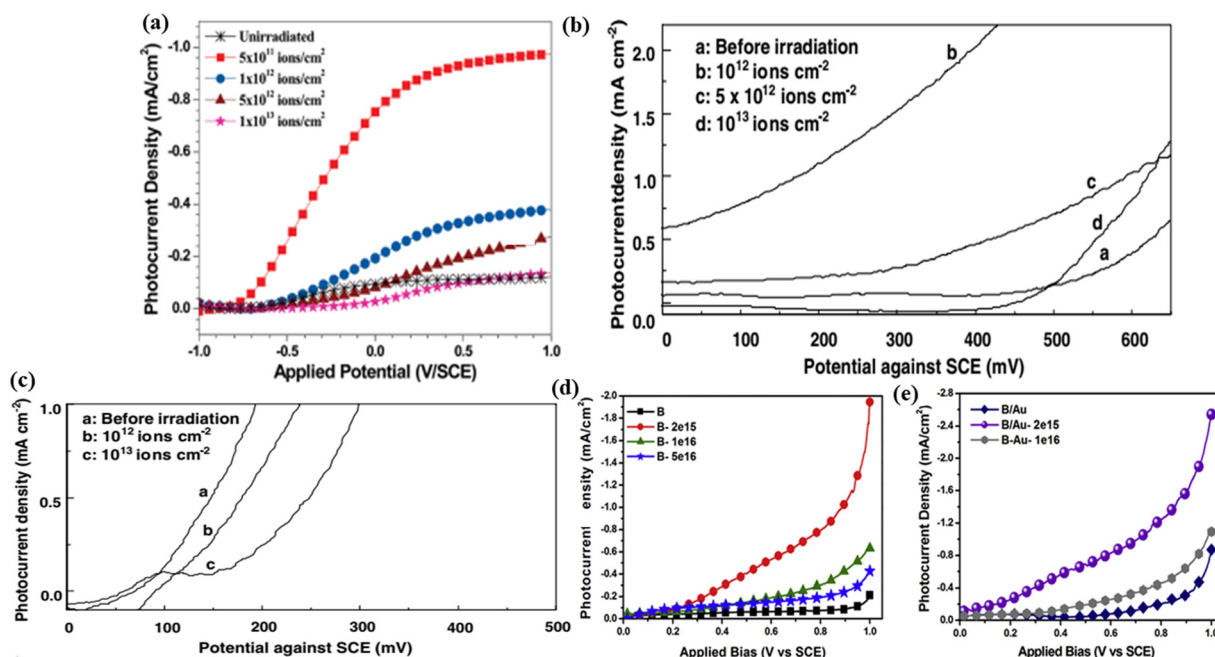


Fig. 12 (a) Photocurrent density curves of non-irradiated and 120 MeV  $\text{Ag}^{9+}$  ion irradiated  $\text{TiO}_2$  thin films at different fluences. Reproduced with permission from ref. 96. Copyright 2010, American Chemical Society. (b) Observed photocurrent density as a function of applied potential with 2% Cr doped  $\text{Fe}_2\text{O}_3$  thin film electrodes before and after SHI irradiation. (c) Observed photocurrent density as a function of applied potential with  $\text{CuO}$  thin film electrodes before and after SHI irradiation. Reproduced with permission from ref. 97. Copyright 2006, Elsevier. (d) and (e) Photocurrent density vs. applied bias for  $\text{BiVO}_4$  and  $\text{BiVO}_4/\text{Au}$  photoanodes. Reproduced with permission from ref. 88. Copyright 2019, Elsevier.

centers for photogenerated charge carriers, resulting in the decrease of the photocurrent.

Srivastav *et al.*<sup>88</sup> investigated the effect of  $N^+$  ion-irradiation on plasmonic Au nanoparticle loaded  $BiVO_4$  photoelectrodes for PEC water splitting. At an irradiated  $2 \times 10^{15}$  ions per  $cm^2$  fluence, the  $BiVO_4/Au$  photoanode shows a photocurrent density of  $2.54 \text{ mA cm}^{-2}$  (an increase of approximately 92% in the photocurrent density) higher than the pristine  $BiVO_4$ . Besides, the onset potential for  $BiVO_4/Au$  is close to 0.1 V/SCE (Fig. 12d and e). The result is due to the improvement in the photon harvesting rate, reduced charge carrier recombination rate, and better separation of charge carriers, after irradiation. Meanwhile, the formation of Au nanoparticles favors the PEC water splitting by optical enhancement and photon scattering.

In a work by Huang,<sup>34</sup> atomic vacancies are introduced onto two-dimensional  $ReS_2$  on Si by using controllable Ar ion-irradiation. The irradiated  $ReS_2/Si$  photocathode demonstrated a highly enhanced catalytic activity for PEC water splitting. A high photocurrent density of  $-18.5 \text{ mA cm}^{-2}$  at 0 V *versus* RHE was achieved in 30 seconds for the irradiated  $ReS_2/Si$  photocathode (Fig. 13a), which is higher than that for non-irradiated  $ReS_2/Si$ . Additionally, the Tafel plot shows that the photocathode of  $ReS_2/Si$ -30s possesses the lowest Tafel slope of  $73.7 \text{ mV dec}^{-1}$  (Fig. 13b). The enhanced HER performance is mainly attributed to the rich and controllable atomic Re vacancies on the  $ReS_2$  nanosheets by ion-irradiation. Also,

the impedance spectroscopy measurements indicated that the photocathode of  $ReS_2/Si$ -30s had a relatively small  $R_{ct}$  at the semiconductor/electrolyte interface, reflecting the fastest electrode-to-electrolyte shuttling of electrons during the HER.

Kumar *et al.*<sup>98</sup> synthesized thin films of hematite ( $\alpha\text{-Fe}_2\text{O}_3$ ) by electrodeposition and after 100 MeV  $Si^{8+}$  SHII, the irradiated film exhibited a significantly improved PEC response than the non-irradiated  $\alpha\text{-Fe}_2\text{O}_3$  sample. The irradiated film at a fluence of  $5 \times 10^{12}$  ions per  $cm^2$  exhibited the highest photocurrent density of  $3.0 \text{ mA cm}^{-2}$  and a hydrogen production rate of  $2.4 \text{ mL h}^{-1} \text{ cm}^{-2}$  (Fig. 13c and d). The explanation might be that the irradiation could induce a decrease in the resistivity, an increase in the donor density and an increase in the flat-band potential for the sample, which may be responsible for its enhanced PEC properties.

The sample group also found that the  $BaTiO_3$  film was a potential candidate photoelectrode for PEC water splitting due to its extraordinary stability in aqueous environments.<sup>99</sup> The fabricated  $BaTiO_3$  thin films are irradiated by 120 MeV  $Ag^{9+}$  irradiation at four different fluences ( $1 \times 10^{11}$ ,  $2 \times 10^{11}$ , and  $3 \times 10^{12}$  ions per  $cm^2$ ). At a fluence of  $1 \times 10^{11}$  ions per  $cm^2$ , the  $BaTiO_3$  film achieved a maximum photocurrent density of  $1.78 \text{ mA cm}^{-2}$  at 0.4 V *vs.* SCE (Fig. 13e), which is ten times larger than that of the pristine sample, and the maximum conversion efficiency of 0.91% was exhibited by the same photoelectrode which is about 30 times larger than that of

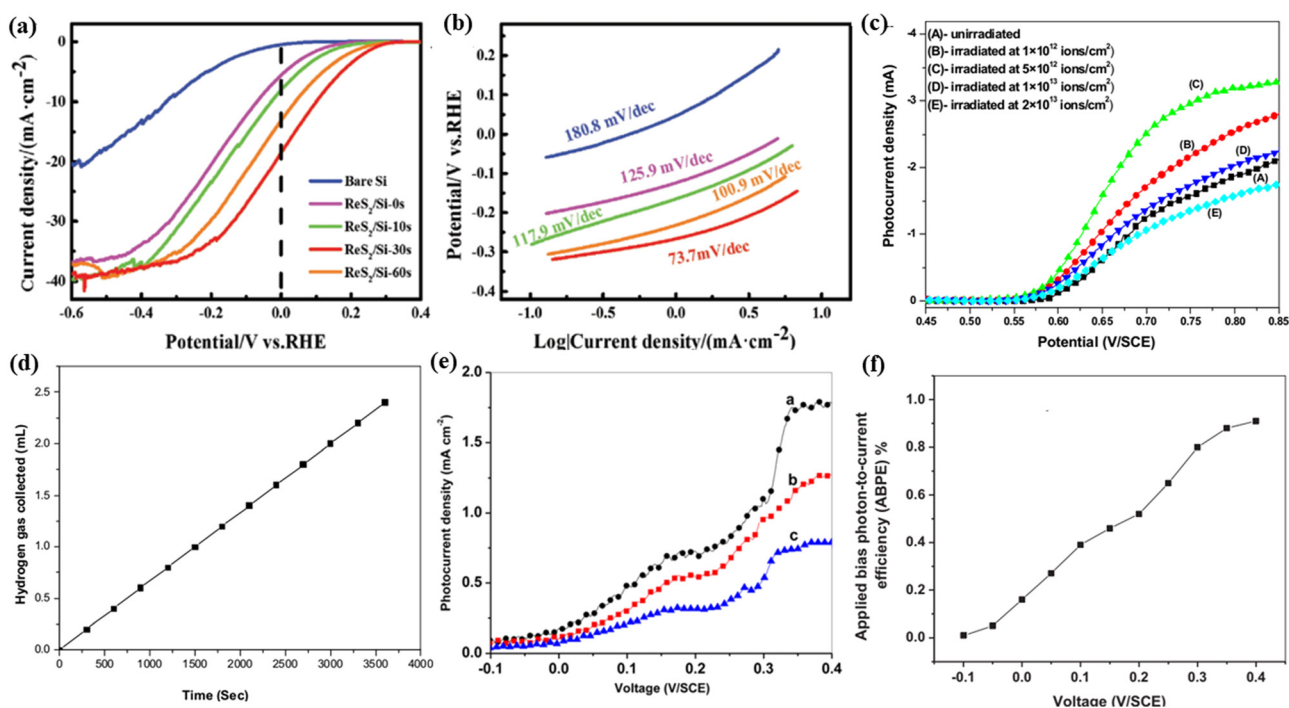


Fig. 13 PEC activity measurements of different photocathodes under  $100 \text{ mW cm}^{-2}$  light energy. (a) LSV curves and (b) Tafel plots. Reproduced with permission from ref. 34. Copyright 2018, Wiley-VCH. The PEC performance of the hematite electrode: (c) Photocurrent density *versus* applied potential curves before and after ion-irradiation and (d) volume of hydrogen generated *versus* time at 0.75 V/SCE for the sample irradiated at  $5 \times 10^{12}$  ions per  $cm^2$ . Reproduced with permission from ref. 98. Copyright 2012, Elsevier. (e)  $I$ - $V$  curve giving photocurrent densities of  $BaTiO_3$  films irradiated at (a)  $1 \times 10^{11}$ , (b)  $2 \times 10^{11}$  and (c)  $3 \times 10^{12}$  ions per  $cm^2$ . (f) Variation of the applied bias photon-to-current efficiency with an applied voltage for the film irradiated at  $1 \times 10^{11}$  ions per  $cm^2$ . Reproduced with permission from ref. 99. Copyright 2013, Elsevier.

the pristine sample (Fig. 13f). The better PEC performance by ion-irradiation is attributed to improved optical absorption capability, increased roughness, efficient separation of charge carriers, and maximum flat-band potential and donor density.

Zheng *et al.*<sup>91</sup> synthesized TiO<sub>2</sub> nanorods by the atomic layer deposition (ALD) method followed by different energies (45, 65 and 85 keV) of N<sup>+</sup> ion-irradiation. They studied the influences of the irradiated ion energy on the morphology and photocurrent density of the nanorods. And it was found that the length of nanorods increases with the increasing ion energy while the photocurrent does not increase continually with the increasing ion energy due to the heavier damage formed in TiO<sub>2</sub> irradiated by N<sup>+</sup> ions with higher energy. The nanorod array photoelectrode displays largely enhanced performance for PEC water splitting compared to that of the non-irradiated TiO<sub>2</sub> thin films; this optimized PEC performance is attributed to the effects of N<sup>+</sup> ion-irradiation, and the nanorod thin films have the larger TiO<sub>2</sub>/electrolyte interfacial area, which facilitates the hole transfer kinetics at the TiO<sub>2</sub>/electrolyte interface. This work proposes that ion-irradiation could be an effective approach for developing nanostructured materials for water splitting.

Normally, the PEC water splitting reaction process consists of another necessary reaction of the OER ( $2\text{H}_2\text{O} + 4\text{H}^+ \rightarrow \text{O}_2 + 4\text{H}^+$ ), which is also known to be the restricting step for the overall reaction. In a work by Zhan,<sup>100</sup> the surface of TiO<sub>2</sub>(110) at different setting heating plate temperatures (300 °C, 400 °C, and 500 °C) and the relationship between the geometric, electronic structure and the PEC activity were investigated by using low-energy Ar<sup>+</sup> irradiation. The irradiated TiO<sub>2</sub> photoanodes showed much enhanced photoresponse with T-500 (setting heating plate temperature 500 °C) showing the highest activity. After 6 h, the photocurrent density of the T-500 photoanode is barely changing, which indicated the high stability of T-500. These optimal results are due to two reasons: one reason is that controlled formations of the oxygen vacancy, Ti<sup>3+</sup>, Ti<sup>2+</sup>, and Ti metallic species were achieved at the surface of TiO<sub>2</sub>, rendering a significant improvement in its light absorption, and charge carrier concentration, and charge transfer properties after Ar<sup>+</sup> irradiation. And the second reason is that Ar<sup>+</sup> irradiation can change the interaction between the TiO<sub>2</sub> surface and water interface, making it more propitious to water decomposition. Thus, low-energy Ar<sup>+</sup> irradiation could significantly improve the PEC performance of the TiO<sub>2</sub> photoanode.

#### 4.3 Application of the ion-irradiation technique for photocatalysts

Photocatalysis is one of the most important topics in chemical research and has attracted tremendous interest in recent years owing to the need for renewable and sustainable energy.<sup>101–103</sup> The most used photocatalysts are semiconductors, whose performance is substantially limited by the lack of visible-light absorption and poor photo-generated charge carrier separation efficiency. Defect engineering, as an effective strategy, can introduce vacancies that improve visible light absorption and charge carrier separation efficiency.<sup>104–107</sup> The ion-irradiation

technique is a novel method to introduce vacancies; due to the cascade collision of irradiation ions with target atoms, vacancies will be introduced into target materials. Importantly, the concentration and distribution of the incident vacancy are controlled by ion fluence and energy.

Ren *et al.*<sup>108</sup> used He<sup>+</sup> ion-irradiation to efficiently introduce both defined C- and N-vacancies (V<sub>C</sub> and V<sub>N</sub>) in graphitic carbon nitride (g-C<sub>3</sub>N<sub>4</sub>) for photocatalytic water splitting. The existence of V<sub>N</sub> introduces the defect state in the band gap, which could improve the visible light absorption and increase the charge carrier separation efficiency thus resulting in the enhanced photocatalytic performance of g-C<sub>3</sub>N<sub>4</sub>. The photocatalytic performances of CN0 (pristine g-C<sub>3</sub>N<sub>4</sub> nanosheet film) and CN2 (He<sup>+</sup> ions fluence of  $1.725 \times 10^{14}$  ions per cm<sup>2</sup>) were evaluated by the photocatalytic hydrogen evolution test (Fig. 14a and b). As shown in Fig. 14b, the CN2 sample with an optimized vacancy concentration showed that a superior hydrogen evolution rate reaches 1271 mmol g<sup>−1</sup> h<sup>−1</sup> which is 2.7 and 19 times higher than the CN0 nanosheets and bulk g-C<sub>3</sub>N<sub>4</sub>, respectively. This work proved that the ion-irradiation technique is an effective approach for improving the photocatalytic performance of materials.

Besides metal-free g-C<sub>3</sub>N<sub>4</sub>, Impellizzeri *et al.*<sup>102</sup> showed the application of Fe<sup>+</sup> ion-implantation to realize an efficient TiO<sub>2</sub> photocatalyst. The increase of the Fe<sup>+</sup> irradiation fluence could enhance the damage of TiO<sub>2</sub>, which could cause strong electron-hole recombination, thus affecting the photocatalytic activity of the as-implanted film. They found that the peak intensity in the X-ray diffraction (XRD) pattern reduces gradually with the increase of the ion-irradiation fluence. The observed reduction in the peak intensity can be related to a damaging process attributed to the ion-implantation process (Fig. 14c and d). Under the fluence of  $5 \times 10^{15}$  ions per cm<sup>2</sup>, the as-implanted film revealed a remarkable photocatalytic efficiency in the degradation of organic compounds in water under visible light irradiation (80% higher than the one obtained for pure TiO<sub>2</sub> films) (Fig. 14e), without the help of any thermal treatments. They further proved that the photocatalytic activity in the degradation of organic compounds strongly depends on the number of defects induced by the ion-implantation process.

The heterojunction semiconductor is also an important sort of material in photocatalysis. In a work by Wang,<sup>109</sup> ZnO NWs were coupled with a narrow bandgap semiconductor material CdTe through a series of aqueous chemical reactions and were further irradiated with different doses of N ions to enhance the photocatalytic activity. They investigated the intensity of PL peaks at various cycles of CdTe/ZnO heterostructures, and it was found that the ZnO–CdTe-20-cycle heterostructures exhibit a lower PL peak intensity than that of other cycles ZnO–CdTe, which indicates the lowest recombination rate of photo-generated carriers. Additionally, in all samples of CdTe/ZnO heterostructures, ZnO–CdTe-20-cycle samples perform the highest photocatalytic activity. The CdTe/ZnO heterojunction nanostructures efficiently enhanced their photocatalytic performance mainly through the introduction of oxygen vacancies by N ion-irradiation, which can narrow the bandgap to improve



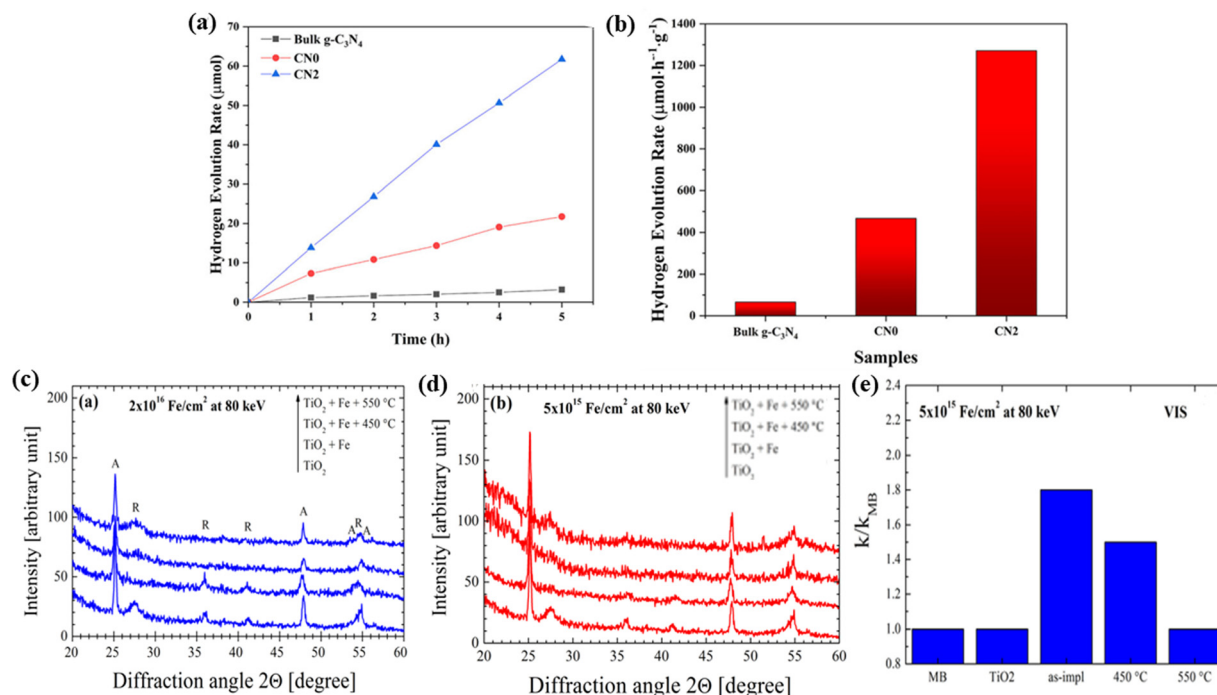


Fig. 14 (a) Hydrogen evolution curves. (b) Photocatalytic hydrogen evolution rates over the bulk g-C<sub>3</sub>N<sub>4</sub>, CN0, and CN2 samples under visible light irradiation. Reproduced with permission from ref. 108. Copyright 2019, Wiley-VCH. XRD patterns of the pure TiO<sub>2</sub> film, as-implanted and after the thermal treatment of the Fe<sup>+</sup> fluence of  $2 \times 10^{16} \text{ cm}^{-2}$  (c) or  $5 \times 10^{15} \text{ cm}^{-2}$  (d) at 80 keV. Photocatalytic rate of MB, normalized to the value obtained for MB in the absence of the photocatalyst, for different Fe<sup>+</sup> implanted films: (e)  $5 \times 10^{15} \text{ cm}^{-2}$  under UV light irradiation. Reproduced with permission from ref. 102. Copyright 2014, AIP Publishing.

the visible absorption ability and effectively separate electrons and holes.<sup>110–112</sup>

#### 4.4 Application of the ion-irradiation technique for lithium-ion batteries

Lithium-ion batteries are regarded as efficient energy storage devices, which have been widely used in every corner of life, such as smart-phones, laptops, and electric vehicles because of their high specific energy and energy densities as well as their long cycling life. Ion-irradiation has appeared as an effective technique for introducing defects into materials to improve performance. Using the ion-irradiation technology to control the evolution of defects in electrodes is an effective method to improve battery performance.<sup>113</sup>

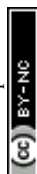
Germanium (Ge) as an anode material has drawn a great deal of interest due to its high specific capacity ( $1623 \text{ mA h g}^{-1}$ ) and Li<sup>+</sup> diffusivity.<sup>114,115</sup> Under high doses of ion-irradiation, the Ge morphology transformed from a nonporous structure to a nanoscale porosity interdigitated network.<sup>116–118</sup> When used as an anode, the network could help in relaxing stress during the cycling process. Rudawski *et al.*<sup>119</sup> fabricated ion-beam modified Ge films for lithium-ion batteries. Compared with the as-deposited Ge film, the ion beam-modified Ge nanostructured electrodes exhibited a specific charge (discharge) capacity of  $\sim 1279$  ( $\sim 1259$ )  $\text{mA h g}^{-1}$  with a coulombic efficiency of 98.4% for the first cycle. After 25 cycles, the specific capacity slightly decreases to  $1352 \text{ mA h g}^{-1}$  which is much higher than that of the deposited film ( $\sim 200 \text{ mA h g}^{-1}$ ) (Fig. 15a).

When the cycling rate increased to C/0.9, the specific capacities were still greater than  $1000 \text{ mA h g}^{-1}$  (Fig. 15b). The result suggests that the ion beam-modified Ge nanostructured electrode has great promise for use in high performance Li ion batteries.

Different from the above mentioned, Rahman *et al.*<sup>120</sup> first utilized *in situ* high-energy Kr ion-irradiation combined with transmission electron microscopy to monitor how the defects and microstructures evolve in Na and Li-layered cathodes. The phase pure crystal structure along with the electrochemical performance shows that Na<sub>2/3</sub>Fe<sub>1/2</sub>Mn<sub>1/2</sub>O<sub>2</sub> and LiNiO<sub>2</sub> are representative and can provide a valuable guideline for designing stable layered cathodes under extreme conditions, such as outer space exploration and nuclear power industries. The quantitative mathematical analysis of the dynamic bright-field imaging shows that defect clusters preferentially align along the Na/Li ion diffusion channels (*a-b* planes), which is likely governed by the formation of dislocation loops. This study provides a valuable guideline for designing stable layered cathodes under extreme conditions, such as outer space exploration and nuclear power industries.

#### 4.5 Application of the ion-irradiation technique for supercapacitors

Besides batteries, supercapacitors can also serve as energy storage devices, which usually include electrical double layer capacitors (EDLCs) and pseudo-capacitors (PC). Compared with batteries, supercapacitors can provide higher power density



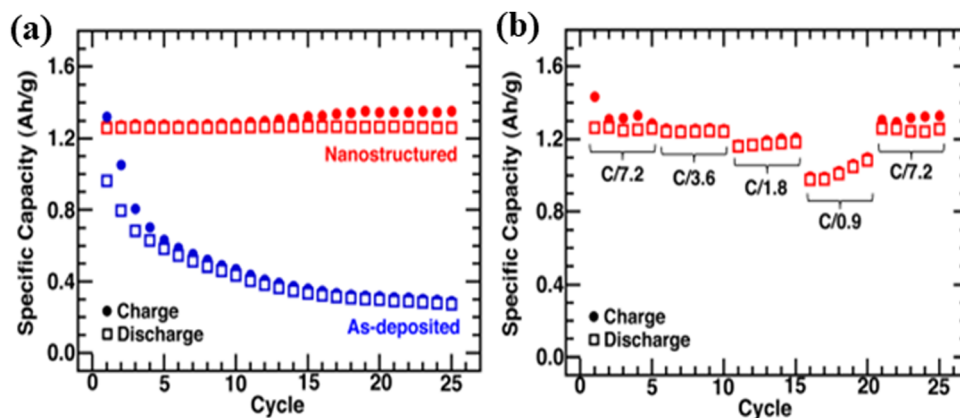


Fig. 15 (a) Specific capacity versus cycle number for a nanostructured Ge electrode cycled at a C/7.2 rate for 25 cycles; the data of the as-deposited Ge electrode cycled at the same rate is provided for comparison. (b) Specific capacity versus cycle number for the nanostructured Ge electrode cycled sequentially at C/7.2, C/3.6, C/1.8, C/0.9, and C/7.2 rates, respectively. Reproduced with permission from ref. 119. Copyright 2012, AIP Publishing.

and long cycling life. However, the restriction of charge storage on the electrode surface makes their capacity much lower than batteries.<sup>121–123</sup> Among different electrode materials, electroactive polymers generally show a higher specific capacitance due to their ability to store charges by faradaic reactions on the surface.<sup>124</sup> Interestingly, the ion-irradiation technique has been proved to be effective in improving the electrical conductivity and generating active sites for the surface physical and chemical modification of the polymers.<sup>125</sup> The irradiated polymers could form new chemical bonds, intermolecular crosslinking, chain scissoring or fragmentation of molecules, bond-breaking, and

formation of unsaturated groups by ion-irradiation, thus improving the performance of supercapacitors.<sup>126</sup>

Hussain *et al.*<sup>127</sup> electrochemically synthesized LiClO<sub>4</sub> doped polypyrrole thin films on an indium tin oxide coated glass substrate, which was subsequently ion-irradiated by 160 MeV Ni<sup>12+</sup> ions at different fluences of  $5 \times 10^{10}$ ,  $5 \times 10^{11}$ , and  $3 \times 10^{12}$  ions per cm<sup>2</sup>. The irradiated polymer films demonstrated an increase in the magnitude of oxidation and reduction peak current (Fig. 16a), which is mainly due to the increase in the conductivity of the polymer films. Then, the conductivity of the polymer films was investigated, and

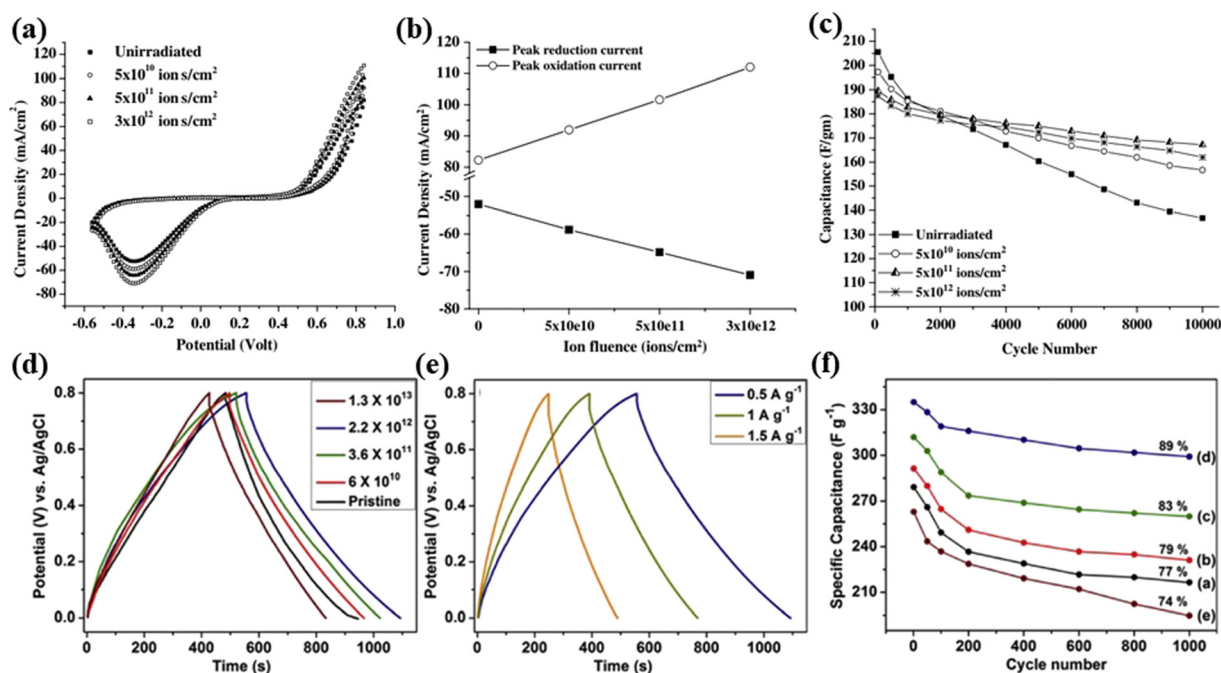


Fig. 16 (a) Cyclic voltammograms of polypyrrole films doped with LiClO<sub>4</sub> before and after ion-irradiation at a potential 50 mV s<sup>-1</sup> scan rate. (b) Peak oxidation and reduction currents versus ion fluence. (c) Stability plots of supercapacitors with LiClO<sub>4</sub> doped polypyrrole electrodes. Reproduced with permission from ref. 127. Copyright 2005, Elsevier. (d) GCD curves of the pristine and ion-irradiated RGO-PPyNT nanocomposites with different fluences. (e) GCD curves of RGO-PPyNT nanocomposites irradiated at a fluence of  $2.2 \times 10^{12}$  ions per cm<sup>2</sup> at different current densities. (f) Variation of the specific capacitance with cycle number of RGO-PPyNT nanocomposites. Reproduced with permission from ref. 121. Copyright 2018, Elsevier.



the results indicated that the conductivity of the films shows an increasing trend with an increase in the ion fluence. The irradiated film exhibits a higher conductivity ( $150\text{--}160\text{ S cm}^{-1}$ ) than the pristine film ( $100\text{ S cm}^{-1}$ ), which is attributed to bond breaking that occurs due to cross-linking and chain scission which creates a large number of free electrons and thus increases in the conductivity of the polymer films. A plot of the oxidation peak current and reduction peak current (Fig. 16b and c) shows that the peak current of oxidation and reduction increases linearly with an increase in the fluence. In addition, the irradiated (fluence  $5 \times 10^{11}$  ions per  $\text{cm}^2$ ) polymer film exhibits a higher capacitance than non-irradiated films after 10 000 cycles.

Devi *et al.*<sup>121</sup> synthesized hybrid nanocomposites of reduced graphene oxide (RGO) and polypyrrole nanotubes (PPyNTs). The electrical properties of RGO-PPyNTs have been impacted by SHII with 85 MeV  $\text{C}^{6+}$  ions at different fluences. The galvanostatic charge-discharge measurements (Fig. 16d and e) show a nearly linear and symmetric discharge-charge curve indicating

a typical EDLC behavior. Besides the highest fluence ( $1.3 \times 10^{13}$  ions per  $\text{cm}^2$ ), the discharge duration of the RGO-PPyNT nanocomposite is slightly prolonged with the increase of irradiation fluences, which indicates an increase in the specific capacitance of the nanocomposites up to  $346\text{ F g}^{-1}$  at a fluence of  $2.2 \times 10^{12}$  ions per  $\text{cm}^2$ . After 1000 cycles, the irradiated nanocomposite retains 89% initial capacity which is higher than that of the non-irradiated nanocomposite (Fig. 16f). It was believed that the ion-irradiation increases the specific surface area and pore size of the electrode, which extends the ion transport from the surface to the bulk of the electrode, and results in an increase in the performance of supercapacitors. The results suggest that ion-irradiation has a great potential for modifying electrodes in supercapacitors.

## Summary and outlook

Ion-irradiation shows great promise for the development of electrode materials for energy conversion and storage devices

**Table 1** Summary of the application of ion-irradiation/implantation technology in energy conversion and storage of (photo) catalyst/electrode materials from recent reports in the literature

| Catalyst/electrode materials                                        | Ion species | Energy           | Fluence (ions per $\text{cm}^2$ )                                                                                                                            | Roles              | Application              | Ref. |
|---------------------------------------------------------------------|-------------|------------------|--------------------------------------------------------------------------------------------------------------------------------------------------------------|--------------------|--------------------------|------|
| Single layer $\text{MoS}_2$                                         | Au          | 500 keV          | $5 \times 10^{11}$<br>$1 \times 10^{14}$                                                                                                                     | Defect             | HER                      | 33   |
| $\text{Bi}_2\text{Te}_3$ nanosheets                                 | Fe          | 320 keV          | $4 \times 10^{14}$                                                                                                                                           | Defect             | HER                      | 35   |
| PtPb nanoplates                                                     | C           | 3 MeV            | $1, 2, 3 \times 10^{16}$                                                                                                                                     | Defect             | MOR<br>EOR<br>ORR<br>MOR | 18   |
| PtPb nanoplate                                                      | Kr          | 1 MeV            | $6 \times 10^{13}$<br>$1 \times 10^{15}$<br>$5 \times 10^{15}$                                                                                               | Defect             | MOR                      | 36   |
| $\text{Co}_3\text{O}_4\text{-O}_x/\text{CoO}/\text{Co}_3\text{O}_4$ | Ar          | 25, 50 keV       | $1, 2 \times 10^{17}$                                                                                                                                        | Defect             | OER                      | 38   |
| ZnO NPs/thin films                                                  | Zn          | 60 keV           | $1 \times 10^{16}$                                                                                                                                           | Doping             | —                        | 40   |
| N-Doped graphene (NG)                                               | N           | 40 keV           | $1 \times 10^{16}$                                                                                                                                           | Doping             | —                        | 41   |
| Polystyrene sphere/hemisphere                                       | Ag          | 15 kV            | $1, 3, 5, 7 \times 10^{16}$                                                                                                                                  | Morphology control | —                        | 48   |
| C-SiC coaxial nanotube                                              | Si          | 200 keV          | $1.1 \times 10^{21}$                                                                                                                                         | Synthesis          | —                        | 50   |
| $\text{WO}_{3-x}$ nanowire                                          | Ar          | 80, 130, 190 keV | $1, 1.5 \times 10^{17}$                                                                                                                                      | Synthesis          | —                        | 52   |
| $\text{MoSe}_2/\text{CC}$                                           | Ar          | 0–100 keV        | $5 \times 10^{14}$<br>$5 \times 10^{15}$<br>$5 \times 10^{16}$                                                                                               | Defect             | HER                      | 66   |
| PS-Ni, NPS-Ni thin films                                            | Ar          | 90 keV           | $1, 2 \times 10^{17}$<br>$5 \times 10^{15}$                                                                                                                  | Synthesis          | HER                      | 67   |
| NiO/CC                                                              | N           | 0–100 keV        | $5 \times 10^{11}$                                                                                                                                           | Defect             | OER                      | 11   |
| $\text{TiO}_2$ thin films                                           | Ag          | 120 MeV          | $1, 5 \times 10^{12}$<br>$1 \times 10^{13}$<br>$1 \times 10^{12}$<br>$1 \times 10^{13}$<br>$2 \times 10^{13}$<br>$2 \times 10^{15}$                          | Defect             | PEC                      | 96   |
| Cr- $\text{Fe}_2\text{O}_3/\text{CuO}$ thin films                   | Si          | 100 MeV          | $1 \times 10^{12}$<br>$5 \times 10^{12}$<br>$1 \times 10^{13}$<br>$2 \times 10^{13}$<br>$2 \times 10^{15}$                                                   | Defect             | PEC                      | 97   |
| $\text{BiVO}_4/\text{Au}$ photoanode                                | N           | 600 keV          | $1 \times 10^{11}$                                                                                                                                           | —                  | PEC                      | 88   |
| $\text{BaTiO}_3$ thin films                                         | Ag          | 120 MeV          | $1 \times 10^{17}$                                                                                                                                           | Defect             | PEC                      | 99   |
| $\text{TiO}_2$ nanorods                                             | N           | 45, 65, 85 keV   | $9.6 \times 10^{17}$                                                                                                                                         | Synthesis/defect   | PEC                      | 91   |
| $\text{TiO}_2(110)$                                                 | Ar          | 1 keV            | $1.725 \times 10^4$                                                                                                                                          | Defect             | PEC                      | 100  |
| g- $\text{C}_3\text{N}_4$                                           | He          | —                | $1, 5 \times 10^{13}$<br>$1, 5 \times 10^{14}$<br>$1.0 \times 10^{16}$<br>$6.25 \times 10^{14}$<br>$1.25 \times 10^{15}$                                     | Defect             | Photocatalysts           | 108  |
| $\text{CdTe}/\text{ZnO}$ heterostructures                           | N           | 190 keV          | $5 \times 10^{10}$<br>$5 \times 10^{11}$<br>$3 \times 10^{12}$<br>$6 \times 10^{10}$<br>$3.6 \times 10^{11}$<br>$2.2 \times 10^{12}$<br>$1.3 \times 10^{13}$ | Defect             | Photocatalysts           | 109  |
| Ge electrodes                                                       | Ge          | 260 keV          | —                                                                                                                                                            | Synthesis          | LIB                      | 110  |
| $\text{Na}_{2/3}\text{Fe}_{1/2}\text{Mn}_{1/2}\text{O}_2$           | Kr          | 5 kV             | —                                                                                                                                                            | —                  | LIB                      | 120  |
| $\text{LiNiO}_2$                                                    | Ni          | 160 MeV          | —                                                                                                                                                            | —                  | Supercapacitor           | 127  |
| Polypyrrole polymer thin films                                      | Ni          | 160 MeV          | —                                                                                                                                                            | —                  | Supercapacitor           | 127  |
| RGO-PPyNT nanocomposites                                            | C           | 85 MeV           | —                                                                                                                                                            | —                  | Supercapacitor           | 121  |



such as water electrolysis cells, lithium-ion batteries and supercapacitors. The recent efforts of ion-irradiation are mainly devoted to the synthesis and modification of electrode materials. The effects of ion-irradiation on the materials include defect introduction, element doping, morphology control, and materials synthesis. For the ion-irradiation technique, the synthesis strategy is relatively simple and is usually used for fabricating nano-materials of semiconductors, carbides, and metal oxides. To date, the catalyst/electrode materials related to ion-irradiation for energy storage and conversion technologies have been involved in electrocatalysis, photoelectrochemistry, photocatalysis, lithium-ion batteries, and supercapacitors. Some catalyst/electrode materials that confer significant improvements by the ion-irradiation technique are summarized in Table 1. A large number of experimental results have proved that the ion-irradiation technique is undoubtedly an effective and promising technology. In addition, ion beam technology is undoubtedly a potential and efficient tool that can be extended to many other fields. For the ion-irradiation technique, opportunities and challenges coexist. With the development in materials science and the progress of characterization technologies, the ion-irradiation technique will obtain great achievements and some more exciting discoveries can be expected in the future. Whereas there still exist challenges that need to be overcome in the following directions:

(1) Ion-irradiation technology can help us to better establish the structure–activity relationship between the defect concentration and material catalytic performance. However, it is necessary to further investigate the mechanism of interaction between the ion-irradiation process and structure change. Some *in situ* characterization techniques such as *in situ* XRD and Raman spectra monitoring can be used during ion-irradiation. Further analysis and research of defect evolution can lay a solid foundation for the future utilization of the ion-irradiation technique for improving material performance.

(2) A comprehensive and in-depth understanding of the functional mechanisms of ion-irradiation using both experimental measurements and theoretical modeling and simulation (such as DFT calculations) for new electrode/electrolyte material design and synthesis. Although the ion-irradiation can result in material surface structure changes for improving the performance, the relationship between the structure and performance is still not fully understood at present. Therefore, some fundamental mechanism understanding is definitely needed. In addition, although ion-irradiation has a positive effect on electrode materials, it also causes different degrees of damage to the materials, and it is equally important to further study the relationship between damage degree and material properties.

(3) For now, the ion-irradiation technology is used in the study of catalytic materials, which can efficiently form a variety of noble metal nanostructures and achieve low-cost preparation with high atomic utilization in the preparation of precious metal-based catalytic materials. The application of ion-irradiation in single atom catalysts will be the direction of the future research directions and development trends.

## Author contributions

Yahui Wang: conceptualization, data curation, and writing – original draft preparation. Xing Cheng and Kai Zhang: writing – review. Ge Chen: methodology, writing – review, and editing. Ruzhi Wang and Jiujun Zhang: supervision.

## Conflicts of interest

There are no potential conflicts of interest to declare.

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