Review

Recent advances in the electrocatalytic application of transition metal nitrides nanocrystalline

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Abstract: The energy crisis and environmental problems are becoming more and more severe due to the long-term consumption of fossil fuels. Therefore, energy conversion devices with high energy density and environmental friendliness (such as fuel cells, metal-air batteries, etc.) have been expected to be reliable alternatives to traditional fossil energy. However, due to the inevitable use of precious metals as the electrode catalysts for these devices, the popularization of these alternatives is seriously hindered. Transition metal nitrides (TMNs) exhibit similar surface and adsorption properties to noble metals since the atomic distance between metal atoms increases and the d-band center of metal atoms downshift after the nitrogen atoms enter the metal lattice. TMNs have become one of the best electrode materials to replace noble metal electrocatalysts in energy storage and conversion devices. In this review, the latest development in the electrocatalytic application of TMNs nanocrystalline is covered. Firstly, we briefly discuss the structure and activity origin of TMNs and introduce the common physical and chemical methods for the preparation of TMNs. Subsequently, we illustrate the applications of unary TMNs and multivariate TMNs in oxygen reduction reaction (ORR) and oxygen evolution reaction (OER). Finally, we summarize the challenges and problems of TMNs encountered at the present stage, and expect its future development.

Keywords: Transition metal nitrides, Preparation methods, Electrocatalytic application, Oxygen reduction reaction, Oxygen evolution reaction.

1. Introduction

With the rapid development of society and the economy, the demand for energy is increasing in recent years. The environmental pollution caused by the use of fossil fuels is becoming increasingly severe. Human beings are facing unprecedented energy shortage and environmental pollution problems.[1-3] Therefore, human society is now facing the challenge to get rid of the dependence on traditional fossil energy, develop new and efficient energy conversion devices, and transform the energy mode. At the same time, the development of energy shortage and conversion devices, such as fuel cells and metal-air batteries, has also provided strategies to solve the energy shortage and environmental pollution.[4-6] Taking the fuel cell as an example, on account of it uses hydrogen as the fuel and the reaction product is water, it has almost no pollution to the environment. Besides, because it gets rid of the restrictions of the Carnot cycle, it also shows a high energy conversion efficiency. Therefore, the fuel cell has become a hot research object for the researchers from many fields.[7, 8] For both fuel cells and metal-air batteries, their internal reaction is generally composed of two parts, one is the fuel oxidation reaction in the anode, the other is ORR in the cathode, and these two reactions occur all need suitable electrocatalysts to reduce reaction barrier.[9, 10] At present, the commercial application of fuel cells and other new energy devices are based on Pt-based materials loaded with carbon black
as the electrocatalysts for both cathode and anode, but due to the scarcity of Pt resources and high price, the cost of these new energy conversion devices is expensive, which significantly restricts their industrialization.[11, 12] In particular, compared with hydrogen oxidation reaction (HOR) in the anode, ORR in the cathode involves two-electron and four-electron transfer process of oxygen, it has higher reaction barrier, more complex reaction process and slower reaction rate. For example, the rate of ORR in the cathode is generally only 1/10 the speed of HOR in the anode in a fuel cell, so the loading of Pt in the cathode is even more.[13-17] Therefore, how to reduce the cost of electrocatalysts and find low-cost electrocatalyst materials to replace precious metals electrocatalysts are of vital importance for the development, application, and popularization of these novel energy conversion devices.

Thanks to the metal-metallic bond rearrangement, the formation of the new chemical bond between the metal and non-metal atom, and the electron transfer between metal and non-metal atoms, TMNs nanocrystal materials are characterized by covalent compounds, ionic crystals, and transition metals, and have great application value in catalysis, energy storage and conversion, super-hard materials, semiconductors and superconductors.[18, 19] Due to the introduction of highly electronegative nitrogen element, the nitrogen atoms fill in the interstitial places of the metal cells, thereby TMNs form interstitial compounds or alloy structures.[20] Besides, nitrogen atoms enter into the interstitial sites of nanocrystal cell, which results in the increase of the transition metal atoms spacing, the expansion of the lattice, the shrinkage of transition metal d-band, and the increase of Fermi level state density. And the redistribute of Fermi level state density makes their electrocatalytic activity and surface properties different from those of the parent metal, and are similar to the eighth group precious metals (such as Pt).[21] These changes give the TMNs excellent conductivity, electrocatalytic activity and optical properties, couple with its corrosion resistance and high-temperature resistance, so that TMNs are available to be used in electrocatalysis, petroleum hydrodesulphurization, ammonia synthesis and decomposition, hydrogen production catalysis, and other fields.[22, 23]

In recent years, researchers have found that downsizing materials to the nanoscale can often give them unusual mechanical, electrical, and optical properties.[24, 25] Hence, the study of nanostructured electrocatalyst materials has aroused great interest. Besides, due to the high price of Pt, the U.S. Department of Energy set the goal of electrocatalyst in 2017, namely, improving the ORR activity of fuel cell cathode electrocatalyst, and further reducing the amount of Pt loaded.[26] Therefore, the design idea of using tiny nanoparticles as electrocatalysts has gradually been considered by the researches. As all we know, the ideal electrode materials should also have an excellent ability for electronic conductivity and ionic conductivity, the design idea of using nanoparticles as the materials for electrocatalyst can not only significantly reduce the consumption of rare and expensive metals, but also improve the ionic conductivity of materials by shortening the diffusion length of ions in catalytic materials. Therefore, the design of nanostructures for electrode materials can significantly improve the electrical properties of materials.[27] Moreover, when the electrocatalyst is discrete nanoparticles or polycrystalline thin films, the dynamics of material activity can be changed almost one order of magnitude. The unique nanostructure will provide new features of the electrocatalyst material surface, which can significantly improve the activity of electrocatalyst materials, and lead to large specific surface area and more adsorption and reaction sites.[26] Therefore, the design of TMNs nanocrystalline is an important research field for electrode catalysts of energy shortage and conversion devices. In this paper, some preparation methods and structural characteristics of TMNs nanocrystals are reviewed. Their applications (ORR and OER) to electrode materials of some new energy devices are briefly reported and summarized. Finally, the development opportunities and challenges of TMNs nanocrystals in the future are discussed.
2. Preparation methods and structural properties of TMNs

Due to its excellent performance as a highly effective electrocatalyst, TMNs have undoubtedly attracted the attention and interest of the scholars. However, the study and discussion on their preparation are still relatively few at present, and the synthetic strategies are relatively limited.[27] The synthetic methods of TMNs can be mainly divided into physical methods and chemical methods (Figure 1), the former include laser ablation, sputtering, arc discharge, physical vapor deposition, etc., and the latter include direct nitriding of transition metals, nitriding of transition metals oxides, ammonolysis of transition metals chloride, solvothermal method, thermal decomposition of polymer precursors, etc.[28]

2.1 Physical synthetic methods

The primary method used for most physical syntheses is deposition, and the sample is mainly in the form of thin film; that is, different physical means are used to load the reactants onto the substrate. Physical vapor deposition (PVD) technology is a relatively popular deposition method at the present stage. PVD refers to using physical processes (such as thermal evaporation of material) to make the surface of source material vaporize, under vacuum conditions, atoms, molecules or partially ionize ions which are then deposited to form a film with some special functions on the surface of the substrate through low pressure, thus realizing the control of atom transfer from source materials to coatings.[29]

PVD technology is often combined with some other physical means (such as sputtering, ion irradiation, etc.) to synthesize TMNs. Sputtering is the most commonly used physical synthesis method due to its high deposition rate, which requires the sputtering gas (argon) and the reaction gas (nitrogen) to prepare high purity thin film samples with controllable stoichiometry and composition by PVD. For instance, Zhu et al.[30] successfully deposited cubic, wurtzic, and explosive high-pressure phases boron nitride films on the metal alloy substrates by tuned substrate radio frequency magnetron sputtering and PVD.
techniques. The percentage of cubic boron nitride phase in the film was about 50% as calculated by Fourier transform infrared measurements. Compared with the cubic phase formed by the traditional low energy ion bombardment, the cubic boron nitride phase prepared by this method had a very low internal stress, which could perfectly solve poor adherence and delamination from the substrates. Wei et al.[31] equipped bimetallic carbon paper supported vanadium-molybdenum nitride (MoVN) thin films by magnetron co-sputtering. Murthy et al.[32] also designed a series of Mo3N2, Ag-Mo3N2, V-Mo3N2 and CuMo3N2 films by magnetron co-sputtering technique. Kang et al.[33] prepared NiN thin films with cauliflower shape and tetrahedral crystal lattice by reactive sputtering in N2 atmosphere.

In addition to sputtering, PVD is also used in combination with ion irradiation, known as ion mixing and vapor deposition (IVD), which can easily change and design the physical and chemical properties of thin films. For example, Uchida et al.[34] used the IVD technique to prepare Titanium aluminum nitride (Ti, Al)N films by depositing Ti and Al metal vapor under simultaneous irradiation N ions. And Liu et al.[29] also successfully prepared a novel quaternary (Ti, Al, Zr)N coating on Si3N4 ceramic substrates by using PVD technology and multi-arc ion plating technique. When the gas pressure was 2.5 Pa, the (Ti, Al, Zr)N coated Si3N4 cutting tools had the most extended lifetime and the best mechanical performance. Although these physical methods can give a definite structure to TMNs, for the majority of TMNs synthesis, the conditions of physical synthetic methods are harsh, the preparation process is cumbersome and inefficient, so it is not suitable for the general synthesis of TMNs.[35]

2.2 Chemical synthetic methods

The chemical synthesis of TMNs is generally divided into two steps. Firstly, the precursors of TMNs are synthesized with different methods. Then the metal precursors are nitrided with NH3 or N2/H2 mixed gases at different temperatures to obtain various forms of TMNs.[36] Therefore, according to the different types of precursor materials, the chemical process for TMNs synthesis can be divided into the transition metal oxide method, transition metal chloride method, metal-organic frameworks method, and other methods.

2.2.1 Transition metal oxide method

Transition metal oxides (TMOs), as the most common compound form of transition metal, are nitrided to prepare TMNs, which is relatively common way to synthesize TMNs at present. [37] The specific experimental procedure is to oxidize the transition metal into TMOs, or directly use TMOs crystals before hot ammonia reduction.[38] The calcination temperature is usually between 300-800 °C, so that TMNs of different dimensions can be obtained. Due to the fact that the oxygen atom is used to replaces the nitrogen atom in the lattice, the lattice volume shrinks. Hence, it is easy to generate many mesoporous structures on the surface of TMNs prepared by TMOs. This method can be used to prepare mesoporous TMNs. For example, Peng et al.[39] nitrided TiO2 nanotubes array into TiN, and used the resulted porous TiN nanotube array as the core to prepare TiN-core porous double-layer MoOx (MoOx/TiN/MoOx) nanotube array with high conductivity. The porous structure facilitated the electrolyte infiltration, and the maximizing of exposure to the reaction sites of the material. Simultaneously, the high conductivity of TiN gave the material a high specific capacitance. Nagai et al.[40] used a temperature-programmed reaction to convert MoOx to Mo3N in the atmosphere of NH3 and N2/H2 mixture. The resulted electrocatalyst showed an irregular surface with abundant nanopores and high specific surface areas, which had excellent electrocatalytic effect in hydrodenitrification, hydrodesulfurization, hydrolysis and hydrogenation. Similarly, Jiang et al.[41] transformed MoOx nanowires into mesoporous Mo3N nanowires with cubic crystal nitrogen-vacancy after annealing at 800 °C in NH3 atmosphere for 6 hours (Figure. 2a, b). It was found that the mechanism of sodium-ion storage devices was a kind of surface redox reaction, so Mo3N nanowires with a large specific surface area used as the electrode material of the
sodium ion storage device would have excellent specific capacity and cycle durability (Figure. 2c, d). In another report, Yang et al.[42] synthesized VN, TiN, NbN, and Ta3N5 by passing ammonia through a pipeline containing the oxides of V, Ti, Nb, and Ta under the protection of argon gas at 450-800 °C. However, these preparation operations are often time-consuming. To optimize the synthesis process of TMNs and shorten the synthesis time, Cheng et al.[43] studied a new rapid nitriding process. They synthesized a series of ordered mesoporous TMNs by using mesoporous TMOs. This ordered mesoporous structure promoted the interaction between ammonia and metal oxides precursors, and effectively shortened the nitriding time. It was pointed out that rapid nitriding also inhibits the closure and collapse of mesoporous structures. Besides, the researchers found that amorphous TMOs could also be directly nitrided to produce TMNs. For example, Wang et al.[44] prepared metastable MoO$_2$ using hydrazine reduction (NH$_4$)$_6$Mo$_7$O$_{24}$ solution. The amorphous MoO$_2$ was converted into face-centered cubic γ-Mo$_2$N at 400 °C ammonia flow, and into hexagonal σ-Mo$_2$N at the different temperature of 600 °C.

The method to prepare TMNs using TMOs as a precursor is relatively mature and widely used; however, general TMOs show relatively more crystal forms. This kind of preparation process usually involves high-temperature conditions. When heated at high temperature, TMOs tends to undergo crystal transformation or transition between crystalline and amorphous states, resulting in incomplete nitriding of the transition metal, and even affecting the final structure of TMNs.

2.2.2 Transition metal chloride method

As the reaction temperature of transition metal chloride and ammonia is lower than that of TMOs, it is also widely used for the preparation of TMNs. For example, Choi et al. [45] prepared tantalum and tungsten nitrides through a two-step process. Firstly, the tan-
tantalum and tungsten chloride was ammonolyzed in anhydrous chloroform at room temperature. Then, the prepared powder was heat-treated in an ammonia atmosphere to obtain tantalum nitride (TaN and Ta₂N) and tungsten nitride (WN). Afanasiev et al. [46] also prepared alkaline-earth chloride nanoparticles through chloridizing scheelite with CCl₄, which were then ammonolyzed to obtain MoN and WN. In this manner, the transition metal nitriding could be carried out at a suitable temperature (500-550 °C) which was much lower than the temperature for the traditional ammoniation of TMOs using pure NH₃ or N₂/H₂ mixtures, and technically easy to achieve. In addition, the transition metal chloride method could also be used to prepare transition metal-nitride nanocomposites. For example, Cheng et al. [47] reported a method for the preparation of transition metal Si₃N₄ composite nanomaterials. They designed the Si₃N₄ complex with palladium chloride and silicon diimide gel as raw materials, and then prepared Pd-Si₃N₄ nanocomposites with high specific surface area and mesoporous structure. This method could also be used for the preparation of other transition metal Si₃N₄ composites, such as Ni, Co, Zr, and Ru. Qiu et al. [48] also reported a new method in which they synthesized various metal nitrides (such as CrN, Fe₃N, etc.) using metal-urea chloride as precursors, and this study also facilitated the understanding of the feasibility of metal-urea chloride as a low-temperature synthesis TMNs. Besides, the reaction mechanism was the nucleation growth mechanism, and the use of the metal-urea complex as a precursor to prepare more complex TMNs was also considered adequately. For example, Weil et al. [49] reported a method to prepare TMNs powders and coatings, they dissolved transition metal chlorides in solvents, such as acetonitrile, and reacted with alkanolamine to form a viscous chelating solution as a precursor, and then generated TMNs through heat treatment.

While using ammonium chloride solution synthesis, the temperature needed by TMNs is lower than those of the transition metal oxide nitride method. Nevertheless, the time it takes to ammonium chloride solution is often long. The synthesis process involves the utilization of a large number of organic reagents, steps and process trivial. At this stage, it is only applicable to be used in the lab to synthesize small-scale TMNs for research.

2.2.3 Metal-organic frameworks method

In addition to transition metal oxides and chloride, metal-organic frameworks (MOFs) are also commonly used as precursors for TMNs. MOFs are crystalline porous materials with periodic network structure formed by the self-assembly of transition metal ions or metal clusters with organic ligands through coordination bonds or other forces. Due to the unique porous structure of MOFs, TMNs prepared by MOFs as precursor materials generally show a relatively rich pore structure. [50-52] As shown in Figure. 3a-c, CoN with various structures were prepared through different MOFs precursors, and such MOFs-derived TMNs all maintained a common porous structure feature. By the way, in terms of the application of electrocatalysts, a rich pore structure can increase the exposure site of the active center, and improve electrocatalysts’ electrocatalytic activity. In recent years, there are many reports on the preparation of TMNs by MOFs. For example, Cheng et al. [53] obtained porous iron nitride (FeN) nanoparticles using cubic crystal MOFs (Prussian blue) as precursors by rapid nitridation process. The FeN samples retained the precursors' structure, and only contracted nominally during the transformation process, thus obtaining a high specific surface area. Zhu et al. [54] synthesized porous CoFeNₓ/N doped carbon nanotube arrays supported by a carbon cloth through simple nitriding of MOFs (ZIF-67) at room temperature. The synthesized samples showed hierarchically porous structures, which increased the exposure of the activity sites, accelerated the charge/mass transfer, and thus improved the electrocatalytic activity. Moreover, Hu et al. [55] reported a preparation method of manganese nitride (MnN) by cracking manganese triazole in an oxygen-free environment. Wang et al. [56] obtained Co@Co₂N/MnO-NC by cracking the Mn-containing molecular sieve-imidazole framework. Liu et al. [57] synthesized FeN using the iron foundation organic frameworks as the hard templates. However, as one of the
most essential advantages of MOFs, high porosity can also bring a few disadvantages. For instance, in the final stage of synthesis, the organic groups of MOFs disappear with nitriding, leaving their spatial structure only, and the resulted TMNs can still be classified as ceramics. Therefore, the porosity means that both the density and the mechanical strength are reduced, which makes it easily destroyed by external forces in practical application. This still needs to be optimized.

Figure 3 (a) Schematic of the synthesis route of Co4N@NC PNCs; (b) Schematic diagram of the formation of porous CoxNiyN composites; (c) Schematic illustration of the synthesis process of mesoporous Co3N@amorphous N-doped carbon NCs. Redrawn with permission of ref. 50, 51 and 52. Copyright 2018 and 2019 American Chemical Society. Copyright 2019 Tsinghua University Press.

2.2.4 Other chemical methods

In addition to the methods mentioned above, many other chemical synthetic ways have been used by researchers to synthesize TMNs. For example, Barker et al.[58] dissolved the metal in liquid zinc, and then the solution reacted with nitrogen to produce binary and ternary nitrides (e.g., CaZnN2). Xiao et al.[59] synthesized two-dimensional MnN2 by lattice matching method. Yao et al.[60] used urea as a nitrogen source to synthesize various transition metal nitride compounds (TiN, NbN, Mo2N, W2N, NbC.N1-x) in different conditions. In conclusion, various nano-morphology TMNs have been prepared by scholars using other precursors and different nitriding methods. There are still many feasible approaches, which need to be discussed in the future.

3. Transition Metal Nitrides as electrocatalyst

Transition metal nitrides are used as the electrocatalysts due to their stable structure, high conductivity, and strong anti-poisoning ability.[61, 62] On one hand, as a kind of interstitial nanocrystal material, TMNs have the characteristics of covalent compounds, ionic crystals, and transition metals due to the rearrangement of metal-metal bonds, the formation of metal and non-metal atoms, and the electron transfer between metal and non-metal atoms. On the other hand, the highly electronegative nitrogen species in the gap results in the increase of the metal atom spacing, shrinkage of the transition metal d-band as well as increase of the Fermi level state density. Therefore, the surface properties and adsorption characteristics of some TMNs (Ti, Ni, Fe, Co, Mo, etc.) are similar to those of noble metal. [63-65] Thus, TMNs have high performance in ORR, OER, and other fields.
3.1. Transition metal nitrides as ORR electrocatalyst

At present, the oxygen reduction electrocatalytic process mainly relies on the Pt-based electrocatalyst. However, due to the shortage of Pt resources, and the continuous increase of prices, the dependence of energy conversion devices on precious metals has greatly restricted its industrialization process. TMNs are considered to be one of the most promising alternatives for noble metal ORR electrocatalysts due to their low price, abundant reserves, and comparable electrocatalytic performance.[66, 67]

3.1.1 Unary TMN

TMN was first discovered as ORR electrocatalyst in 1964.[68] Jasinski experimentally found that cobalt, cooperating with nitrogen, showed activity for ORR. Since then, TMNs have been widely studied. Yang et al.[69] put forward a simple method to prepare mesoporous CrN by the ammonolysis of a bulk K$_2$Cr$_2$O$_7$, and synthesized Pt/CrN electrocatalyst with mesoporous CrN as the support. The Pt/CrN electrocatalyst exhibited higher corrosion resistance and electrochemical active area (ECSA) than the Pt/C electrocatalysts, as well as higher ORR activity. This high electrocatalytic activity attributed to the high conductivity and high corrosion resistance of CrN support, as well as the synergy between CrN support and Pt.

Similarly, Zheng et al.[70] also made a study about TMN serving as the support of precious metal, and synthesized 3D layered porous Pt/TiN electrocatalyst by a simple approach. The porous 3D support with higher corrosion resistance and mechanical stability played a key role in improving the conductivity and chemical durability of Pt. Therefore, Pt/TiN showed better electrochemical performance than Pt/C with a 40 mV negative shift in the half-wave potential ($E_{1/2}$). In another report, Yue et al.[71] comparatively studied the electrocatalytic activity and durability of Pt/TiN and Pt/C, and found that Pt/TiN electrocatalyst reduced about 22% of the initial ECSA, which was superior to 66% ECSA of Pt/C after 1000 cycles. At the same time, the $E_{1/2}$ of Pt/TiN (0.85 V) was also higher than that of Pt/C (0.846 V). This brilliant performance was attributed to the fact that the energy levels between Pt-d and N-p states were similar, which facilitated the electron transfer to form Pt-N-Ti bonds.

Besides, TMN is also used as electrocatalyst, Qi et al.[72] synthesized MoN/C and Mo$_2$N/C electrocatalysts and studied the relationship between their structure and ORR activity. This work revealed that the energy gap between the HOMO of MoN/C and the LUMO of O$_2$ was excessively small which brought fast electron transfer between them. As a result, MoN/C exhibited higher ORR electrocatalytic activity compared with Mo$_2$N/C in O$_2$-saturated 0.5 M HClO$_4$.

In addition to the MoN electrocatalyst, Zhang et al.[73] reported a series of novel heterostructure Ni$_3$N quantum dots, which were uniformly distributed on the surface of NiO nanosheets. According to the electronic structure characterizations, Ni$_3$N QD showed a strong synergy with NiO nanosheets forming a unique electronic structure around the Fermi level, which not only increased the carrier concentration and conductivity, but also enriched the active sites of oxygen. Thus, the ORR activity of the favorable Ni$_3$N QD/NiO electrocatalyst was comparable to that of Pt/C.

3.1.2 Multivariate TMNs

Although the unary TMN displays impressive electrocatalytic activities, to further enhance the ORR activity, many scholars have conducted on TMN doping. Luo et al.[74] explored the limitations on early-transition-metal nitrides (ScN, TiN, CrN, and VN) as competitive electrocatalysts. Then, this work proposed a strategy to enhance its ORR activity by doping with transition metals and synthesized V$_{0.95}$Mo$_{0.05}$Ns (M=Cu, Ni, Co, Fe, Mn, Cr, and Ti). Among them, V$_{0.95}$Co$_{0.05}$N exhibited the most outstanding ORR performance, with its electron transfer number of the ORR of above 3.8, and the value was competitive with the performance of 20 wt.% Pt/C in alkaline medium.

Tian et al.[75] investigated the effects of the doping elements (Fe, Co, and Ni) on TiN electrocatalytic performance. The experimental data indicated that the doping might have...
impact on the microstructure of the TiN with the vacancy of the d-band of the Ti atom reduction, and its ability to provide electrons to oxygen is increased. Most importantly, the performance of Ti_{0.95}Ni_{0.05}N was almost comparable to that of commercial Pt/C in 0.1 M KOH solution. Later, Tian et al.[76] continuously studied the effects of the doping cobalt on TiN in electrocatalytic performance. They prepared Ti_{0.8}Co_{0.2}N nanosheets by the solvothermal with post-nitriding approach and found the electronic modification effect caused by Co doping endowed the Ti_{0.8}Co_{0.2}N electrocatalyst with a remarkable activity with E_{1/2} of 0.85 V (Figure. 4a-f). The obtained electrocatalyst revealed high durability with only 4mV shifting of the half-wave potential after the durability test for 5000 cycles (Figure. 4g).

Different from Tian and co-workers who prepared TMNs electrocatalysts with Co doping on TiN, Li et al.[77] synthesized Ti_{0.95}Fe_{0.05}N support with large specific surface area and high conductivity with hydrothermal method followed by post-nitriding treatment. According to the electrochemical data, Pt/Ti_{0.95}Fe_{0.05}N displayed higher activity and stability than that of Pt/C. In other words, it was obvious that the ECSA of Pt/C and Pt/Ti_{0.95}Fe_{0.05}N NTs was decreased by 29% and 7% after 1000 cycles and 5000 cycles, respectively, and it diminished 90% and 30%. Furthermore, introducing Fe into TiN NTs enhanced the synergy between metals, which resulted in an enhanced activity of the Pt/Ti_{0.95}Fe_{0.05}N NTs electrocatalyst.

In addition, Varga et al.[78] prepared the nitrogen-doped graphene electrocatalyst decorated with FeN and Fe_{2}N using the freeze-dried mixture of graphene oxide and iron salt as raw material. According to the report, the electrocatalyst with the iron content of

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**Figure 4.** (a) Schematic of the formation of titanium nitrides assemblies; (b) SEM image of Ti_{0.8}Co_{0.2}N assemblies; (c-d) TEM image of Ti_{0.8}Co_{0.2}N assemblies; (e) XRD pattern of TiO\textsubscript{2}, TiN and Ti_{0.8}Co_{0.2}N assemblies; (f) LSV curves of different electrocatalysts in 0.1 M KOH electrolyte; (g) potential cycling performance of Ti_{0.8}Co_{0.2}N assemblies. Redrawn with permission of ref.76. Copyright 2018 American Chemical Society.
20 wt.% achieved the highest current density (3.65 mA cm\(^{-2}\)), which was better than that of the pristine iron nitride and the undecorated nitrogen-doped graphene. Moreover, the electrocatalysts showed improved methanol tolerance compared with the Pt-based electrocatalyst.

In general, it can be seen from these results that TMNs are a promising to improve the ORR electrocatalyst, especially in an alkaline environment. However, the poor performance of TMNs in acidic medium is also an urgent problem to be solved.

### 3.2 Transition metal nitrides as OER electrocatalyst

There is no doubt that the noble metal (Ir and Ru) based materials perform excellently in OER. However, the relative scarcity and high cost make their application in this field unsustainable. Therefore, it is important to search for new OER electrocatalytic materials with lower overpotential, superior stability, and lower cost. In recent years, TMNs have attracted a great deal of attention based on their electronic structure similar to that of the precious metals.[22, 23, 79, 80]

#### 3.2.1 Unary TMN

The Co-based nitride electrocatalyst exhibits strong electron donating ability and high durability due to the change of the M-N bond as well as state density of the metal d-band. Chen et al.[81] prepared Co\(_4\)N porous nanowire based on carbon cloth, and the electrocatalyst showed superior activity for OER with 10 mA cm\(^{-2}\) at an overpotential of 257 mV, which benefited from the synergy effect of metallic property, 1D porous nanowire arrays, and unique 3D electrode configuration. Later, Zhang et al.[82] proposed a new method to prepare cobalt nitride nanowires through N\(_2\) radio-frequency plasma treatment. In this way, it took only 1 min to prepare cobalt nitride.

In addition to cobalt-based TMNs electrocatalysts, nickel-based and iron-based TMNs electrocatalysts are also used in OER. Ni\(_3\)N nanosheets as OER electrocatalyst was first studied by Xu and co-workers.[83]. The Ni\(_3\)N nanosheets performed well in OER with an overpotential of 350 mV and Tafel slope of 45 mV/dec for abundant active sites as well as favorable conductivity. Surface plasmon resonance (SPR) was a recognized mechanism to enhance the activity of OER, Zeng et al.[84] developed a cation exchange strategy to prepare Fe\(_2\)N nanoparticles embedded in mesoporous TiO\(_x\)N\(_y\) nanoshells which could be used as a plasmonic material to induce hot electrons and possess high porosity and electrical conductivity. The Fe\(_2\)N nanoparticles served as an efficient OER electrocatalyst in alkaline medium with an extremely small overpotential of 270 mV at the current density of 10 mA cm\(^{-2}\).

#### 3.2.2 Multivariate TMNs

To improve the OER activity of TMNs, many scholars try to introduce heteroatoms into unary TMNs to change the electronic structure, optimize the coordination of the active metal center environment, and improve the intermediate adsorption as well as desorption on the interface.
Wang et al. [85] grew nickel-cobalt nitride nanosheets on macroporous Ni foam (NF) by electrodeposition. The obtained NiCo-N-NF electrocatalyst achieved good stability and high activity for OER with overpotential of 290 mV at 10 mA cm$^{-2}$ due to its 3D interconnected porous structure and the synergistic effect of bimetallic active sites. Gao et al. [86] also studied the synergistic effect of nickel-cobalt and a surface nitridation strategy. They synthesized nickel-cobalt nitride (Ni$_2$Co-N) nanocactoids on carbon cloth that exhibited a remarkable activity with an overpotential of 214 mV and Tafel slope of 53 mV dec$^{-1}$ in alkaline media lower than those of IrO$_2$. (Figure 5a-g). Li et al. [87] offered a facile approach to synthesize holey cobalt-iron nitride nanosheet based on Ni foam substrate. The obtained CoFeN-HNAs/NF served as attractive OER electrocatalyst with a large ECSA, low charge transfer resistance, and rapid gas diffusion. Besides, the electrocatalyst showed a low overpotential of 260 mV. Liu et al. [88] studied the effect of molar ratio of iron atom and cobalt atom on Fe-CoN electrocatalyst. According to this report, CoFe(3:1)-N showed a remarkable activity for OER with an overpotential of 200 mV and excellent durability. Sun et al. [89] took advantage of the synergistic effect of Co-N-CeO$_2$ to prepare Co$_3$N-CeO$_2$ hybrid nanosheet array grown on a graphite plate (Co$_3$N-CeO$_2$/GP) by anion intercalation enhanced electrodeposition route and subsequent nitridation (Figure. 6a-c). The treatment of CeO$_2$ coupling Co$_3$N porous nanosheet significantly enhanced the OER activity, which was evidenced by a low overpotential of 239 mV to reach a current density of 10 mA cm$^{-2}$ and long-term durability at a large current density of 500 mA cm$^{-2}$ for 50 h (Figure. 6d, e).
For nickel-iron bimetallic TMNs electrocatalyst, Kwag et al. [90] synthesized two-dimensional and nanocrystalline FeNi3N/rGO nanohybrid sheets. The electrocatalyst showed superior activity with overpotential of 290 mV at 10 mA cm\(^{-2}\) and robust long-term stability for 24 h, due to the synergistic effect between nanocrystalline FeNi3N and graphene nanosheets as well as its unique nanoarchitectures. Liang et al. [91] prepared FeNi3N-NiS\(_2\) electrocatalyst by hydrothermal and nitridation process (Figure.7a,b), and the obtained electrocatalyst reveals an extremely brilliant OER properties and long-term stability with a overpotential of 230 mV and a small Tafel slope of 38 mV dec\(^{-1}\) (Figure.7c-e). The high ECSA and the electron effect between FeNi3N and NiS\(_2\) were also explained, which reduced the activation energy of OER, and thus enhanced the intrinsic activity of the electrocatalyst. To further prove that the Fe doping modified the electronic structure of NiS\(_2\), and improved the redox activity of the electrocatalyst surface, thereby significantly improving the electrocatalytic OER performance. Li et al. [92] prepared transition metal nitride, NiFeN, which grew on a 3D network-like support of the strutted graphene foam (NiFeN/SG). The electrocatalyst showed enhanced OER activity, as evidenced by a
low overpotential of 226 mV at 10 mA cm\(^{-2}\). It also indicated that the 3D structure increased the ECSA and promoted active site dispersion, which helped the diffusion of the reactants and accelerated the electrocatalytic OER process. Similarly, Wang et al.[93] compounded a NiFeOOH/NiFeN/Ni heterojunction and stressed the importance of heterojunction in improving OER activity. They found the as-obtained electrocatalyst showed attractive performance for OER, and NiFeN/Ni was largely beneficial for charge transfer, such electrocatalyst only required overpotential of 200 mV to actuate 10 mA cm\(^{-2}\), which was superior to most recently reported electrocatalysts and commercial RuO\(_2\).

Jia et al.[94] synthesized Ni-doped molybdenum nitride nanorods, which not only performed a remarkable small overpotential of 218 mV but also maintained long-term stability for 110 hours. It also indicated that TMNs in combination with multiple active components could produce heterostructure that enhanced the electrocatalyst activity.
Table 1 Summary of TMNs activity as OER electrocatalysts.

| Electro catalysts | Electrolyte | $E_{\text{onset}}$ [V vs. RHE] | Tafel slope [mV dec$^{-1}$] | Ref. |
|-------------------|-------------|-------------------------------|--------------------------|------|
| Co$_4$N          | 1 M KOH    | 1.49                          | 44                       | [81] |
| CoN               | 1 M KOH    | 1.52                          | 70                       | [82] |
| Ni$_3$N          | 1 M KOH    | 1.58                          | 45                       | [83] |
| NiCo$_2$N/NF     | 1 M KOH    | 1.52                          | 65                       | [85] |
| Ni$_3$Co-N       | 1 M KOH    | 1.44                          | 53                       | [86] |
| CoFeN$_x$HNAs/NF | 1 M KOH    | 1.49                          | 57                       | [87] |
| CoFe$_{2\text{3\text{3}}}$-N | 1 M KOH | 1.43                          | 42                       | [88] |
| Co$_3$N-CeO$_2$/GP | 1 M KOH | 1.47                          | 46                       | [89] |
| Fe$_5$Ni$_2$/rGO | 1 M KOH    | 1.52                          | 49                       | [90] |
| FeNi$_3$N-NiS$_2$ | 1 M KOH | 1.46                          | 38                       | [91] |
| Ni$_3$Fe$_2$/SG  | 1 M KOH    | 1.46                          | 43                       | [92] |
| NiFeOOH/Ni$_3$FeN/Ni | 1 M KOH | 1.43                          | 36                       | [43] |
| Nifoam@Ni-N$_{2\text{Ni}}$Mo$_{0.8}$N | 1 M KOH | 1.45                          | 39                       | [93] |

3.3 Transition metal nitrides electrocatalyst for ORR&OER

So far, Ir-based electrocatalysts are considered to be the best OER electrocatalysts. In the meantime, Pt-based electrocatalysts are the best choices for ORR. However, pure Ir-based or Pt-based materials are too expensive, and not sufficiently active for bifunctional OER and ORR. Hence TMNs serving as highly active and long-term stable bifunctional electrocatalysts have been widely investigated.[95, 96]

A novel bifunctional electrocatalyst (Ni$_3$FeN/NRGO) was prepared by Fan et al.[97], which substituted for the commercial noble-metal electrocatalysts. The NRGO not only dispersed the Ni$_3$FeN nanoplates, but also provided a conductive framework for the high retention of the ECSA and improvement of the synergistic effect in the OER and ORR processes. By coupling the theoretical and experimental approaches, Ni$_3$FeN/NRGO was confirmed to show high performance in both OER and ORR with the lowest onset overpotential of 150 mV for OER and the onset potential of 0.9 V for ORR. In addition, Wang et al.[98] delivered a novel bifunctional Ni$_3$FeN/Co,N-CNF electrocatalyst driving both OER and ORR. Because of Co,N-CNF as the support of the electrocatalyst, the Ni$_3$FeN nanoparticles achieved highly dispersed. At the same time, as a result of the synergistic effect of Ni$_3$FeN/Co and N-CNF, the as-prepared electrocatalyst had better OER and ORR activities than noble metal electrocatalysts with the low overpotential of 0.27 V for OER and $E_{1/2}$ of 0.81 V for ORR. Zhang et al.[99] also reported ultra-small nanoparticles (Fe$_2$N) grown on CNTs. Compared with the pure Fe$_2$N nanoparticles, the Fe$_2$N nanoparticles in Fe$_2$N-CNTs was extremely smaller (5 nm). Thus, The $E_{1/2}$ of Fe$_2$N-CNTs for ORR was 0.71V, and the overpotential for OER was 240mV. This work might bring us a novel, environmentally friendly approach for the preparation of ultra-small TMNs. To further improve the stability of nickel-iron nitride electrocatalyst, Fu et al.[100] prepared mesoporous nickel-iron nitride bifunctional electrocatalyst without carbon as support (Figure.8a). The NiFeN electrocatalyst was microsphere with hierarchically porous structure, which provided generous interparticle void space and high specific surface area (Figure.8b-g). In this manner, it showed excellent ORR activity with $E_{1/2}$ of 0.78 V and OER activity with a low overpotential of 355 mV at 10 mA cm$^{-2}$ in alkaline solution (Figure.8h,i). A trimetallic (NiFeMn) nitride electrocatalyst was produced by Wu et al.[101] in a molecular sheet form, which was stabilized by Ti metal on titanium carbide (Ti$_3$C$_2$) sheets. Intimate contact
between the two sheets produced a strong force at the interface, thus effectively avoiding the accumulation of a nitride sheet. This work unlocked a high electrocatalytic performance of bifunctional trimetallic nitride electrocatalyst, and provided a new way for the application of two-dimensional sheets in flexible and wearable devices. Different from the above, Cui et al.[102] prepared FePt electrocatalyst with porous nickel-iron nitride as the support. On the one hand, unlike the widely studied Pt-M disordered alloys, ordered FePt intermetallic nanoalloy had definite composition and structure, which was beneficial for the formation of highly dispersed active sites.[103] On the other hand, NiFeN support showed brilliant chemical stability and high conductivity. Therefore, FePt/NiFeN electrocatalyst released excellent electrocatalytic activity for OER and ORR at the same time, and achieved a long-term cycling performance of over 480 h at 10 mA cm\(^{-2}\) in Zn-air batteries.

![Figure 8](https://example.com/figure8.png)

**Figure. 8** (a) Schematic of the formation of the porous NiFeN hierarchical microspheres; (b-d) SEM images of the NiFe LDH hierarchical microspheres; (e-g) SEM images of the porous NiFeN hierarchical microspheres; (h) ORR LSV curves in 0.1 M KOH electrolyte; (i) OER LSV curves in 0.1 M KOH electrolyte. Redrawn with permission of ref.99. Copyright 2017 Elsevier.
In addition to nickel-iron nitride, Davari et al. [104] produced a mixture of two MnN phases (Mn$_4$N and Mn$_6$N$_{2.58}$) as bifunctional electrocatalysts for rechargeable Zn-air batteries. Although the obtained MnN particles were irregular in shape with the size of 100 µm, the high covalent nature of the Mn-N bond provided a greater driving force for the electrocatalytic reaction [105], thus making the electrocatalyst show superior bifunctional activity. At the same time, it exhibited good ORR and OER stability for at least 1500 cycles in 0.1 M KOH.

Figure. 9 (a) Schematic of the formation of Co$_4$N@NC-m and Co$_4$N@NC; (b) ORR LSV curves in 0.10 M KOH electrolyte; (c) OER LSV curves in 0.1 M KOH electrolyte. Redrawn with permission of ref.105. Copyright 2019 Elsevier.

Chen et al. [106] obtained Co$_4$N@N-doped carbon with the assistance of melamine. Besides, they cleverly added melamine to help increase conductivity, the amount of nitrogen doping, and ECSA of the electrocatalyst, and regulate the size and distribution of Co$_4$N nanocrystals (Figure.9a). Therefore, Co$_4$N@NC-m served as a high activity and long-term stability electrocatalyst for both ORR/OER and air cathode of rechargeable Zn-air batteries. Besides, their work provided an effective approach to synthesize non-noble metal electrocatalysts with controllable morphology (Figure.9b, c). Guan et al. [107] studied the cobalt nitride nanoparticles supported on nitrogen-doped reduced graphene oxide sheet (O-S-Co$_5$N@N-RGO). Such flexible material showed marvelous electrocatalytic activity towards both OER and ORR with low overpotentials 380 mV at 10 mA cm$^{-2}$ current density and $E_{1/2}$ of 0.82 V in 1 M KOH solution. It was believed that the Co-N$_4$ sites in the RGO sheet and the Co sites on the surface of O-S-Co$_5$N crystal were the active sites for ORR and OER, respectively. Gao et al. [108] developed cobalt-iron bimetallic nitrides with
N-doped multi-walled carbon nanotubes (Co-Fe-N@MWCNT). It was worth noting that MWCNTs acted as a bridge to connect nanoparticles to optimize the conductivity of the electrocatalyst, while regulating the uniform dispersion of nanoparticles to increase ECSA.[109] In addition, there was a strong interaction between iron and cobalt, and iron could effectively regulate the conversion of cobalt from cobalt (III) to cobalt (IV).[110] Thus, Co-Fe-N@MWCNT performed well for both OER and ORR.

Table 2 Summary of TMNs activity as OER&ORR electrocatalysts (the voltage gap: $\Delta E = E_{i=10} - E_{1/2}$)

| Electrocatalysts | Electrolyte | $E_{1/2}$ for ORR [V vs. RHE] | $E_{i=10}$ for OER [V vs. RHE] | $\Delta E$ [V] | Ref. |
|-----------------|-------------|-------------------------------|-------------------------------|---------------|-----|
| Ni₃FeN/NRGO     | 0.1 M KOH   | 0.72                          | 1.38                          | 0.77          | [96]|
| Ni₃FeN/Co₃N-CN  | 0.1 M KOH   | 0.81                          | 1.50                          | 0.69          | [97]|
| Fe₃N/N-CNTs     | 0.1 M KOH   | 0.71                          | 1.66                          | 0.95          | [98]|
| NiFeN           | 0.1 M KOH   | 0.78                          | 1.58                          | 0.7           | [99]|
| NiFeMnN         | 0.1 M KOH   | 0.84                          | 1.52                          | 0.68          | [100]|
| FePt/Ni₃FeN     | 0.1 M KOH   | 0.93                          | 1.60                          | 0.72          | [101]|
| Co₄N@NC-m       | 0.1 M KOH   | 0.87                          | 1.63                          | 0.81          | [105]|
| Co₆.47N         | 0.1 M KOH   | 0.82                          | 1.61                          | 0.80          | [106]|
| Co-Fe-N@MWCNT   | 0.1 M KOH   | 0.92                          | 1.52                          | 0.72          | [107]|

3.4 Design principles of TMNs electrocatalysts for oriented applications

TMNs play a key role in electrocatalytic application especially in OER and ORR. The oxygen-involved reactions require extremely abundant accessible electrocatalytic sites due to its slow kinetics. Thus the reasonably designing TMNs is an effective approach to improve the performance for electrocatalytic application.

In the aspect of doping agents, the electrocatalyst is desired to adjust the electronic structure to improve the ability of donating electrons to adsorbed oxygen. Furthermore, the doping agents show great influence on the microstructures of electrocatalysts. With the defects producing, active centers are formed, which is beneficial to capture electrons and speed up mass electron transport.

On the other hand, the electrocatalyst is expected to possess a large number of exposed active sites since the electrocatalytic reaction takes place on the electrochemical surface of the electrocatalyst. Hence nanostructure with high specific surface area such as nanoflowers, hierarchically macropores and mesopores is required to take full advantage of exposed active sites. The strong synergy between the intrinsic activity and nanostructures is also demanded to achieve the improvement of electrocatalytic performance.

4. Conclusions and perspectives

In this review, we summarized the recent advances in the synthesis and application of transition metal nitrides for oxygen reduction reaction and oxygen evolution reaction. Selecting different transition metal ions (Ti, Ni, Fe, Co, Mo) or introducing heteroatoms into the unary transition metal could make the TMNs produce similar properties to noble metal. Similarly, high-efficiency multiscale mass transfer structure was constructed by using TMNs nanowires and mesoporous structure, which had been proved to greatly enhance the electrocatalytic performance of TMNs. To improve the preparation and application of TMNs, the following challenges need to be studied later.

Firstly, the preparation of TMNs electrocatalysts has been restricted to laboratory preparation, which cannot be produced on a large scale due to its complicated preparation
and high cost. Therefore, simple, low-cost, and environmentally friendly preparation methods should be investigated as a part of the subsequent research.

Secondly, in acidic media or under the conditions with extreme pH value, the activity of TMNs electrocatalysts will severely reduce due to the dissolution of transition metals, protonation of active sites, and corrosion of conductive substrates. Therefore, TMNs electrocatalysts with robust stability need to be developed.

Thirdly, to further improve the electrocatalytic activity, the electronic structure can be adjusted based on the d-band center theory by doping with metal or non-metal elements, forming a heterostructure, or using a synergistic effect.

Finally, the in-situ techniques like XRD/ND-analysis, TEM, and XPS are used to monitor the changes of TMNs during the preparation process and the electrocatalytic reaction process, which is beneficial for the understanding the principle of TMNs catalysis and provides a theoretical basis for subsequent electrocatalyst research.

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