The effect of \textit{in situ} nitrogen doping on the oxygen evolution reaction of MXenes†

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The development of non-noble metal electrocatalysts with high performance for the oxygen evolution reaction (OER) is highly desirable but still faces many challenges. Herein, we report a facile and controllable strategy to fabricate N-doped titanium carbide flakes (Ti$_3$C$_{1.8}$N$_{0.2}$ and Ti$_3$C$_{1.6}$N$_{0.4}$) using an \textit{in situ} nitrogen solid solution, followed by an etching process. The introduction of nitrogen is beneficial to the Ti$_3$C$_{1.6}$N$_{0.4}$ flakes for more exposed active sites, accelerated charge transfer upon an electrochemical reaction, and improved wettability for more accessible sites. As a result, the as-obtained Ti$_3$C$_{1.6}$N$_{0.4}$ catalyst exhibits enhanced electrocatalytic properties for OER, which include a small $\eta_{\text{ onset }}$ of 245.8 mV, low Tafel slope of 216.4 mV dec$^{-1}$, and relatively good catalytic stability. The present work not only deepens the understanding of \textit{in situ} N-doped MXene electrocatalysts, but also provides a guideline for the preparation of other N-doped MXene-based hybrid materials for other renewable energy applications.

1. Introduction

Due to the increasing energy consumption and serious pollution, we are eager to develop sustainable energy sources to lessen the reliance on fossil fuels. Renewable energy storage and conversion techniques such as fuel cells, metal-air batteries, and water splitting systems, which have high efficiency, excellent reliability, and zero pollution, are promising alternatives for traditional fossil fuels.\textsuperscript{1-4} The oxygen evolution reaction (OER) is one of the most fundamental and important reactions in the above-mentioned renewable energy technologies. However, this reaction is currently constrained by sluggish kinetics and large overpotential; thus, efficient electrocatalysts are in demand. To date, precious metal-based catalysts, such as IrO$_2$ and RuO$_2$, are the best due to their optimum binding with essential intermediates, but their applicability is hampered due to their high cost and scarcity.\textsuperscript{5} Hence, tremendous efforts have been devoted to search for alternative electrocatalysts constructed from low-cost and earth-abundant elements to efficiently conduct OER with a minimum overpotential.\textsuperscript{6}

MXenes, as an emerging family of two-dimensional (2D) transition metal carbide, nitride and carbonitride nanomaterials, have attracted significant attention. Their formula is M$_{m+1}$X$_n$T$_z$, where M is an early transition metal, X represents C and/or N, and T represents surface termination groups (–O, –OH, and –F).\textsuperscript{7-9} Moreover, MXenes can be achieved by the selective etching of the A element from the MAX phases, where A is a group IIIA or IVA element. For the sake of simplicity, MXenes are usually expressed as M$_{m+1}$X$_n$. MXenes are regarded as promising supporting materials by virtue of their high electrical conductivity, unique hydrophilicity, relatively good chemical stability, and high surface area as well as their ultralow work function.\textsuperscript{7} In particular, in the field of electrocatalysis, MXenes ensure fast electron transport during electrochemical reactions and may alter the electrophilicity of the active centers of the catalysts, thus tuning the catalytic properties of multicomponent catalyst systems. Ti$_3$C$_2$, as a promising member of MXenes, is commonly used in the field of electrocatalysis. Nevertheless, it should be mentioned here that the inherent catalytic activity of Ti$_3$C$_2$ is relatively low, which may be due to its low number of catalytic active sites and the tendency to stack.

To improve its catalytic activity, one commonly used strategy is to combine Ti$_3$C$_2$ with materials that have high catalytic activity. On the basis of this principle, Ti$_3$C$_2$-based hybrid electrocatalysts, such as Ni$_{1-x}$Fe$_x$PS@Ti$_3$C$_2$,\textsuperscript{8} NiCoS/Ti$_3$C$_2$T$_x$,\textsuperscript{8} FeNi-LDH/Ti$_3$C$_2$-MXene,\textsuperscript{9} MoS$_2$/Ti$_3$C$_2$-MXene@C,\textsuperscript{10} and Ti$_3$C$_2$T$_x$-CoBDC,\textsuperscript{7} appear constantly and exhibit superb OER/HER activity. Additionally, the chemical doping of heteroatoms is an effective strategy to improve the electrochemical properties of materials. Notably, nitrogen-containing functional groups can significantly enhance the electrochemical performance of the materials by increasing the electrical conductivity for faster ion transport and improvement in surface wettability for more active sites.\textsuperscript{12,13} For example, our previous studies have...
demonstrated that nitrogen-doped Ti$_3$C$_2$ nanosheets possess superior electrochemical performance.\textsuperscript{14-16} Jiang and coworkers have demonstrated that the synthesized nitrogen-doped ultrathin carbon nanosheets (NCNs) with an ultrathin sheet structure, ultrahigh specific surface area, and rich edge defects exhibit extraordinary trifunctional activity toward ORR, OER, and HER.\textsuperscript{17} Sheng et al. have shown that the developed self-supported nitrogen-doped carbon hydrogel films also have outstanding OER activity.\textsuperscript{18} Moreover, dual-atom (N and P) co-functionalized porous carbon materials possess superior electrolytic activity.\textsuperscript{19-20}

In this work, nitrogen-doped few-layered Ti$_3$C$_2$ nanosheets with different nitrogen concentrations (Ti$_3$C$_{1.8}$N$_{0.2}$ and Ti$_3$C$_{1.6}$N$_{0.4}$) have been synthesized using an \textit{in situ} nitrogen solid solution, followed by an HCl and LiF-assisted etching process. The results prove that the introduction of nitrogen is beneficial for electrical conductivity and wettability. Besides, the fresh Ti$_3$C$_{1.6}$N$_{0.4}$ electrocatalysts with relatively higher electrical conductivity, better wettability, and more active sites are favorable for OER activity. This work expands the understanding of the effect of \textit{in situ} nitrogen doping on the catalytic performance of MXenes and also contributes to the preparation of N-doped MXene hybrid structures with higher catalytic activity.

2. Results and discussion

A previous work has indicated that nitrogen can be easily introduced into the Ti$_3$C$_2$ flakes by achieving the corresponding nitrogen solid solution (Ti$_3$AlC$_{1.8}$N$_{0.2}$ and Ti$_3$AlC$_{1.6}$N$_{0.4}$ powders).\textsuperscript{15} Additionally, Ti$_3$C$_2$, Ti$_3$C$_{1.8}$N$_{0.2}$ and Ti$_3$C$_{1.6}$N$_{0.4}$ flakes are obtained using a wet-chemical method, as presented in Fig. S1,\textsuperscript{†} and the atomic structure of the Ti$_3$C$_{1.6}$N$_{0.4}$ phase is schematically presented in Fig. 1. It can be seen that a certain number of nitrogen atoms dispersed into the Ti$_3$C$_{1.6}$N$_{0.4}$ atom structure not only provide abundant active sites, but also effectively improve the electrical conductivity and wettability, thus enhancing the electrocatalytic activity. All of these advancements endow Ti$_3$C$_{1.6}$N$_{0.4}$ with enhanced electrocatalytic performance for OER in an alkaline solution (4OH$^-$ $\rightarrow$ 2H$_2$O(l) + O$_2$(g) + 4e$^-$).

Actually, the flexible and free-standing films produced by the vacuum filtration of few-layered MXene flake suspensions are beneficial for the characterization of their own features. Scanning electron microscopy (SEM) and transmission electron microscopy (TEM) were employed to observe the morphological and microstructural properties of the Ti$_3$C$_2$, Ti$_3$C$_{1.8}$N$_{0.2}$, and Ti$_3$C$_{1.6}$N$_{0.4}$ films. From the top-view SEM images (Fig. 2a-c), similar morphologies with considerable wrinkles can be found on the surfaces of the Ti$_3$C$_2$, Ti$_3$C$_{1.8}$N$_{0.2}$, and Ti$_3$C$_{1.6}$N$_{0.4}$ films. Fig. 2d shows the cross-sectional view SEM image of the Ti$_3$C$_{1.8}$N$_{0.2}$ film. It can be clearly seen that the flexible Ti$_3$C$_{1.8}$N$_{0.2}$ film with a well-aligned layered structure self-assembles from the Ti$_3$C$_{1.6}$N$_{0.4}$ flakes via the vacuum filtration method, which is consistent with previous results.\textsuperscript{6,21} Further observation of the Ti$_3$C$_{1.6}$N$_{0.4}$ flakes shows that they have a lateral size of several microns and thickness of several nanometers, as shown in Fig. 2e. The corresponding EDS elemental mappings (Fig. 2f) reveal that Ti, C, and especially N are evenly distributed in the Ti$_3$C$_{1.8}$N$_{0.2}$ film, which is also good evidence for the successful introduction of nitrogen. X-ray diffraction (XRD) was employed to examine the crystal structures of the three ceramic powders and films (Fig. S2 and 2g). It can be seen that the XRD patterns of the three ceramic powder samples show similar characteristic peaks, which can be assigned to the diffraction peaks of Ti$_3$AlC$_2$ (PDF card #52-0875). Moreover, with the increase in the nitrogen doping content, the typical peaks of (104) gradually shift to the right, indicating the successful introduction of nitrogen (Fig. S2†). The XRD patterns of the three films are similar, indicating that the three films have similar crystal structures. However, the inset shows that the characteristic peaks of the (002) plane of the Ti$_3$C$_{1.8}$N$_{0.2}$ and Ti$_3$C$_{1.6}$N$_{0.4}$ films shift to a lower angle, which confirms the enhancement in the d-spacing caused by nitrogen doping. Besides, the shift in the typical (002) peak of the Ti$_3$C$_{1.6}$N$_{0.4}$ film is relatively inconspicuous compared with that of the Ti$_3$C$_{1.8}$N$_{0.2}$ film, which may be due to different types of nitrogen. The above-mentioned results demonstrate that nitrogen has been successfully introduced into the Ti$_3$C$_{1.8}$N$_{0.4}$ flakes.

To detect the chemical composition and bonding states of the as-synthesized samples, X-ray photoelectron spectroscopy (XPS) was subsequently performed for these samples. The survey region (0–1200 eV) shown in Fig. 3a indicates the presence of the common peaks of C 1s, N 1s, O 1s, F 1s, and Ti 2p. Specifically, additional N 1s peaks for the Ti$_3$C$_{1.8}$N$_{0.2}$ and Ti$_3$C$_{1.6}$N$_{0.4}$ films further indicate the successful introduction of nitrogen into these films, which is in good agreement with the EDS results. The elemental composition \textit{via} XPS presented in Table S1† reveals that the concentrations of N 1s are 3.76 at% and 5.72 at% for Ti$_3$C$_{1.8}$N$_{0.2}$ and Ti$_3$C$_{1.6}$N$_{0.4}$, respectively. More nitrogen atoms improve the electronegativity of Ti$_3$C$_{1.8}$N$_{0.4}$, which is beneficial for the electron-withdrawing effect. To further analyze the location of nitrogen, the deconvolutions of the N 1s region spectra are shown in Fig. 3b. The three peaks located at 397.1 eV, 399.8 eV, and 401.9 eV correspond to the N–Ti (nitride), N–5 (pyrrolic nitrogen), and N–Q (quaternary nitrogen) species.\textsuperscript{21} The presence of the N–Ti peak provides confirmation of the \textit{in situ} substitution of C atoms with N atoms in the octahedral interstice of the Ti$_3$C$_{1.8}$N$_{0.2}$ structure. The N–Ti bond in the N 1s spectra is consistent with the Ti–N peak centered at 457.3 eV in the Ti 2p spectra (Fig. S3†). In particular, the Ti$_3$C$_{1.8}$N$_{0.2}$ structure exhibits more N species, including of N–Ti, N–5, N–Q species, and an extra N–6 (pyridinic nitrogen) bond at 398.5 eV.\textsuperscript{22} In addition, the concentrations of the different nitrogen species in the two samples are quantified based on the integrated peak areas and presented in Table S2.† It can be seen that N–Ti is the dominant N species in the Ti$_3$C$_{1.8}$N$_{0.2}$ (52.46 at%) and Ti$_3$C$_{1.6}$N$_{0.4}$ (61.75 at%) samples; in contrast, compared with that of the Ti$_3$C$_{1.8}$N$_{0.2}$ sample, the loading of N–5 in the Ti$_3$C$_{1.6}$N$_{0.4}$ sample diminishes markedly accompanied by an increase in N–Q and the emergence of N–6.
Based on the above-mentioned results and previously reported findings,\textsuperscript{19,23–27} the properties of the various N bonds are as follows: (1) the N–Ti and N-5 species have little effect on the improvement in the electrocatalytic performance. (2) N–Q, where N is bonded with three C neighbors, provides a larger contribution to enhance the electrocatalytic activity. (3) N-6, where N is incorporated into a six-membered heterocyclic ring, brings about electron donor properties, which may be the main reason for the enhanced OER activity. Therefore, higher contents of both N–Q (13.52 at%) and N-6 (2.88 at%) in the Ti\textsubscript{3}C\textsubscript{1.6}N\textsubscript{0.4} sample endow it with higher catalytic activity.

Fig. 3c shows the Raman spectra of the three films and the peaks are consistent with those reported in our previous works.\textsuperscript{6,15} In particular, the characteristic peak at \( \sim 1364 \text{ cm}^{-1} \) is related to the D-band and is attributed to the disordered graphite formed by the defects in carbon-based materials. The other characteristic peak located at \( \sim 1549 \text{ cm}^{-1} \) is related to the G-band, which corresponds to the vibration of the sp\textsuperscript{2}-hybridized carbon atoms in a 2D hexagonal lattice. The values of \( I_D/I_G \) calculated from the intensities of the D-band and G-band peaks were 0.89 for Ti\textsubscript{3}C\textsubscript{2}, 0.98 for Ti\textsubscript{3}C\textsubscript{1.8}N\textsubscript{0.2}, and 1.06 for Ti\textsubscript{3}C\textsubscript{1.6}N\textsubscript{0.4}. A higher value of \( I_D/I_G \) implies the insertion of N atoms and indicates the formation of more disorder and defects (heterogeneous atoms and vacancies) in the carbon structure, which contribute to additional active sites for the improvement in catalytic activity.\textsuperscript{28} In addition, the distinct redshift in the D peak of the Ti\textsubscript{3}C\textsubscript{1.6}N\textsubscript{0.4} sample indicates the restoration of the conjugate structure after the nitrogen atoms are doped into the graphitic lattice. As indexed in the Raman spectra, the rich defect sites observed among the crystal lattices of Ti\textsubscript{3}C\textsubscript{1.6}N\textsubscript{0.4} can tune its electronic and surface properties and thus optimize the adsorption energies of the electrochemical catalysis steps.

It has been reported that the structural defects caused by heterogeneous atom doping in materials have a significant influence on their electrical properties.\textsuperscript{29,30} The electrical conductivities of the films are shown in Fig. 3d and S5.\textsuperscript{†} As expected, doping nitrogen atoms is indeed helpful to improve the conductivity of the sample. The electrical conductivities were 81.74 \( \pm \) 1.16 mS cm\textsuperscript{-1} for Ti\textsubscript{3}C\textsubscript{1.6}N\textsubscript{0.4}, 28.42 \( \pm \) 0.43 mS cm\textsuperscript{-1} for Ti\textsubscript{3}C\textsubscript{1.8}N\textsubscript{0.2}, and 21.90 \( \pm \) 0.39 mS cm\textsuperscript{-1} for Ti\textsubscript{3}C\textsubscript{2}. It is worth mentioning that Ti\textsubscript{3}C\textsubscript{1.6}N\textsubscript{0.4} has the highest electrical conductivity among the three measured films, which can result in efficient electron transfer during the electrocatalytic reaction process. In addition, the contact angle of a dynamic 1 M KOH aqueous electrolyte was measured for the three films. As illustrated in Fig. 3e, the introduction of nitrogen has a positive effect on wettability. Obviously, the Ti\textsubscript{3}C\textsubscript{1.6}N\textsubscript{0.4} film shows a smaller contact angle of 9.8° with the electrolyte, which indicates that Ti\textsubscript{3}C\textsubscript{1.6}N\textsubscript{0.4} is more hydrophilic compared with Ti\textsubscript{3}C\textsubscript{1.8}N\textsubscript{0.2} and Ti\textsubscript{3}C\textsubscript{2}. Better hydrophilicity can improve the accessibility of the catalyst to the electrolyte and increase the available sites, thus leading to enhanced catalytic performance.

To obtain further insights in the effect of nitrogen doping on electrocatalytic performance, we studied the fresh Ti\textsubscript{3}C\textsubscript{2},
Ti$_3$C$_{1.8}$N$_{0.2}$ flakes towards OER in a 1 M KOH alkaline solution in a typical three-electrode system, where the potential was compensated by solution resistance ($iR_s$). Before the electrochemical measurements, the electrode underwent an activation treatment using several linear sweep voltammetry (LSV) until a stable curve was achieved. Fig. 4a presents the LSV curves of the Ti$_3$C$_2$, Ti$_3$C$_{1.8}$N$_{0.2}$, and Ti$_3$C$_{1.6}$N$_{0.4}$ electrodes at a scan rate of 5 mV s$^{-1}$. The bare carbon fiber paper (CFP) substrate showed no catalytic activity. In contrast, the fresh Ti$_3$C$_2$ flakes showed minimal catalytic activity, while the fresh Ti$_3$C$_{1.8}$N$_{0.2}$ flakes exhibited enhanced catalytic activity. Apparently, the fresh Ti$_3$C$_{1.6}$N$_{0.4}$ flakes showed the best OER activity, which may be because of the synergistic effect of more exposed active sites, higher electrical conductivity, and better wettability due to the introduction of nitrogen. However, it is also worth mentioning here that N doping improves activity, but it can result in reduced stability of the material structure at the same time, as reflected in the literature. The corresponding Tafel plots are also provided to investigate the OER kinetics of the electrocatalysts. As shown in Fig. 4b, the Tafel slopes are 243.4 mV dec$^{-1}$, 234.7 mV dec$^{-1}$, and 216.4 mV dec$^{-1}$ for the fresh Ti$_3$C$_2$, Ti$_3$C$_{1.8}$N$_{0.2}$, and Ti$_3$C$_{1.6}$N$_{0.4}$ catalysts, respectively. Obviously, the Ti$_3$C$_{1.6}$N$_{0.4}$ flakes exhibited the smallest Tafel slope, indicating the best OER catalytic kinetics.

The overpotential is an important index to evaluate the electrocatalytic performance of the as-fabricated electrocatalysts. The onset overpotential is defined as the overpotential when the current density reaches 1.0 mA cm$^{-2}$. As shown in Fig. 4c, the onset overpotentials ($\eta_{\text{onset}}$) are 449.0, 418.7, and 245.8 mV for the fresh Ti$_3$C$_2$, Ti$_3$C$_{1.8}$N$_{0.2}$, and Ti$_3$C$_{1.6}$N$_{0.4}$ electrocatalysts, respectively. The smaller $\eta_{\text{onset}}$ of the Ti$_3$C$_{1.6}$N$_{0.4}$ electrocatalyst implies its superior electrocatalytic activity, which means that a low additional potential is required to sustainably drive an electrochemical reaction from its reversible potential. In addition, the mass activity, which is the current normalized by the loaded mass of a catalyst, is currently gaining importance for the evaluation of nanostructured electrocatalysts for HER. The mass activity of the Ti$_3$C$_{1.8}$N$_{0.2}$ electrocatalyst at $\eta = 350$ mV was 14.0 A g$^{-1}$, which was 7.7-fold and 5.6-fold higher than those of the Ti$_3$C$_2$ and Ti$_3$C$_{1.8}$N$_{0.2}$ electrocatalysts, respectively. The much higher mass activity (A g$^{-1}$) at the same overpotential ($\eta$) further demonstrates that Ti$_3$C$_{1.6}$N$_{0.4}$...
is the best among the three electrocatalysts. Besides indispensable higher activity, the excellent stability of electrocatalysts toward OER is another critical factor for their practical application. The long-term durability of the electrocatalyst was estimated via chronopotentiometry at 10 mA cm$^{-2}$. It can be observed from Fig. 4d that the Ti$_3$C$_{1.6}$N$_{0.4}$ electrode shows a fluctuating potential curve over a period of 12 h. The Ti$_3$C$_{1.6}$N$_{0.4}$ electrode requires $\sim$1.68 V to afford an OER current density of 10 mA cm$^{-2}$. The initial high potential is due to incomplete electrode activation and partial wettability of the electrode surface. However, it is worth mentioning that the continuous increase in potential after 9.0 h is due to the reduced stability of the Ti$_3$C$_{1.6}$N$_{0.4}$ flakes, which might be due to the combined effects of the prolonged alkaline environment, released oxygen and electricity.

To gain a better understanding of the superior electrocatalytic activity of the Ti$_3$C$_{1.6}$N$_{0.4}$ catalyst, the electrochemically active surface area (ECSA) of the catalyst was estimated by determining the electrochemical double layer capacitance ($C_{dl}$) since ECSA is generally proportional to $C_{dl}$.$^{22}$ A small potential

Fig. 3  (a) XPS spectra of the Ti$_3$C$_2$, Ti$_3$C$_{1.8}$N$_{0.2}$, and Ti$_3$C$_{1.6}$N$_{0.4}$ films. (b) High-resolution XPS N 1s spectra of the Ti$_3$C$_{2}$N$_{0.2}$ and Ti$_3$C$_{1.6}$N$_{0.4}$ films. Raman spectra (c) and electrical conductivities (d) of the three films. (e) The contact angles of the dynamic 1 M KOH electrolyte for the three films.
range from 1.17 V to 1.27 V without an apparent faradaic reaction was selected to calculate \( C_{dl} \) (Fig. S4†). Thus, the final \( C_{dl} \) was obtained by extracting the slope of the linear relationship between the current density difference and the scan rate. As shown in Fig. 5a, the \( C_{dl} \) values are 2.51 mF cm\(^{-2}\) for the Ti\(_3\)C\(_2\) catalyst, 3.09 mF cm\(^{-2}\) for Ti\(_3\)C\(_{1.8}\)N\(_{0.2}\), and 6.08 mF cm\(^{-2}\) for Ti\(_3\)C\(_{1.6}\)N\(_{0.4}\). Compared with Ti\(_3\)C\(_2\) and Ti\(_3\)C\(_{1.8}\)N\(_{0.2}\), Ti\(_3\)C\(_{1.6}\)N\(_{0.4}\) has a larger \( C_{dl} \) value, indicating that it has more active sites, which are useful for catalyzing OER.

It is widely accepted that the OER performance in the electrolyte is dominated by ionic and transport resistances.\(^{33}\) As shown in Fig. 5b, to further probe the possible reason for the enhanced performance of the Ti\(_3\)C\(_{1.6}\)N\(_{0.4}\) catalyst, the corresponding Nyquist plots at 1.62 V have been recorded to pinpoint the resistance during experimental OER. The equivalent circuit model is leveraged to obtain the values of the ionic and charge transfer resistances of the OER catalyst in a 1 M KOH electrolyte (inset of Fig. 5b). In a high frequency region, the diameter of the semicircular arc is related to the charge transfer resistance (\( R_{ct} \)), which originates from the ion transfer across the entire interface between the electrode material and the electrolyte accessible area. The real-axis intercept of the impedance spectra in the high frequency region corresponds to the internal resistance (\( R_s \)) of the test device. The simulated \( R_s \), \( R_{ct} \), and CPE values of the Ti\(_3\)C\(_2\), Ti\(_3\)C\(_{1.8}\)N\(_{0.2}\), and Ti\(_3\)C\(_{1.6}\)N\(_{0.4}\) catalysts are shown in Table S3.† Notably, the Ti\(_3\)C\(_{1.6}\)N\(_{0.4}\) catalyst has relatively small \( R_s \) (3.12 ohm) compared to the Ti\(_3\)C\(_2\) (\( R_s = 4.88 \) ohm) and Ti\(_3\)C\(_{1.8}\)N\(_{0.2}\) (\( R_s = 5.39 \) ohm) catalysts, indicating a decrease in the internal resistance and thus a better electronic response. Furthermore, the Ti\(_3\)C\(_{1.6}\)N\(_{0.4}\) catalyst exhibits the lowest \( R_{ct} \) (279.20 ohm), which can reduce the potential barrier to drive the current transport across the catalyst. To this end, the resistances acquired from the Nyquist plots of the three catalysts reveal that the Ti\(_3\)C\(_{1.6}\)N\(_{0.4}\) catalyst with significantly lower...
impedance markedly enhances OER kinetics. In a word, the improved OER activity of the Ti$_3$C$_{1.6}$N$_{0.4}$ catalyst is ascribed to the synergetic effect of high electrical conductivity, good wettability, and more exposed active sites.

3. Conclusions

In summary, we have successfully synthesized N-doped titanium carbide nanosheets (Ti$_3$C$_{1.8}$N$_{0.2}$ and Ti$_3$C$_{1.6}$N$_{0.4}$) using an in situ nitrogen doping solid solution method with the merits of facile and controllable features. The results indicate that the introduction of nitrogen can increase exposed active sites and improve electrical conductivity and wettability, thus enhancing the electrocatalytic activity. As a result, the Ti$_3$C$_{1.6}$N$_{0.4}$ catalyst with a higher nitrogen concentration exhibits better catalytic properties for OER, including a small $\eta_{\text{onset}}$ of 245.8 mV, low Tafel slope of 216.4 mV dec$^{-1}$, and relatively good catalytic stability. The enhanced electrocatalytic activity of the Ti$_3$C$_{1.6}$N$_{0.4}$ catalyst is attributed to the synergetic effects of abundant catalytic active sites, high conductivity, low resistance, and good wettability. This work contributes to the field of in situ nitrogen-doped MXene electrocatalysts and also provides a guideline for the preparation of other N-doped MXene-based hybrid materials for renewable energy applications.

4. Experimental section

Detailed experimental information is available in the ESL.$^\dagger$

Conflicts of interest

There are no conflicts to declare.

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References


Fig. 5 (a) Double-layer capacitances ($C_{dl}$) of the Ti$_3$C$_2$, Ti$_3$C$_{1.8}$N$_{0.2}$, and Ti$_3$C$_{1.6}$N$_{0.4}$ electrocatalysts. (b) Nyquist plots of the above-mentioned electrocatalysts measured at a potential of 1.62 V vs. RHE; the inset shows the equivalent electrical circuit image.