High-Thermal-Stability and High-Thermal-Conductivity Ti$_3$C$_2$T$_x$ MXene/Poly(vinyl alcohol) (PVA) Composites

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ABSTRACT: MXenes, a new family of two-dimensional materials, have recently attracted increasing attention due to their unique properties for a wide range of potential applications. Herein, we synthesize Ti$_3$C$_2$T$_x$/poly(vinyl alcohol) (PVA) composites and investigate the effects of the thermal properties of MXene by temperature-dependent Raman spectroscopy and polarized-laser power-dependent Raman spectroscopy. Compared to the Ti$_3$C$_2$T$_x$ MXene, the PVA significantly improves the thermal stability of Ti$_3$C$_2$T$_x$ by reducing the thermal coefficient of the E$_2^\text{g}$ mode from −0.06271 to −0.03357 cm$^{-1}$/K, which is attributed to the strong Ti−O bonds formed between the MXene and PVA polymer confirmed by the X-ray photoelectron spectroscopy results. Meanwhile, the thermal conductivities of Ti$_3$C$_2$T$_x$ and Ti$_3$C$_2$T$_x$/PVA composites reach as high as about 55.8 and 47.6 W/(m K), respectively. Overall, this work will contribute to extend the MXene applications in terms of polymer-based nanocomposites and improve the reliability of the related devices effectively.

1. INTRODUCTION

Since the discovery of graphene, two-dimensional (2D) atomic crystals have attracted increasing scientific research interest owing to their high surface areas and unique properties that differ from the corresponding three-dimensional analogues. Among these materials, MXenes, a recently discovered family of 2D transition-metal carbides or carbonitrides, shed new light on the electronic properties of these materials can be electronically tuned by intercalation. MXene/N-doped graphene demonstrates a great potential in water purification and sensors. In addition, theoretical studies suggest that several members of this family, such as Ti$_3$C and Mo$_2$C, can become potential candidates for hydrogen storage and thermoelectric applications. Meanwhile, MXenes can also be used to produce functional nanocomposites due to their functionalized surfaces and 2D morphology. The key factor to introduce polymer matrices into MXenes is the utilization of cationic-charged or hydrogen-bond-forming polymers to achieve a strong interaction with negatively charged OH/F/O-terminated MXene surface. Previously, Ti$_3$C$_2$T$_x$ MXene was attempted to mix with charged diallyldimethylammonium chloride, polydiallyldimethylammonium, electrically neutral poly(vinyl alcohol) (PVA), or polyacrylamide (PAM) to produce Ti$_3$C$_2$T$_x$/polymer nanocomposite systems. Using dimethylsulfoxide, uniform and flexible Ti$_3$C$_2$T$_x$/PAM composites with only 6 wt % (1.7 vol %) MXene loading were obtained with an electrical conductivity of about 3.3 × 10$^{-2}$ S/m. Ling et al. reported that the Ti$_3$C$_2$T$_x$/PVA composites showed electrical conductivity as high as 2.2 × 10$^2$ S/m, whereas the measured electrical conductivity of pure Ti$_3$C$_2$T$_x$ films was about 2.4 × 10$^5$ S/m. In addition, the tensile strength of the Ti$_3$C$_2$T$_x$/PVA composites was significantly enhanced from 22 ± 2 to 30 ± 10 MPa compared to the pure Ti$_3$C$_2$T$_x$ films. The introduction and confinement of the polymer between the MXene flakes not only increased flexibility, but also stimulated cationic intercalation, offering an impressive volumetric capacitance of ~530 F/cm$^3$ for MXene/PVA–KOH composite films at 2 mV/s. This implies that MXenes are promising fillers in multifunctional polymer composites, which in turn can be employed in many...
applications such as flexible and wearable energy-storage devices, radio frequency shielding, water filtration system, and so on. Raman spectroscopy, a nondestructive method, is typically used to investigate the strain distribution and thermal stability of a material by assessing its Raman frequency shifts and related thermal coefficients. In light of the importance of MXene/polymer composite systems, in this study, we fabricated uniform Ti$_3$C$_2$Tx/PVA composites and explored their thermal stability and thermal conductivity by temperature-dependent and polarized-laser power-dependent Raman measurements. The results demonstrate that the introduction of PVA significantly improves the thermal stability of the composites in comparison to the pure Ti$_3$C$_2$Tx. Meanwhile, the thermal conductivity of Ti$_3$C$_2$Tx/PVA composites also stays at high level. These results will have a great impact on the reliability of the related devices.

2. EXPERIMENTAL SECTION

Figure 1 provides a general overview of the fabrication process of Ti$_3$C$_2$Tx/PVA composites. A piece of the Ti$_3$AlC$_2$ monolith (1 g) was immersed into 10 mL of 50 wt% hydrofluoric acid solution for 18 h at room temperature to dissolve the Al atoms. The resulting suspension of multilayered Ti$_3$C$_2$Tx was repeatedly washed with deionized water six to eight times, centrifuged at 1500 rpm for 10 min, and decanted until the supernatant reached a pH of ~6. The separated wet sediment was dried under vacuum for 24 h before further experiments. PVA with a molecular weight of about 67000 g/mol in 1 wt% aqueous solutions was prepared. The Ti$_3$C$_2$Tx/PVA composites were characterized by a scanning electron microscope (SEM) (Hitachi SU-70) and a transmission electron microscope (TEM) (Tecnai G2 F20 S-Twin) operated at an accelerating voltage of 200 kV. Thermogravimetric analysis was performed by a thermal gravimetric analyzer (Netzsch, TG 209F1) in nitrogen atmosphere and oxygen atmosphere. The heating rate was 10 K/min. Raman spectroscopy measurements were carried out in a confocal microscopy setup with a 532 nm solid-state green laser for excitation. The temperature ranging from 300 K (room temperature) to 480 K was controlled by a heating stage. Polarized-laser power-dependent Raman measurements were performed at 0.01P$_0$, 0.1P$_0$, 0.25P$_0$, 0.5P$_0$, 0.75P$_0$, and P$_0$ where P$_0$ = 2.5 mW. And thermal conductivities of the pristine Ti$_3$C$_2$Tx MXene and Ti$_3$C$_2$Tx/PVA composites were also confirmed by a laser thermal conductivity test instrument (Netzsch, LFA 427) at room temperature. X-ray photoelectron spectroscopy (XPS) studies were carried out in a VG ESCALAB 220i-XL system equipped with a monochromatized Al Kα X-ray source (1486.6 eV), a multichannel plate, and a delay-line detector under a vacuum of 1 × 10$^{-7}$ mbar. High-resolution (HR) spectra were collected at a constant pass energy of 20 eV and quantified using empirically derived relative sensitivity factors provided by Kratos Analytical. Binding energies were referenced to the C 1s peak of the C−C bond at 284.80 eV. The data were analyzed by commercially available software CasaXPS.

3. RESULTS AND DISCUSSION

Figure 2 shows the XRD patterns of Ti$_3$AlC$_2$, as-prepared Ti$_3$C$_2$Tx powders, and the corresponding Ti$_3$C$_2$Tx/PVA composites. It can be readily observed that the (002) peak of the Ti$_3$AlC$_2$ MAX phase shifts toward lower angles from about 9.60$^\circ$–8.60$^\circ$ (Ti$_3$C$_2$Tx MXene) and 7.80$^\circ$ (Ti$_3$C$_2$Tx/PVA composites), which indicates the increase of the c lattice parameter. According to the Bragg formula 2d sin $\theta = n\lambda$, where $\theta$ is the Bragg angle, d is the interplanar spacing, and $\lambda$ is the wavelength of the incident X-ray (0.15406 nm), the d parameters of the Ti$_3$AlC$_2$ MAX, Ti$_3$C$_2$Tx MXene, and Ti$_3$C$_2$Tx/PVA composites are about 18.34, 20.54, and 22.10 Å, respectively. This demonstrates that the PVA can interact with Ti$_3$C$_2$Tx MXene. The surface functional groups of MXene are changed, which enlarges the d-spacing of MXene sheets.

Figure 1. Schematic diagram of the fabrication process of Ti$_3$C$_2$Tx/PVA composites.

Figure 2. XRD patterns of Ti$_3$AlC$_2$, Ti$_3$C$_2$Tx, MXene, and Ti$_3$C$_2$Tx/PVA composites.
addition, the broadening of the (002) peaks in the Ti$_3$C$_2$Tx MXene compared to that in the MAX phase mainly attributes to the decreasing structure order.

The as-synthesized Ti$_3$C$_2$Tx has a layered morphology resembling exfoliated graphite, as shown in Figure 3a. The enlarged view of Ti$_3$C$_2$Tx (Figure 3b) indicates the clear layered structure. However, in Figure 3c, after introduction of PVA, the sheets stack together because the hydroxyl in the PVA reacted with the −OH-terminated MXene surface (−OH + −OH = −O− + H$_2$O) and formed Ti−O bonds at the surface, which is confirmed by the XPS results. The enlarged view of Ti$_3$C$_2$Tx/PVA composites (Figure 3d) indicates compact layered structure. To further explore the structure changes of Ti$_3$C$_2$Tx MXene after the PVA reaction, the material is also characterized by TEM, as displayed in Figure 4. Figure 4a,d exhibit the two-dimensional nanosheets of Ti$_3$C$_2$Tx and Ti$_3$C$_2$Tx/PVA, respectively. The exfoliated slices are very thin and disperse as single layer or stack of several layers. The inset selected area electron diffraction (SAED) patterns in Figure 4b,e show that the crystal planes of the MXenes are hexagonal systemic, indicating that the PVA does not change the crystal type. Furthermore, the space parameters of both the Ti$_3$C$_2$Tx and Ti$_3$C$_2$Tx/PVA composites corresponding to (1010) planes are measured to be about 0.2656 and 0.2595 nm, respectively. On the basis of the above values, the lattice constant $a$ is about 0.3070 nm for Ti$_3$C$_2$Tx and 0.2982 nm for Ti$_3$C$_2$Tx/PVA. Therefore, it demonstrates that the original basal plane structure of the Ti$_3$C$_2$Tx MXene shrinks a little after the PVA treatment coupled with the lattice expansion along the c axis. Figure 4c,f shows the cross sections of the multilayered Ti$_3$C$_2$Tx MXene and Ti$_3$C$_2$Tx/PVA composites, respectively. The nanosheets are stacked less uniformly in Ti$_3$C$_2$Tx/PVA than in Ti$_3$C$_2$Tx due to the polymer reaction.

Figure 5a shows the TG curve of Ti$_3$C$_2$Tx in nitrogen atmosphere. The temperature is from room temperature to 950 °C. There are three weight-change steps. The first step is due to the release of free water adsorbed on the surface of Ti$_3$C$_2$Tx. The second step is due to the release of bonding water and functional groups on the surface of Ti$_3$C$_2$Tx. The third step is due to the decomposition of Ti$_3$C$_2$Tx MXene. The decomposition temperature of pure Ti$_3$C$_2$Tx MXene is about 785 °C. Figure 5b shows the TG curve of Ti$_3$C$_2$Tx/PVA composites in nitrogen atmosphere. The temperature is also from room temperature to 950 °C. There are four weight-change steps compared to Ti$_3$C$_2$Tx MXene. The temperature from 220 to 480 °C is the decomposition temperature of PVA. The content of PVA is 12.71% in the composites in our experiment. The decomposition temperature of Ti$_3$C$_2$Tx/PVA composites is about 823 °C, which is higher than that of Ti$_3$C$_2$Tx MXene. So, the thermal stability of Ti$_3$C$_2$Tx/PVA composites is better than that of Ti$_3$C$_2$Tx MXene in nitrogen atmosphere. Figure 5c shows the TG curve of Ti$_3$C$_2$Tx in oxygen atmosphere. The temperature is from room temperature to 950 °C. There are four weight-change steps. The oxidation temperature of Ti$_3$C$_2$Tx MXene is about 322 °C, the decomposition temperature of PVA is 12.71%, and the decomposition temperature of MXene is about 729 °C. Figure 5d shows the TG curve of Ti$_3$C$_2$Tx/PVA composites in oxygen atmosphere. The temperature is from room temperature to 950 °C. There are five weight-change steps. The oxidation temperature of the composites is about 530 °C, and the decomposition temperature of the composites is about 811 °C. The result indicates a better thermal stability of Ti$_3$C$_2$Tx/PVA composites than pure Ti$_3$C$_2$Tx MXene in oxygen atmosphere. So, from our TG results, the introduction of PVA can improve the thermal stability of Ti$_3$C$_2$Tx in both nitrogen and oxygen atmosphere.

Raman spectroscopy is a powerful tool to investigate the atomic bonds, thermal stability, and thermal conductivity of two-dimensional materials, such as black phosphorus, MoS$_2$,
In this work, temperature-dependent Raman measurements ranging from 300 to 480 K were carried out for both Ti$_3$C$_2$T$_x$ MXenes and Ti$_3$C$_2$T$_x$/PVA composites. Raman frequency spectra with increasing temperature are plotted in Figure 6a,c. In Ti$_3$C$_2$T$_x$, the Raman frequency of about 156 cm$^{-1}$ (E$_g^1$ mode) at room temperature is mainly from the in-plane vibrations of Ti$_2$ and C atoms, as illustrated in Figure 6e. Besides, the Raman peaks at 606 cm$^{-1}$ (E$_g^2$ mode) and 440 cm$^{-1}$ in the Ti$_3$C$_2$T$_x$/PVA composite are attributed to the in-plane vibrations of C and O atoms, respectively. However, the E$_g^1$ mode shifts to a lower wave number at about 146.4 cm$^{-1}$ in the Ti$_3$C$_2$T$_x$/PVA composite. It can be inferred that the PVA polymer weakens the in-plane motion of the Ti$_2$ and C atoms. In comparison to the bare Ti$_3$C$_2$ nanosheet, the T-terminated Ti$_3$C$_2$ normally has a smaller $a$ parameter and a larger $d$ spacing while the atoms remain in the same Wyckoff positions. The XRD and TEM results have demonstrated that the $c$ and $a$ lattice parameters are enlarged about 1.56 Å and reduced 0.088 Å, respectively, after the PVA treatment. As a result, the Ti$_2$ atoms are pushed...
out a little from the plane, then the bond length between the Ti1 and C atoms shrinks, whereas that between the Ti2 and C atoms is elongated, and in turn weakens the in-plane vibrations of Ti2 and C atoms. From Figure 6b,d, the Raman phonon mode $E_g^1$ obviously exhibits red shift from 300 to 480 K and can be fitted by a linear equation: $\omega = \omega_0 + \chi \Delta T$, where $\omega_0$ is the mode frequency at room temperature, $\chi$ is the first-order temperature coefficient, and $\Delta T$ is the temperature difference relative to 300 K. The measured $\chi$ value for the Ti$_3$C$_2$Tx/PVA composites is about $-0.03357 \pm 0.00023$ cm$^{-1}$/K, which is much smaller than the corresponding value $-0.06271 \pm 0.00045$ cm$^{-1}$/K of the Ti$_3$C$_2$Tx MXenes, indicating a better thermal stability in the former. Several data point dispersions are attributed to slight variation in the laser spot focused on the material. The observed softening of the Raman peaks with increasing temperature arises from an anharmonicity, which is related to the harmonic potential constant, the phonon occupation, as well as the thermal expansion of the crystal.$^{36}$ The improvement of thermal stability in the Ti$_3$C$_2$Tx/PVA composites is primarily contributed to the surface modification introduced by the PVA treatment. Therefore, it is essential to explore the change of the chemical bonding.

XPS investigations were performed to characterize the chemical compositions of the prepared MXene powders and composites. A detailed survey spectrum for both samples shows the presence of Ti, oxygen (O), fluorine (F), and carbon (C) (not shown here). The relative high-resolution XPS images of Ti 2p and O 1s are shown in Figure 7. The Ti 2p component centered at 455.9 eV is associated with the Ti–C bonds, whereas that at about 461 eV can be assigned to the Ti–O and Ti–F bonds, which are totally referred as Ti–O$_x$F$_y$.$^{37,38}$ Compared to the pure Ti$_3$C$_2$Tx MXene, the peak integrated intensity ratio of Ti–O$_x$F$_y$ to Ti–C increases from about 0.425 to 0.693 in the Ti$_3$C$_2$Tx/PVA composite, which indicates that more Ti–O$_x$F$_y$ bonds are formed after PVA intercalation. Similarly, the O 1s XPS images with Ti–O bond at about 529.0 eV, Ti–OH bond at 532.0 eV,$^{39}$ and C–O bond at 533.1 eV are presented in Figure 7b,d. The intensity ratio of Ti–O bond to Ti–OH bond is about 0.657 in Ti$_3$C$_2$Tx MXene, whereas that in the Ti$_3$C$_2$Tx/PVA composites increases to about 0.859, which indicates that more Ti–O bonds are formed after the PVA treatment. The result is consistent with the results of Ti 2p XPS. Overall, the PVA polymer can conjugate with the Ti$_3$C$_2$Tx MXene effectively in terms of Ti–O bond formation Ti$_3$C$_2$Tx/PVA composites. In Ti$_3$C$_2$Tx MXenes, the bond lengths of Ti–O, Ti–F, and Ti–OH are 1.97, 2.17, and 2.19 Å, respectively, indicating that Ti–O bond is the most stable one among the terminal groups.$^{40}$ Generally, the strong terminal Ti–O bond significantly weakens the vibrations in which the surface Ti atoms are involved and increases the material thermal stability greatly, especially in the Ti$_3$C$_2$Tx/PVA composites where the PVA polymer profoundly increases the
relative intensity of Ti−O bond declared by the XPS results. Otherwise, one PVA molecular (C2H4O)n includes much −OH groups, which promotes PVA polymer cladding on the Ti3C2Tx MXene effectively. This is also helpful to improve the thermal stability and prevent Ti3C2Tx MXene from oxidation and rolling into conical shape. Thus, the PVA polymer has great impacts on the thermal properties of MXenes, and it is reasonable to assume that it would be possible to tune the thermal stability or thermal conductivity of the Ti3C2Tx/PVA composites by controlling the Ti3C2Tx-to-PVA ratio.

To obtain the thermal conductivity, polarized-laser power-dependent Raman spectra were recorded, as displayed in Figure 8a. The maximum laser power \( P_0 \) was about 2.5 mW measured at the exit of the microscope lens and can be reduced using an optical attenuator. Obvious frequency red shift of the Raman peaks with increasing laser power was discovered, which arose from the local laser heating effect. Figure 8c,d show the red shift of the \( E_g^1 \) mode frequency derived from the power-dependent Raman spectra. The \( \partial \omega/\partial p \) coefficients for the Ti3C2Tx MXene and Ti3C2Tx/PVA composite films are about 968 and 1081 cm−1/W, respectively. For two-dimensional materials with thin-film structure, the thermal conductivity can be expressed as

\[
k = \chi (1/2 \pi h) (\partial \omega/\partial p)^{-1}
\]

where \( \chi \) is the first temperature coefficient of the \( E_g^1 \) for the Ti3C2Tx MXene and Ti3C2Tx/PVA composite films and \( h \) is the thickness of the film. Taking the related values into eq 1, the thermal conductivities of the Ti3C2Tx and Ti3C2Tx/PVA composite films are about 55.8 and 47.6 W/(m K), respectively.

To confirm the thermal conductivity results, the laser flash method was also performed. The samples were cut into circular...
Figure 7. High-resolution Ti 2p and O 1s XPS images of (a, b) Ti$_3$C$_2$T$_x$ MXene and (c, d) Ti$_3$C$_2$T$_x$/PVA composite.

Figure 8. Polarized-laser power-dependent Raman spectra of (a) Ti$_3$C$_2$T$_x$ MXene and (b) Ti$_3$C$_2$T$_x$/PVA composite films under different laser power. Power dependence of E$_{g}^{1}$ mode of (c) Ti$_3$C$_2$T$_x$ MXene and (d) Ti$_3$C$_2$T$_x$/PVA composite films including linear fits of the experimental data.
specimens with a diameter of 20 mm and a thickness of 3 mm. The experiments were carried at room temperature. The thermal conductivities of the Ti$_3$C$_2$Tx and Ti$_3$C$_2$T$_x$/PVA composite films are tested for 55.2 ± 1.7 and 47.3 ± 1.2 W/(m K), respectively, which are in accordance with the polarized-laser power-dependent Raman test results. With the introduction of PVA, the conductivity of the Ti$_3$C$_2$T$_x$/PVA composites decreases little from pure Ti$_3$C$_2$Te MXene. And the conductivity of the Ti$_3$C$_2$T$_x$/PVA composites can still surpass that of Fe (47.4 W/(m K)), SiO$_2$ (7.6 W/(m K)), Al$_2$O$_3$ (45 W/(m K)), stainless steel (40 W/(m K)), and most of the other two-dimensional materials. Our results show that the Ti$_3$C$_2$T$_x$/PVA composites exhibit high thermal stability and thermal conductivity and can be used in many areas, such as supercapacitors, Li-ion batteries, solar cells, electromagnetic shielding, and so on.

4. SUMMARY

In conclusion, high-thermal-stability and high-thermal-conductivity Ti$_3$C$_2$T$_x$/PVA composites is prepared in our work. The PVA polymer on the thermal properties of Ti$_3$C$_2$T$_x$ MXene are investigated. The PVA polymer significantly improves the thermal stability of the MXene by introducing strong Ti–O bonds and cladding on the surface of the Ti$_3$C$_2$T$_x$ MXene without changing the crystal structure largely. The thermal conductivities of Ti$_3$C$_2$T$_x$ and Ti$_3$C$_2$T$_x$/PVA are about 55.8 and 47.6 W/(m K), respectively, which are even higher than those of some metals and most of the other two-dimensional materials. This study makes a crucial step to explore the potential use of MXene in polymer-based nanocomposites for a host of applications (e.g., energy-storage devices, water filtration system, thermoelectric devices, and Aeronautics and Astronautics materials) in harsh environment.

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Notes
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