In situ preparation of an anatase/rutile-TiO$_2$/Ti$_3$C$_2$T$_x$ hybrid electrode for durable sodium ion batteries†

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Herein, a facile one-step method is developed to in situ prepare crystalline anatase and rutile TiO$_2$ nanocrystals on Ti$_3$C$_2$T$_x$ by regulating the metastable Ti ions. The combination of TiO$_2$ nanocrystals and Ti$_3$C$_2$T$_x$ not only introduces extensive accessible sites for Na$^+$ storage, but also promotes the charge transport by efficiently relieving the collapse of Ti$_3$C$_2$T$_x$. Compared with pristine Ti$_3$C$_2$T$_x$, the optimized TiO$_2$/Ti$_3$C$_2$T$_x$ hybrid electrode (anatase/rutile-TiO$_2$/Ti$_3$C$_2$T$_x$, A/R-TiO$_2$/Ti$_3$C$_2$T$_x$) exhibits a desirable specific surface area (22.5 m$^2$ g$^{-1}$), an ultralow charge transfer resistance (42.46 Ω) and excellent ion diffusion (4.01 x 10$^{-14}$). Remarkably, rich oxygen vacancies are produced on TiO$_2$/Ti$_3$C$_2$T$_x$ which is beneficial to enhance the insertion/de-insertion of Na$^+$ during the charge/discharge process. As a result, the A/R-TiO$_2$/Ti$_3$C$_2$T$_x$ delivers a high average capacity of 205.4 mA h g$^{-1}$ at 100 mA g$^{-1}$ and a desirable capacitance retention rate of 84.7% can be achieved after 600 cycles at 500 mA g$^{-1}$.

1. Introduction

Cost-effective energy storage systems have been one of the most promising candidates to face the challenges of the worldwide energy issues. Among various electrochemical energy storage techniques, the rechargeable lithium-ion batteries (LIBs) and sodium-ion batteries (SIBs) have attracted great interest due to their high energy density and flexible characteristics. However, the limited lithium resources increase the fabrication cost, thus hindering the development of LIBs. Compared to LIBs, SIBs have been considered as the most hopeful next-generation secondary battery due to their advantages of abundant sodium reserves. Unfortunately, the large diameter of Na$^+$ (116 pm) is always accompanied by the issues of slow reaction kinetics, low capacity and poor cycle stability when making SIB.

MXene, as a newly emerged 2D material, has exhibited prominent advantages, i.e. outstanding electronic conductivity and large surface area. Essentially, the MXene is proposed as an ideal matrix body to hybridize electrochemical active materials for SIB by inducing a synergistic effect. For example, Lv et al. developed Ti$_3$C$_2$T$_x$ by intercalation of ethanol or dimethyl sulfoxide. When employed as the anode for SIB, it achieved 110 mA h g$^{-1}$ at a current density of 100 mA g$^{-1}$, which was 16% higher than that of the original Ti$_3$C$_2$T$_x$ (95 mA h g$^{-1}$). However, due to the easy collapsing characteristics of MXene, the Na$^+$ storage capacity of the MXene anode is still far away from the theoretical value. To address this issue, various strategies have been proposed to alleviate the structure damage of MXene in the Na$^+$ storage process. For example, Sun et al. provided a surface regulation method, replacing the O-terminal with the PO$_2$-terminal, which greatly enhanced the specific capacity of Nb$_5$C$_3$ MXene and effectively relieves the collapse of MXene. Besides, the preparation of MXene based nanocomposites is also an efficient way to address this issue. The obvious advantage of the nanocomposite is to reduce the volume expansion during charge/discharge and thereby improve the stability of the battery. For example, Yang et al. in situ grew anatase TiO$_2$ on MXene towards the enlarged spacing layer of MXene. Meanwhile, the anatase TiO$_2$ nanoparticles also supply additional adsorption sites for sodium ions, which gives rise to a reversible capacity of 52 mA h g$^{-1}$ to 156 mA h g$^{-1}$ with a large current span of 50–2000 mA g$^{-1}$.

Due to the network structure enabled by the stacking and edge-sharing of TiO$_6$ octahedra, TiO$_2$ provides many ideal insertion sites for accommodating Na$^+$ and the suitable path for the diffusion of Na$^+$. Furthermore, among all kinds of crystallized TiO$_2$, rutile-TiO$_2$ exhibits good cycling stability due to its robust structure. Regarding TiO$_2$, it is known that the TiO$_2$ anode has the advantages of being non-toxic, low cost, high specific capacity and good rate performance. Nevertheless, the large band gap (3.0 eV) associated with TiO$_2$ causes the inferior electronic conductivity, which brings a negative impact on rate capability.

In this work, we aim to construct hybrid structures composed of Ti$_3$C$_2$T$_x$ and TiO$_2$ in different crystallinity for...
durable SIBs. It is explored that the coupling of TiO₂ with Ti₃C₂Tx not only benefits to support the 2D structure of Ti₃C₂Tx, but also introduces additional capacity enabled by TiO₂. Meanwhile, a large number of oxygen vacancies can be generated, which facilitates the insertion/de-insertion of sodium ions. Further, the agglomerate of TiO₂ can be relieved during the preparation. Owing to the synergistic effect, the A/R-TiO₂/Ti₃C₂Tx anode has proved superior Na⁺ storage performance.

2. Experimental

2.1 Materials

Ti₃AlC₂ (MAX phase, 325 mesh) powder was purchased from Shanghai Bowei applied material technology Co., Ltd. Hydrofluoric acid (HF-35%), hydrogen peroxide (H₂O₂), Na₂SO₄ and metallic sodium were purchased from Aladdin (China). All reagents are analytically pure and can be used as is without further purification.

2.2 Synthesis of Ti₃C₂Tx

1 g Ti₃AlC₂ powder was added into a 100 ml plastic beaker. Meanwhile, 12 ml hydrofluoric acid (HF-35%) was dropped into the beaker under an ice bath environment, and then continuously stirred at 45 °C for 48 hours. After that, the black precipitate was washed repeatedly with deionized water (DI) until it reached a neutral pH value. Finally, it was collected and dried at 60 °C to obtain the Ti₃C₂Tx powder.

2.3 Synthesis of A-TiO₂/Ti₃C₂Tx and A/R-TiO₂/Ti₃C₂Tx

0.2 g Ti₃C₂Tx powder, 80 ml DI, 0.5 ml H₂O₂ and 1 ml HCl (36%) were sent into a 100 ml stainless-steel autoclave with Teflon lining and maintained at 110 °C for 12 h. Subsequently, the precipitate was washed repeatedly with DI until the pH equals 7. Afterwards, the product was collected and dried at 60 °C to obtain the Ti₃C₂Tx. In terms of the synthesis of A/R-TiO₂/Ti₃C₂Tx, we followed the same procedure except for setting the reaction at 200 °C.

2.4 Synthesis of A-TiO₂ and A/R-TiO₂

0.2 g Ti₃C₂Tx powder, 80 ml DI, 4 ml H₂O₂ and 1 ml HCl (36%) were sent into a 100 ml stainless-steel autoclave with Teflon lining and maintained at 110 °C for 12 h. Subsequently, the precipitate was washed repeatedly with DI until the pH equals 7. Afterwards, the product was collected and dried at 60 °C to obtain the A-TiO₂. In terms of the synthesis of A/R-TiO₂, we followed the same procedure except for setting the reaction at 200 °C.

2.5 Material characterization

The morphology, structure and elemental distribution were characterized by scanning electron microscopy (SEM, Hitachi SU5000) and transmission electron microscopy (TEM, FEI Talos F-200X), X-ray diffraction (XRD, SmartLab) with Cu Kz radiation at 45 kV and 200 mA over a 2θ range of 5-85° was utilized to examine the internal crystalline structures of samples. The Raman spectra were measured on a spectrometer (DXR, USA) equipped with an optical microscope at room temperature. Various functional groups associated with samples were classified by Fourier transform infrared spectroscopy (FTIR) on a high-resolution infrared spectrometer (WQF-520A). The binding energy of the samples confirmed by X-ray photoelectron spectroscopy (XPS, Thermo ESCALAB 250Xi) was measured using the Al Kz ray (hv = 1486.6 kV) with a working voltage of 14.6 kV. The Brunauer–Emmett–Teller (BET) experiment was performed with a surface area analyzer (JW-BK200C). The pore size distribution plot was obtained according to the Barrett–Joyner–Halenda (BJH) method. The oxygen vacancy signal was obtained using a paramagnetic resonance spectrometer (EPR, Emsmicro).

2.6 Electrochemical characterization

The electrochemical behavior and SIB performance of Ti₃C₂Tx, A-TiO₂, A/R-TiO₂, A-TiO₂/Ti₃C₂Tx and A/R-TiO₂/Ti₃C₂Tx were studied by making a coin-type battery (CR2032). Initially, the working electrode was prepared by mixing the active materials, superconducting black and polyvinylidene fluoride (PVDF) in N-methyl-2-pyrrolidone (NMP) according to a mass ratio of 8 : 1 : 1, and then pasting evenly on copper foam. The average mass loading of the active material was about 2.0 mg cm⁻². Subsequently, the battery was assembled in a glove box filled with argon with H₂O and O₂ whose concentrations were below 0.1 and 0.3 ppm, respectively. The constant current charge discharge measurements were performed at room temperature using a battery program-controlled test system (CT2001A, LANHE, China) with a voltage range of 0 to 3 V. The cyclic voltammetry (CV) and electrochemical impedance spectroscopy (EIS) curves were measured using an electrochemical workstation (PARSTAT 3000A-DX, Princeton Applied Research, USA).

3. Results and discussion

3.1 Synthesis and materials characterization

Fig. 1a shows the schematic illustration of the structure of the TiO₂/Ti₃C₂Tx nanocomposite. There are a large number of metastable Ti ions present on both sides of Ti₃C₂Tx, which guarantees the building of the TiO₂/Ti₃C₂Tx hybrid.²²,²⁶,²⁷ By regulating the state of metastable Ti ions, the crystallinity of TiO₂ can be rationally controlled. Fig. 1b is the SEM image of Ti₃C₂Tx, exhibiting a typical accordion-like layered structure. Fig. 1c shows the SEM image of the A-TiO₂/Ti₃C₂Tx hybrid. Apparently, the accordion-like layered structure remains, while the surface of Ti₃C₂Tx is occupied by a large number of A-TiO₂ nanoparticles. When introducing R-TiO₂ into A-TiO₂/Ti₃C₂Tx, as shown in Fig. 1d, regular octahedral nanocrystals have been observed which are deemed as R-TiO₂. Fig. S1† shows the SEM images of A-TiO₂ and A/R-TiO₂, where the TiO₂ particles agglomerate together. The elemental mapping image of A/R-TiO₂/Ti₃C₂Tx, as shown in Fig. 1e, clearly demonstrates the co-existence of Ti, C and O. Fig. 1f shows the high-resolution transmission electron microscope (HRTEM) image of A/R-TiO₂/Ti₃C₂Tx with the enlarged HRTEM image of the selected area as well.
lattice fringes of $d = 0.342$ nm, 0.33 nm and 0.21 nm are associated with the (101) plane of A-TiO$_2$, (110) plane of R-TiO$_2$ and (105) plane of Ti$_3$C$_2$Tx, respectively, demonstrating the coexistence of A-TiO$_2$, R-TiO$_2$ and Ti$_3$C$_2$Tx-MXene. Consequently, the A/R-TiO$_2$/Ti$_3$C$_2$Tx hybrid is successfully synthesized.

Fig. 2a shows the X-ray diffraction (XRD) patterns of Ti$_3$C$_2$Tx, A-TiO$_2$/Ti$_3$C$_2$Tx and A/R-TiO$_2$/Ti$_3$C$_2$Tx. Basically, the characteristic peaks of Ti$_3$C$_2$Tx can be indexed in all samples, particularly two peaks at 9.08° and 18.22°. As a supplementary, the XRD patterns of A-TiO$_2$ and A/R-TiO$_2$ are given in Fig. S2.† In terms of the A-TiO$_2$/Ti$_3$C$_2$Tx, except for the characteristic peaks of Ti$_3$C$_2$Tx, a new peak located at 25.31° has emerged, which corresponds to the (101) plane of A-TiO$_2$. Upon the increase of the oxidation degree, a new peak appearing at 27.2° is assumed as the characteristic peak of the (110) plane of R-TiO$_2$, proving the presence of the A/R-TiO$_2$/Ti$_3$C$_2$Tx nanocomposite.† By using quantitative full pattern Rietveld refinement from the XRD diffraction patterns of each phase, we have calculated the mass ratio of MXene in TiO$_2$/MXene nanocomposites, which is 1 : 0.17 for MXene : A-TiO$_2$ in the A-TiO$_2$/Ti$_3$C$_2$Tx composite and 1 : 1.05 : 4.06 for MXene : A-TiO$_2$ : R-TiO$_2$ in A/R-TiO$_2$/Ti$_3$C$_2$Tx, respectively.

Fig. 2b and c presents the Raman and Fourier transform infrared spectroscopy (FTIR) spectra of all samples. In the Raman spectrum of Ti$_3$C$_2$Tx, a distinct peak between 140 cm$^{-1}$ and 180 cm$^{-1}$ refers to the typical C–C plane and the metastable Ti–Ti vibration modes. Besides, Ti–O vibration bands of surface groups arise at 202 cm$^{-1}$, 232 cm$^{-1}$, 256 cm$^{-1}$ and 267 cm$^{-1}$, which consist of Ti–OH, Ti–O bond and Ti–O–Ti stretching vibration of R-TiO$_2$ (O–Ti–O). Beyond that, the peak at 650 cm$^{-1}$ would be assigned to the (Ti–O) tensile zone, which is associated with A-TiO$_2$. Remarkably, in A/R-TiO$_2$/Ti$_3$C$_2$Tx, the band at 840 cm$^{-1}$ is attributed to the lattice stretching vibration of R-TiO$_2$ (O–Ti–O). Differing from the FTIR spectrum of Ti$_3$C$_2$Tx, a strong absorption band emerges from 500 cm$^{-1}$ to 900 cm$^{-1}$ in A-TiO$_2$/Ti$_3$C$_2$Tx due to the existence of A-TiO$_2$. Comparing the A/R-TiO$_2$/Ti$_3$C$_2$Tx, the band at 840 cm$^{-1}$ is attributed to the lattice stretching vibration of R-TiO$_2$ (O–Ti–O). Beyond that, the peak at 650 cm$^{-1}$ would be assigned to the (Ti–O) tensile zone, which is associated with A-TiO$_2$. Based on the above results, we have successfully synthesized the MXene/TiO$_2$ nanohybrids, in which the crystallinity of TiO$_2$ can be precisely controlled by regulating the metastable Ti ions through a facile one-step method.

Fig. 2d shows the O 1s spectra of Ti$_3$C$_2$Tx, A-TiO$_2$/Ti$_3$C$_2$Tx and A/R-TiO$_2$/Ti$_3$C$_2$Tx. Apparently, the C–O bond doesn’t have a violent change in all samples. In sharp contrast, the Ti–O bond is significantly enhanced in A/R-TiO$_2$/Ti$_3$C$_2$Tx, suggesting the presence of a large amount of TiO$_2$.† Fig. 2e gives the C 1s spectra, which can be split into C–C, Ti–C and C–O bonds. The C–C and Ti–C bonds reveal the signature of the Ti–C–Ti structure in all samples. The attenuation of Ti–C peaks in A-TiO$_2$/Ti$_3$C$_2$Tx and A/R-TiO$_2$/Ti$_3$C$_2$Tx is indicative of oxidation of surface Ti ions into TiO$_2$. In the Ti 2p spectrum (Fig. 2f), one pair of strong peaks located at 456.47 eV and 465.57 eV are correlated with the Ti$^{4+}$, while two peaks centered at 455.27 eV and 461.27 eV indicate the existence of the Ti–C bond in Ti$_3$C$_2$Tx. There was a controversy; a new peak of peaks occur
at about 458.57 and 464.27 eV, which are associated with the Ti-O bonds in A-TiO$_2$/Ti$_3$C$_2$T$_x$ and A/R-TiO$_2$/Ti$_3$C$_2$T$_x$. In addition, the peak area ratio of Ti-O/Ti-C ($\approx 2.57$) in A/R-TiO$_2$/Ti$_3$C$_2$T$_x$ is larger than that of A-TiO$_2$/Ti$_3$C$_2$T$_x$ ($\approx 0.81$), demonstrating the presence of a large amount of TiO$_2$.\textsuperscript{8,35,32}

Fig. 2g and h show the typical nitrogen adsorption–desorption isotherm and pore size distribution of Ti$_3$C$_2$T$_x$, A-TiO$_2$/Ti$_3$C$_2$T$_x$ and A/R-TiO$_2$/Ti$_3$C$_2$T$_x$, respectively. The relatively apparent hysteresis loop is examined on the isotherms of the three samples, implying the mesoporous characteristic. Compared with pure Ti$_3$C$_2$T$_x$, the formation of the TiO$_2$/Ti$_3$C$_2$T$_x$ heterojunction structure would bring in larger mesopores, which contributes to larger specific surface area. Incorporation of TiO$_2$ with different phases into MXene would further reinforce this effect, leading to increased specific surface area.\textsuperscript{39,40} Apparently, the N$_2$ adsorption capacities of the samples follow the order of Ti$_3$C$_2$T$_x$ < A-TiO$_2$/Ti$_3$C$_2$T$_x$ < A/R-TiO$_2$/Ti$_3$C$_2$T$_x$. The formation of TiO$_2$ nanocrystals on MXene provides more oxygen vacancies of A-TiO$_2$/Ti$_3$C$_2$T$_x$, which demonstrates to improve the electronic structure, charge adsorption sites for Na ions. Furthermore, we made BET measurements on the three samples, which show that the specific surface area (SSA) of Ti$_3$C$_2$T$_x$, A-TiO$_2$/Ti$_3$C$_2$T$_x$ and A/R-TiO$_2$/Ti$_3$C$_2$T$_x$ is 5.0, 6.0 and 22.5 m$^2$ g$^{-1}$, respectively. In strong contrast to Ti$_3$C$_2$T$_x$ and A-TiO$_2$/Ti$_3$C$_2$T$_x$, the dominant pore size of A/R-TiO$_2$/Ti$_3$C$_2$T$_x$ concentrates on 3–5 nm. The improved SSA with rational pore size makes A/R-TiO$_2$/Ti$_3$C$_2$T$_x$ possess more sodium ion storage sites, which could adapt to strain/stress during cycling and promote electrode/electrolyte contact.\textsuperscript{41,42}

Fig. 2i shows the electron paramagnetic resonance (EPR) spectra of Ti$_3$C$_2$T$_x$, A-TiO$_2$/Ti$_3$C$_2$T$_x$ and A/R-TiO$_2$/Ti$_3$C$_2$T$_x$. The signal response at 3515 gauss has been observed in all samples, corresponding to a g factor of 2.003, which is a characteristic of oxygen vacancies.\textsuperscript{44} In Ti$_3$C$_2$T$_x$, the oxygen vacancies may be related to the surface functional groups. After oxidation, the oxygen vacancies of A-TiO$_2$/Ti$_3$C$_2$T$_x$ have significantly increased, and the signal intensity is about 4.34 times higher than that of Ti$_3$C$_2$T$_x$. Additionally, the signal factor of A/R-TiO$_2$/Ti$_3$C$_2$T$_x$ is the strongest among all samples, with ~7.38 times higher signal intensity of Ti$_3$C$_2$T$_x$. Notably, oxygen vacancies have been demonstrated to improve the electronic structure, charge transfer, and surface properties of TiO$_2$ by narrowing the band gap, resulting in the accelerated Na$^+$ adsorption and diffusion kinetics.\textsuperscript{43–45}

3.2 Electrochemical performance and ion diffusion kinetics

The cycling performances of the Ti$_3$C$_2$T$_x$, A-TiO$_2$/Ti$_3$C$_2$T$_x$ and A/R-TiO$_2$/Ti$_3$C$_2$T$_x$ electrodes at 100 mA g$^{-1}$ are exhibited in Fig. 3a. The three electrodes realize good cycling stability. Remarkably, the A/R-TiO$_2$/Ti$_3$C$_2$T$_x$ electrode delivers a reversible capacity of 183.5 mAh g$^{-1}$ after 100 cycles, while the reversible capacities of Ti$_3$C$_2$T$_x$ and A-TiO$_2$/Ti$_3$C$_2$T$_x$ are separately 96.8 and 107.9 mAh g$^{-1}$. As a comparison, Fig. S3† shows the reversible capacities of A-TiO$_2$ and A/R-TiO$_2$ after 100 cycles, which are 55.6 and 118.7 mAh g$^{-1}$, respectively. Both are far less than the capacity of the A/R-TiO$_2$/Ti$_3$C$_2$T$_x$ anode. Further, Fig. 3b shows that the rate capacities of the A/R-TiO$_2$/Ti$_3$C$_2$T$_x$ electrode are 205.4, 135.7, 103.5, 77.5 and 40.6 mAh g$^{-1}$ at 100, 500, 1000, 2000 and 5000 mA g$^{-1}$. These values are considerably higher than those for the A-TiO$_2$/Ti$_3$C$_2$T$_x$ and Ti$_3$C$_2$T$_x$ electrodes, which are 132.4 mAh g$^{-1}$, 78.3 mAh g$^{-1}$, 50.1 mAh g$^{-1}$, 25.6 mAh g$^{-1}$, and 11.4 mAh g$^{-1}$ and 109.9 mAh g$^{-1}$, 59.7 mAh g$^{-1}$, 31.1 mAh g$^{-1}$, 11.4 mAh g$^{-1}$, and 12.6 mAh g$^{-1}$, respectively.

Fig. 3c displays the first four continued CV curves of the A/R-TiO$_2$/Ti$_3$C$_2$T$_x$ with potential windows of 0.01–3.0 V at 0.1 mV s$^{-1}$. As illustrated, the irreversible oxidation peak in the anodic process for the first cycle can be ascribed to the decomposition of the electrolyte, the formation of the solid electrolyte interface (SEI), or the irreversible muti-electrochemical reactions.\textsuperscript{46} In the case of the first sodiation process, two broad anodic peaks occurred at 1.39 V and 2.03 V, respectively, and their intensity decreased compared with the subsequent cycles, corresponding to the extraction of sodium ions from the Ti$_3$C$_2$T$_x$ electrode. No significant shift was observed in subsequent cycles, suggesting that the charge storage in Ti$_3$C$_2$T$_x$ was due to the intercalation of Na$^+$ rather than a conversion reaction.\textsuperscript{32,34} In the following three subsequent cycles, a cathodic peak near 2.35 V can be clearly observed due to the sodiation, while an anodic peak near 0.45 V is attributed to the de-sodiation. This reversible process is accompanied by the conversion of Ti$^{4+}$/Ti$^{3+}$ couples in the A/R-TiO$_2$/Ti$_3$C$_2$T$_x$.\textsuperscript{14,37–46} Beyond the first cycle, the subsequent cycles are almost overlapping, implying the high reversibility of the electrode. The selected charge–discharge curves of A/R-TiO$_2$/Ti$_3$C$_2$T$_x$ within 0–3.0 V at a current density of 100 mA g$^{-1}$ are drawn in Fig. 3d. The initial discharge and charge capacity were 774.7 and 270.9 mAh g$^{-1}$, respectively, delivering a Coulomb efficiency (CE) of 35%. The low CE and huge irreversible capacity loss are mainly resulted from the formation of SEI film and irreversible electrochemical reaction.\textsuperscript{14,47} Interestingly, the charge–discharge curves after the first cycles are going to overlap, suggesting the improved reversibility of the A/R-TiO$_2$/Ti$_3$C$_2$T$_x$.\textsuperscript{14,47}
Ti$_3$C$_2$T$_x$ electrode. Fig. 3e illustrates the long cycling performance of A/R-TiO$_2$/Ti$_3$C$_2$T$_x$ at 500 mA g$^{-1}$ in 600 cycles. Essentially, the reversible initial and final capacities are 128.8 and 109.1 mA h g$^{-1}$, respectively, deducing a capacitance retention rate of 84.7%.

To figure out the charge-storage mechanism of the A/R-TiO$_2$/Ti$_3$C$_2$T$_x$ electrode, cyclic voltammetry (CV) profiles at various scan rates were tested. Fig. 3e illustrates the CV curves of A/R-TiO$_2$/Ti$_3$C$_2$T$_x$ at scan rates of 0.1, 0.3, 0.5, 0.7, 1.0 and 2.0 mV s$^{-1}$, respectively. Classically, the charge-storage mechanism can be investigated by exploring the relationship between the peak current ($i_p$, mA) and the scan rates ($v$, mV s$^{-1}$):

$$i = av^b$$

$$\log(i) = b \log(v) + \log(a)$$

where $a$ and $b$ are adjustable parameters. When $b$ equals 1, the capacity is fully governed by the capacitive-controlled process; if $b$ equals 0.5, it corresponds to a diffusion-controlled process. Fig. S4† shows that the $b$ values of A/R-TiO$_2$/Ti$_3$C$_2$T$_x$ are 0.888 and 0.910, which realizes the mixed behavior of diffusion-controlled and capacitive-controlled process. Further, the ratio of contribution of capacitive-controlled process and diffusion-controlled process can be quantitatively estimated through the following equations:

$$i(v) = k_1v + k_2v^{1/2}$$

$$i(v)/v^{1/2} = k_1v^{1/2} + k_2$$

where $i(v)$, $k_1$, $k_2$, $v^{1/2}$ and $v$ denote the current at a fixed potential, the capacitive-controlled current, the diffusion-controlled current and the scan rate, respectively. The capacitance contribution of the capacitive-controlled process at a given scan rate can be figured out by obtaining $k_1$. Through linear fittings of the $v^{1/2}$ vs. $i(v)/v^{1/2}$ plots, $k_1$ values at different scan rates are determined.**48** As shown in Fig. 3g and h, as the scan rate increased from 0.1 mV s$^{-1}$ to 2 mV s$^{-1}$, the capacitive contribution grew from 56.1% to 84.0% of the total stored charge, which implies that the electrochemical process is determined by both intercalation and capacitive reactions and the capacitive charge storage contributes most to the total capacity at high rates.

Fig. 4a shows the Nyquist plot and equivalent circuit diagrams of the three electrodes at a frequency of 0.01 to 10$^6$ Hz. As shown in Fig. 4a, all plots exhibit similar shapes composed of a straight line in the low frequency area and a semicircle in the high frequency range.**48** The equivalent circuit is given in the inset, where $R_n$, $R_c$, and $W_0$ represent the electrolyte resistance, the interfacial charge transfer resistance and the Warburg impedance related to the Na$^+$ diffusion within the electrode.**50** As shown in Table 1, $R_c$ significantly decreases when TiO$_2$ is incorporated in Ti$_3$C$_2$T$_x$ regardless of the crystal type. This can be attributed to the improved conductivity and increased electroactive sites. Moreover, the $R_c$ of A/R-TiO$_2$/Ti$_3$C$_2$T$_x$ is 42.46 $\Omega$, which is much lower than that of A-TiO$_2$/Ti$_3$C$_2$T$_x$ (119 $\Omega$), indicating the high interfacial charge transfer which is beneficial to enhance the rate capability. Except for the resistance, we calculated the diffusion coefficient of sodium ions ($D_{Na}$) from the low frequency domain, which can be expressed by the following equations:

$$D = \frac{R^2T^2}{2A^2n^2F^2C^2\sigma^2}$$

$$Z' = R_c + R_{ct} + \sigma\omega^{1/2}$$

$R$ denotes the molar gas constant (8.314 J K$^{-1}$ mol$^{-1}$), $T$ is the absolute temperature (298.15 K), $A$ is the area of the electrode, $n$ is the number of electrons, $F$ is the Faraday constant (96485 C mol$^{-1}$), $C$ is the absolute concentration of Naions ($10^{-3}$ mol L$^{-1}$), $\sigma$ is the sodium ion diffusion coefficient ($10^{-5}$ cm$^2$ s$^{-1}$), and $\omega$ is the angular frequency ($s^{-1}$).

![SEM images of (a and b) the pure Ti$_3$C$_2$T$_x$ and (c and d) A/R-TiO$_2$/Ti$_3$C$_2$T$_x$ electrode before/after 100 charging–discharging cycles, respectively.](image_url)

**Table 1** The charge transfer impedance, weber factor and diffusion coefficient of Ti$_3$C$_2$T$_x$, A-TiO$_2$/Ti$_3$C$_2$T$_x$ and A/R-TiO$_2$/Ti$_3$C$_2$T$_x$.

| Electrode          | $R_{ct}$/\Omega | $\sigma$/ohm cm$^2$ s$^{-0.5}$ | $D_{Na}$/cm$^2$ s$^{-1}$ |
|--------------------|-----------------|-------------------------------|--------------------------|
| Ti$_3$C$_2$T$_x$   | 289.8           | 57                            | 2.858 $\times$ 10$^{-15}$ |
| A-TiO$_2$/Ti$_3$C$_2$T$_x$ | 119             | 55.1                          | 8.602 $\times$ 10$^{-15}$ |
| A/R-TiO$_2$/Ti$_3$C$_2$T$_x$ | 42.46           | 48                            | 4.01 $\times$ 10$^{-14}$  |

**Fig. 4** (a) EIS spectra and Nyquist plots, and (b) relationship between $Z'$ and $\omega^{-1/2}$ at low frequencies.
is the number of electrons transferred during the reaction, \( C \) is the sodium ion concentration and \( P \) is the Faraday constant (96 486 C mol\(^{-1}\)). Fig. 4b shows the relationship between \( Z \) and \( \omega^{-1/2} \) at low frequencies.\(^{32}\) The values of \( \sigma \) and the \( D_{\text{Na}^+} \) are summarized in Table 1. The \( D_{\text{Na}^+} \) of A/R-TiO\(_2\)/Ti\(_3\)C\(_2\)T\(_x\) is 4.01 \( \times \) \( 10^{-14} \) cm\(^2\) s\(^{-1}\) which is approximately 14.01 times and 4.66 times higher than the \( D_{\text{Na}^+} \) of Ti\(_3\)C\(_2\)T\(_x\) (2.858 \( \times \) \( 10^{-15} \) cm\(^2\) s\(^{-1}\)) and A-TiO\(_2\)/Ti\(_3\)C\(_2\)T\(_x\) (8.602 \( \times \) \( 10^{-15} \) cm\(^2\) s\(^{-1}\)), respectively. This is presumably ascribed to the incorporation of long rod-like R-TiO\(_2\) nanocrystals, which can effectively relieve the collapse of Ti\(_3\)C\(_2\)T\(_x\) layers.

Fig. 5a and b show the SEM images of the Ti\(_3\)C\(_2\)T\(_x\) electrode before and after the cycling, respectively. Obviously, the layered structure of pure Ti\(_3\)C\(_2\)T\(_x\) seriously collapsed after cycling. In sharp contrast, the structure of A/R-TiO\(_2\)/Ti\(_3\)C\(_2\)T\(_x\) remains largely unaffected (Fig. 5c and d), illustrating its desirable stability.

4. Conclusions

Through the surface regulation of Ti\(_3\)C\(_2\)T\(_x\), we generated A-TiO\(_2\) and R-TiO\(_2\) \textit{in situ} from metastable Ti ions on the surface of Ti\(_3\)C\(_2\)T\(_x\) by a one-step method to obtain nano hybrid structures. The growth of crystalline A-TiO\(_2\) and R-TiO\(_2\) is beneficial to improve the collapse of Ti\(_3\)C\(_2\)T\(_x\). The 2D structure can provide a fast channel for the transmission of Na\(^+\). A-TiO\(_2\) and R-TiO\(_2\) crystals can introduce more active sites for Na\(^+\). Moreover, the crystals can provide a supporting effect for the layered structure and maintain the integrity and firmness of the structure. Meanwhile, a large number of oxygen vacancies can be introduced into the composite which are conducive to the insertion and extraction of Na\(^+\). Due to the advantages of the A/R-TiO\(_2\)/Ti\(_3\)C\(_2\)T\(_x\) hybrid structure, the prepared A/R-TiO\(_2\)/Ti\(_3\)C\(_2\)T\(_x\) electrode has proved high capacity, great rate performance and long-term cycle stability. We believe that these findings will provide new ideas for the design of nanostructured electrodes for advanced rechargeable batteries.

Conflicts of interest

There are no conflicts to declare.

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