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Structural, optical and mechanical characterization of PMMA-MXene composites functionalized with MEMO silane

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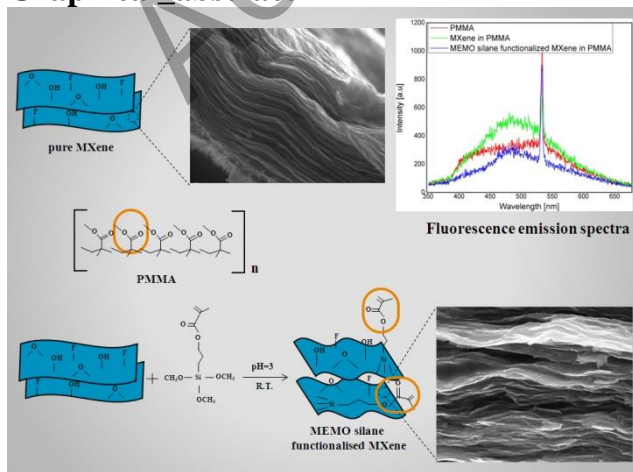
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Abstract

Processing and characterization of PMMA-MXene composites were investigated. γ -Methacryloxypropyltrimethoxy (MEMO) silane was used to modify the surface of MXenes and improve the compatibility between MXenes and the polymer. The FTIR analysis revealed the formation of a chemical bond between MXene and MEMO silane, while the XPS analysis confirmed the presence of silicon in the functionalized MXene. PMMA composites with non-functionalized and functionalized MXene were prepared using a solution casting method. Tensile tests showed that, compared to neat PMMA, Young's modulus increased in both composites by 22.1% and 27.6%, respectively. As a result of coupling between the PMMA matrix and the surface-modified MXenes, the tensile strength also increased by about 37%. In addition, optical spectroscopy showed higher absorption for the composite with surface-modified MXenes and short-lived fluorescence with emission intensity sensitive to the crumpling of functionalized MXene nanosheets.

Graphical abstract



Keywords: composites, mechanical properties, electron microscopy, X-ray photoelectron

spectroscopy, optical spectroscopy, MXenes

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1. Introduction

MXenes are a growing group of 2D materials with a unique combination of properties. MXenes can be synthesized by selective etching of the corresponding MAX phase (top-down) or by chemical vapor deposition (bottom-up). MAX phases are a large family of layered transition metal carbides or nitrides. They can be represented with the general formula $M_{n+1}AX_n$ ($n=1-4$), where M stands for the transition metal (usually an element from the 3rd to the 5th group of the PTE such as Ti, Zr, V, Nb, Ta, or Cr), A for the element from the 13th or the 14th group (the most common are Al and Si), and X for C or N [1-4]. MXenes have a similar general formula $M_{n+1}X_nT_z$, where M and X have the same meaning, and T represents functional groups attached to the MXene surface. It can be seen from the latter formula that there is a missing A atom which was selectively etched [5,6]. There are over 30 predicted MXene structures, but only some of them have been experimentally confirmed [7,8]. Based on their properties, MXenes can be used for different applications such as hydrogen storage, dye absorbents, ion exchange, and antibacterial agents. In addition, MXenes can be used as stable water (or some other) dispersions or as vacuum-filtered films.

Poly(methyl methacrylate) (PMMA) is an acrylate derivative and one of the most interesting industrial thermoplastic polymers. It is weather-resistant, biocompatible, chemically stable, and has good optical transparency from the ultraviolet to the near-infrared regions [9-12]. Polymeric waveguides are mostly made of this polymer because of these properties. Moreover, PMMA-based composites for a wide range of optical and optoelectronic applications can be obtained by embedding optically active media like quantum dots, carbon nanotubes, graphene, nanowires, or single crystals [13-15]. However, one of the drawbacks of this polymer is its brittleness. To enhance its properties, PMMA can be blended with other polymers, or nano- or micro-inorganic fillers can be added to it. The results depend on the type of interaction between the polymer chains and the filler [12, 16, 17].

Until now, there have been some theoretical descriptions of the mechanical properties of MXene single flakes [18]. In short, MXene single flakes have excellent mechanical properties, very similar to graphene sheets [19], which makes them a good candidate for a load-bearing role in composite materials. According to density functional theory, Young's modulus of the single flake $Ti_3C_2T_x$ MXene is about 0.33 TPa [20]. Since there are a lot of functional groups on the MXene flake surface, they can be modified with different kinds of materials such as nano cellulose, soybean phospholipid, metals (Pt, Ag, et cetera), or inorganic compounds (Fe_2O_3 , Mn_3O_4) [21-23] to improve or change their properties or just to prevent the oxidation. MXenes can also be incorporated into polymers to create composite materials with enhanced mechanical [24], electrical [25], or optical [26] properties, even with a small concentration of flakes. MXene surface plays a vital role here as well because of its functional groups that can interact with a polymer structure. The problem occurs when MXene functional groups and a polymer are not compatible, which may lead to MXene agglomeration and, consequently, to the deterioration of properties.

MXenes possess promising electron transport properties because of the high density of states at the Fermi level. Thus, they can be considered as materials for various optical applications, like light harvesting materials, and as functional mediums in polymer composites in optical, electromagnetic interference (EMI) shielding, or sensing applications [27-31]. The optical properties of MXenes are influenced by way of their surface termination. One of the possible sites

for surface termination is on top of transition metal atoms; the second is between top metal atoms, and the third is between stacking X-atoms layers [29, 31].

The research interest in MXene polymer composites increased over the past few years. It can be concluded that the processing method also influences the properties of polymer composites with MXene [32-36]. Highly transparent $Ti_3C_2T_x$ MXene film on flexible PET substrate was prepared by Ying et al. [34]. The processing by in situ polymerization was also performed with very good results, and the hybrid nanocomposite of PMMA with MXene/ZnO nanoparticles with improved dielectric properties was obtained [35]. Melt blending proved to be a suitable method when working with a thermoplastic polymer matrix, e.g., for preparing thermoplastic polyurethane (TPU) nanocomposites with homogeneously dispersed Ti_3C_2 MXene nanosheets [36]. All these methods were developed to achieve composites with high performances.

Various MXene modifications may lead to better compatibility between the flakes and the polymer matrix and suppress the aggregation of MXene nanosheets, as evidenced by the use of polyethylene glycol (PEG) [37], dopamine [38], or aniline [39]. There is also a well-described method of MXene functionalization with amino silane [40], which can serve as the basis when utilizing silanes.

The idea to incorporate MXenes in PMMA originated from our attempts to produce an optically-sensitive transparent thin film. There are only a few studies of PMMA-MXene nanocomposites [41], including our previous work [42] and the synthesis of specific MXene structures using PMMA as a template [43]. To our knowledge, there were no previous attempts to synthesize polymer composites with MXenes functionalized with silane. γ -Methacryloxypropyltrimethoxy (MEMO) silane was chosen for MXene functionalization because of its very similar structure to PMMA, which results in better compatibility and stronger interfacial bonding in the composite [40,41].

In our research, we analyzed the impact of functionalized MXenes on the optical and mechanical properties of the PMMA. Two types of composites were prepared: the first one, where the MXenes were directly added to PMMA, and the second one, where the MXenes were functionalized with MEMO silane before they were added to PMMA. The obtained composites were tested in order to determine Young's modulus and tensile strength and prove that the optical properties of MXenes are preserved when added to PMMA.

2. Experimental section

2.1. Materials

For the preparation of Ti_3C_2 MXene, concentrated HCl (Fisher Scientific UK) and LiF (325 mesh powder Alfa Aesar) were used. γ -Methacryloxypropyltrimethoxy (MEMO) silane (Dynasylan MEMO) was used to functionalize MXenes. DMF was purchased from Sigma Aldrich. Dimethyl sulfoxide (DMSO) was purchased from Fisher Scientific UK. PMMA pellets (Acryrex®, Chi Mei Corporation, Taiwan) were used to prepare the composites.

2.2. Preparation of delaminated Ti_3C_2 MXene

Ti_3C_2 MXene was prepared by etching the Ti_3AlC_2 MAX phase using LiF/HCl method. The powdered MAX phase was gradually added into a mixture of LiF and concentrated HCl. After 24 hours at $35^\circ C$, the dispersion was rinsed with DI water until pH 5, and the supernatant turned black. The centrifuge tube was shaken by hand for 1 min after adding water during the rinsing procedure. After the last washing step, sediment was again dispersed in DMF and sonicated for 1 hour under argon bubbling. Finally, the obtained dispersion was centrifuged for 1 hour at 3000 rpm, and the supernatant was collected.

2.3. Modification of MXene with MEMO silane

MEMO silane was first hydrolyzed in 50 ml of water/ethanol solution at room temperature for 1 hour, and the pH was adjusted with acetic acid [41].

MEMO silane functionalized MXene was prepared through a similar method as pure MXene. Ti_3C_2 MAX phase was etched using LiF/HCl method. The powdered MAX phase was gradually added into a mixture of LiF and concentrated HCl. After 24 hours at $35^\circ C$, the dispersion was rinsed with DI water until pH 5, and the supernatant turned black. The centrifuge tube was shaken by hand for 1 min after adding water during the rinsing procedure. The obtained sediment was dried for 1 hour at $70^\circ C$ and mixed with DMSO for 18 hours in a magnetic stirrer. After the mixing, sediment was separated from the excess DMSO by centrifugation. Afterward, it was redispersed in deionized water and sonicated for 1 hour under argon bubbling. Finally, the obtained dispersion was centrifuged for 1 hour at 3000 rpm, and the supernatant was collected.

The mass ratio of MEMO silane to MXene was 3:1. Hydrolyzed MEMO silane and MXene supernatant were mixed with a magnetic stirrer. The obtained mixture was centrifuged for 10 minutes at 4000 rpm to sediment unreacted silane. The supernatant was then vacuum filtered to get the thin film of functionalized MXene, and used in further synthesis.

2.4. Sample preparation

Three series of samples were prepared: pure PMMA, PMMA with unmodified MXenes (PMMA-MX), and PMMA with MEMO silane-modified MXenes (PMMA-MXS).

Series 1 - PMMA:

A pure PMMA matrix was prepared by dissolving the PMMA pellets in dimethylformamide (DMF) for 24 hours in a magnetic stirrer. The solution was then poured into Teflon molds and left partially covered in the dryer for another 24 hours at $65^\circ C$.

Series 2 - PMMA-MX:

The supernatant was used as a solvent for the series. PMMA pellets were put into the supernatant and left in the magnetic stirrer for 24 hours. The concentration of MXenes in the composite was 1 wt%. The solution was poured into Teflon molds and dried for 24 hours at $50^\circ C$, followed by 24 hours in a vacuum drier.

Series 3 - PMMA-MXS:

PMMA was dissolved in DMF for 24 hours in a magnetic stirrer. After that, the previously

made thin film was measured, chipped, and added to the solution to obtain the concentration of 1 wt% of the functionalized MXene in PMMA. Next, the solution was treated in an ultrasonic bath for several minutes to break down the chips, followed by stirring for another 24 hours. The solution was then poured into Teflon molds and left in the oven at 65 °C overnight.

The schematic representation of the experiment is shown in Figure 1.

2.5. Material characterization

FTIR spectra of functionalized MXene and composites were collected through single-beam Fourier-Transfer Infrared Spectroscopy (FTIR) using a Nicolet 6700 spectrometer.

SEM images were obtained by using FESEM Tescan MIRA 3 XMU electron microscope.

Tensile testing was performed at room temperature with the Shimadzu EZ-LX universal testing instrument equipped with a 5 kN load cell. The machine resolution was ± 0.5 % of the indicated value (within 1/500 to 1/1 of load cell rated capacity), and crosshead position detection accuracy was 0.1%. The samples (12 mm wide, 60 mm long, and 0.5 mm thick) were tested at a 3 mm/min speed. All the samples were conditioned at room temperature and in a desiccator for 48 hours before measurements. Three identical samples were tested for each series.

XPS analysis of the samples was carried out on the SPECS Systems with XP50M X-ray source for Focus 500 and PHOIBOS 100 energy analyzer using a monochromatic Al K α X-ray source (1486.74 eV) at 12.5 kV and 12 mA. All samples were fixed onto an adhesive copper foil to provide strong mechanical attachment and good electrical contact. All survey XPS spectra (0–1300 eV BE) were recorded with a constant pass energy of 40 eV, energy step of 0.5 eV, and dwell time of 0.2 s, while high-resolution XPS spectra of the corresponding lines were taken with a pass energy of 20 eV, energy step of 0.1 eV and a dwell time of 2 s.

The optical characteristics of MXenes were measured using a time-resolved laser-induced optical spectroscopy system. It is based on Nd-YAG Vibrant OPO (Optical Parametric Oscillator) laser as an excitation source. The OPO output is continuously tunable in the range between 320 nm and 475 nm. The fourth harmonic (266 nm) and the second harmonic (532 nm) outputs are also available. The time duration of the laser pulse is about 5 ns, and its repetition rate is 10 Hz. The samples' time-resolved emission spectra were obtained by using the Hamamatsu streak camera system equipped with the spectrograph. Absorbance spectra were obtained using the Ocean Optics HR2000 spectrometer. The samples were excited by a 12 V, 55 W halogen light bulb.

3. Results and discussion

The Fourier transformation infrared spectroscopy (FTIR) was used to confirm the functionalization of MXene with MEMO silane (Figure 2). The characteristic broad peak around 3500 cm⁻¹, which corresponds to hydroxyl groups or entrapped water, can be seen in the spectrum of unmodified MXene. The disappearance of this peak in the spectrum of modified MXene can be attributed to the termination of OH groups by silanol groups of MEMO silane [40, 44].

FTIR measurements for the composite samples are shown in Figure 3. Characteristic peaks

for PMMA occur at 1718 cm^{-1} ν (C = O) and 1430 cm^{-1} ν (C - O). The bands at 2953 cm^{-1} correspond to the C-H elongation of the methyl group (CH₃). The peak at 1136 cm^{-1} corresponds to the vibrations of the ester group C - O [44]. The peak at 980 cm^{-1} corresponds to the ester bonds, and peaks at 751 and 846 cm^{-1} represent (C - C) stretching [45]. Spectra of series with MXene have a new peak that corresponds to Ti-O stretching vibration and shifts to 552 (596) cm^{-1} [46]. Also, a new broad feature centered at $\sim 3100\text{ cm}^{-1}$ is present in these spectra and can be attributed to the Ti-OH bond [47].

Scanning electron microscopy (SEM) was employed to examine the structure of the particles used in the experiment. Both MXenes and MEMO silane functionalized MXenes were vacuum filtered to obtain the samples for the microscope. Results are shown in Figure 4.

At first glance, a compact structure is observed in both cases. It is assumed that nanosheets were stacked during vacuum filtration. On the other hand, as shown in Figure 4b, which represents functionalized MXenes, the structure is looser and less compact. Since the pH of hydrolyzed MEMO silane used in synthesis was 3, this caused the 'crumpling' of MXene flakes, as reported in [48]. In addition, after functionalization, the silane molecules bonded to the flakes' surface in the perpendicular direction, which also leads to an enlarged spacing between MXene layers [48,49] and is, along with the crumpling, represented in Figure 4.

In order to confirm our assumptions about the improved compatibility between functionalized MXenes and PMMA, we analyzed the SEM images of both types of composites, which are presented in Figures 4c and 4d, respectively. Figure 4c represents the cross-section of the PMMA-MX sample. Clusters of MXene nanoparticles can be observed due to the rigorous mixing of the polymer solution. On the other hand, no noticeable agglomeration can be observed in the cross-section of the PMMA-MXS sample (Figure 4d). That is, MXene nanoparticles are more evenly distributed.

The representative curves from the tensile test of all three series are shown in Figure 5 and summarized in Table 1.

From the shapes of tensile curves, it can be seen that Young's modulus, calculated as the slope of the linear section of the stress-strain curve in the 0.5 - 2.5 MPa stress range, increases for both PMMA-MX and PMMA-MXS. The strain at break, however, is lower for PMMA-MX. This can be explained by the incompatibility between the MXene surface and PMMA, which leads to MXene flakes serving as stress concentrators and causing micro ruptures during deformation. Better bonding between MEMO silane functionalized MXene and PMMA is achieved in the PMMA-MXS composite, which results in improved load transfer from the matrix to reinforcements. As a result, the strain at break and tensile strength increase for PMMA-MXS, and consequently, its toughness improves.

The average values of mechanical properties determined by tensile tests, along with corresponding standard deviations (s.d.), are shown in Table 1. Compared to the results of some other MXene-polymer nanocomposites and PMMA nanocomposites with other nanoparticles, our materials follow the trend of the increased values of Young's modulus and tensile strength as obtained in [50-52]. Looking at the relative improvement of Young's modulus, in this work, an increase of 22.1% for PMMA-MX and 27.6% for PMMA-MXS were achieved, which is in accordance with other studies, e.g., an increase of 38.9% in [50] and 33% in [52]. The increase in tensile strength was 9% for PMMA-MX and almost 40% for PMMA-MXS. Compared with other

MXene-polymer composites with a similar wt% of filler content, thermoplastic polyurethane (PTU) with 1 wt% has a 25% increase in tensile strength [53], while for 0.5 wt% of MXenes in Waterborne polyurethane and Ultrahigh molecular weight polyethylene the increase was 25% and 65%, respectively [53]. For nanocomposites with 1 wt% of graphene oxide in PTU and 5.4 wt% in PVA, the increase of tensile strength was 40% and 87%, respectively [54, 55]. In addition, the theoretical prediction of the modulus of elasticity can be calculated using Halpin–Tsai's micromechanical model leading to the value of 2152 MPa, which is comparable to the values that we observed for PMMA-MX and PMMA-MXS samples.

In order to express the reliability of the results of mechanical testing, standard deviation and measurement uncertainty were calculated. Measurement accuracy analysis for tensile strength (TS) was performed with the method defined by Klysz [55, 56]. Based on machine accuracy, the standard uncertainty for TS was calculated as $u(\text{TS})=0.18$ MPa [57]. For strain, the standard uncertainty was calculated as $u(\text{Str})=0.21\%$ [57]. Uncertainty of measurements is a parameter for error probability distribution in the context of instrument accuracy's influence. The uncertainty of measurements is lower than the standard deviation, as can be seen from Table 1, so the instrument's influence on the error can be considered as low. Actually, both the standard deviation and uncertainty of the measurements are considerably influenced by the number of samples in a testing procedure [57].

Survey spectra of MXene and functionalized MXene with MEMO silane are presented in Figures 6a and 6b, respectively. The main photoelectron lines of Ti, O, and C constituting elements are clearly observed and marked together with their Auger lines. The photoelectron lines detected at around 685 eV, 270 eV, and 200 eV belong to F1s, Cl2s, and Cl2p, respectively, which originate from the preparation procedure of Ti_3C_2 MXene using the LiF/HCl method. It was found that after functionalization of MXene with MEMO silane (Figure 7b), the additional lines that correspond to silicon (Si 2s and Si 2p lines, respectively), which are marked with arrows, were observed at about 154 eV and 102 eV. The region of Si 2p recorded in better resolution is presented in Figure 6c, confirming the successful functionalization of MXene with MEMO silane.

Absorbance spectra of PMMA, PMMA-MX, and PMMA-MXS are shown in Figure 7. Due to the low signal levels in UV and IR regions, corresponding parts of measured spectra are very noisy. Therefore, the truncated wavelength range is presented in Figure 7 of spectra that were measured originally in the range between 300 nm and 1100 nm. Absorbance spectra of MXene samples have similar shapes as absorbance spectra obtained by Maleski et al.[58], where the spectrum of MXene on the sapphire surface was presented. In the 300-400 nm range, the spectrum is with a lot of noise because of the dispersion of MXenes in PMMA and some signal attenuation. It can be seen that functionalized MXene in PMMA shows higher absorbance in the 800 nm region, similarly to [58], while for pure MXene, absorbance in this region is lower. The absorbance of PMMA is almost flat in the entire region. The absorption peak at 800 nm observed in PMMA-MXS is ascribed to the inherent out-plane interband transitions [58].

To show the changes in the optical appearance of the composites, we photographed them on a piece of paper, along with neat PMMA, so they could be observed together and compared. The resulting image is shown in Figure 8.

Emission intensity (integrated-in-time) of time-resolved optical spectra of PMMA, MXene in PMMA, and MEMO silane functionalized MXene in PMMA using the 266 nm excitation are

shown in Figure 9. The very strong peak at 532 nm actually corresponds to spectrometer diffraction grating second-order “ghost” of laser excitation at 266 nm. Because of their relatively small absorption, as MXene materials have moderate quantum yield (8.9% in Desai et al.[59]), they are also characterized by small fluorescence intensity. As shown in [39], MXene materials have wideband fluorescence emission in the 400-600 nm range. Regardless, the emission peak of PMMA-MX is the highest and slightly red-shifted to higher wavelengths. However, the peak of PMMA-MXS again blue shifts and appears at nearly the same wavelength as in the spectrum of neat PMMA. This influence of silane modification was already observed in our earlier work [42] and can be attributed to the silane modification influencing a better spatial organization of modified MXenes in the polymer matrix. The MXene fluorescence spectra have shapes very similar to spectral shapes presented in Mittal et al.[59], where MXene nanosheets were used for selective and sensitive fluorescence detection of Ag^+ and Mn^{2+} ions. In this research, the disturbance of MXene nanosheets with the embedding of these ions also occurred. As shown in Figure 10, which represents streak images of time-resolved spectra of PMMA, PMMA-MX, and PMMA-MXS, MXene fluorescence is very short-lived. It lasts only as long as the laser pulse lasts. Therefore, with the laser excitation duration of 5 ns and the repetition rate of 10 Hz, we obtained relatively small intensities of MXene fluorescence compared to [59], where continuous excitation was used.

Presented results suggest that the fluorescence emission is sensitive to the surface modification of MXene and the crumpling of MXene nanosheets. This results in fluorescence quenching, that is, the decreasing of the fluorescence intensity in PMMA-MXS that can be associated with the MXene-silane bonding. The broad absorption properties of MXenes in the 400-1200 nm range and their intrinsic fluorescence quenching ability make them a promising material as a quencher in fluorescence resonance energy transfer (FRET) [60, 61]. Thus, our PMMA-MXS thin film can potentially be used as a biosensing element in fluorometric assays.

4. Conclusion

A new type of MXene-based PMMA composites with improved toughness was developed via a simple solution casting method. To improve the bonding between the reinforcement and the polymer matrix, MXenes were functionalized with MEMO silane. Composites with unmodified and functionalized MXenes were processed, and structural, optical, and mechanical properties were investigated and analyzed in order to pave the way for understanding the significance of MEMO silane addition to the MXenes. Functionalization was confirmed with the comparative FTIR analysis of neat and functionalized MXenes and the XPS survey spectra. SEM images and tensile tests confirmed improved compatibility of functionalized MXenes with PMMA, as higher tensile strength, strain at break, and modulus of elasticity were obtained in PMMA-MXS. Optical characterization of composites revealed that functionalized MXenes in PMMA showed higher absorption in some parts of the visible spectrum. The silane modification influences a better spatial organization of modified MXenes in the PMMA matrix, resulting in better fluorescence quenching and leading to the potential use of PMMA-MXS thin films as enzyme reaction sensors. This research helps to disclose some of the properties of this type of nanocomposites that are novel and are yet to be fully understood.

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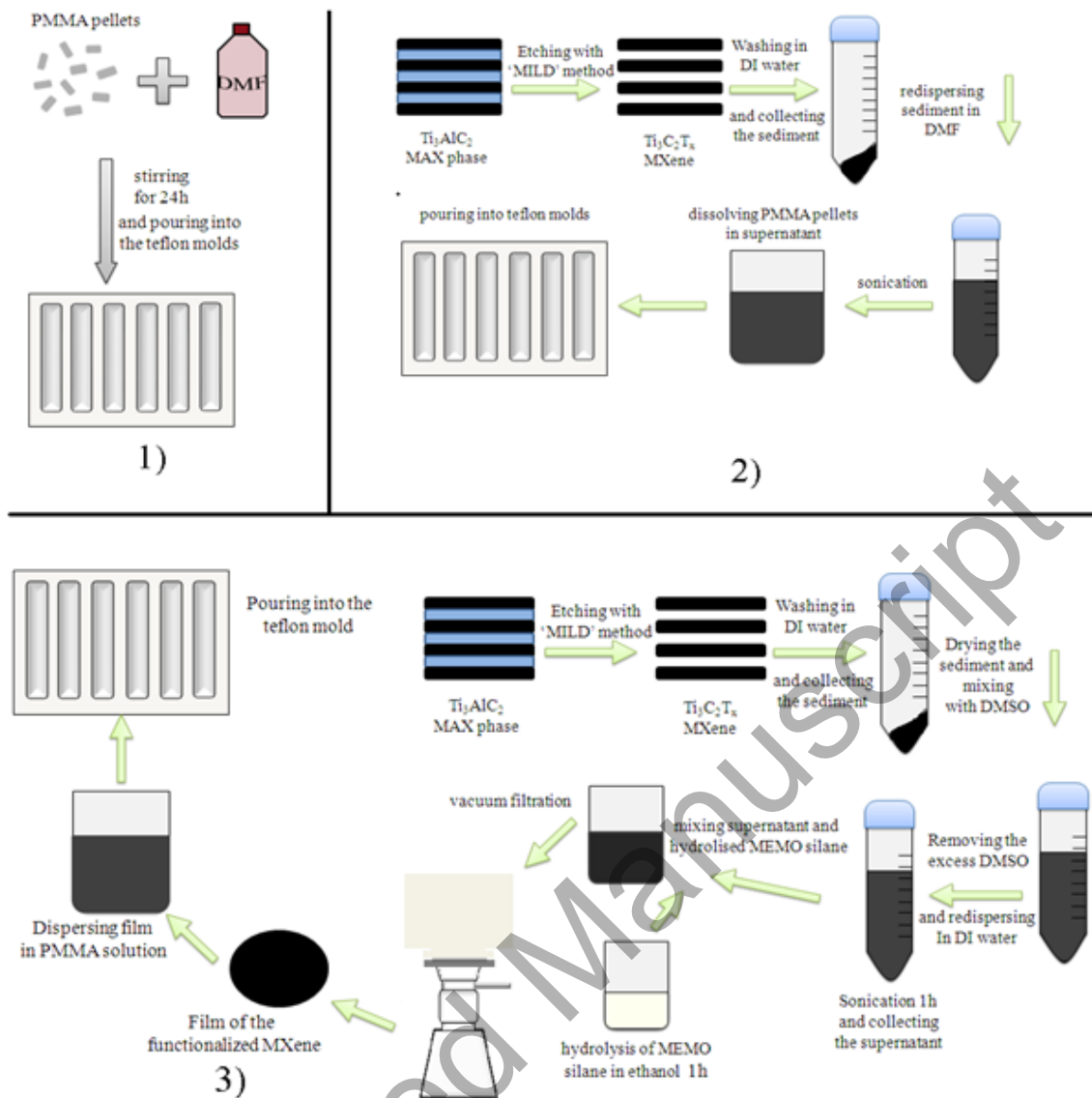


Figure 1. Schematic representation of the experiment: 1) preparation of PMMA, 2) preparation of unmodified MXene in PMMA (PMMA-MX), and 3) preparation of MEMO silane-modified MXene in PMMA (PMMA-MXS).

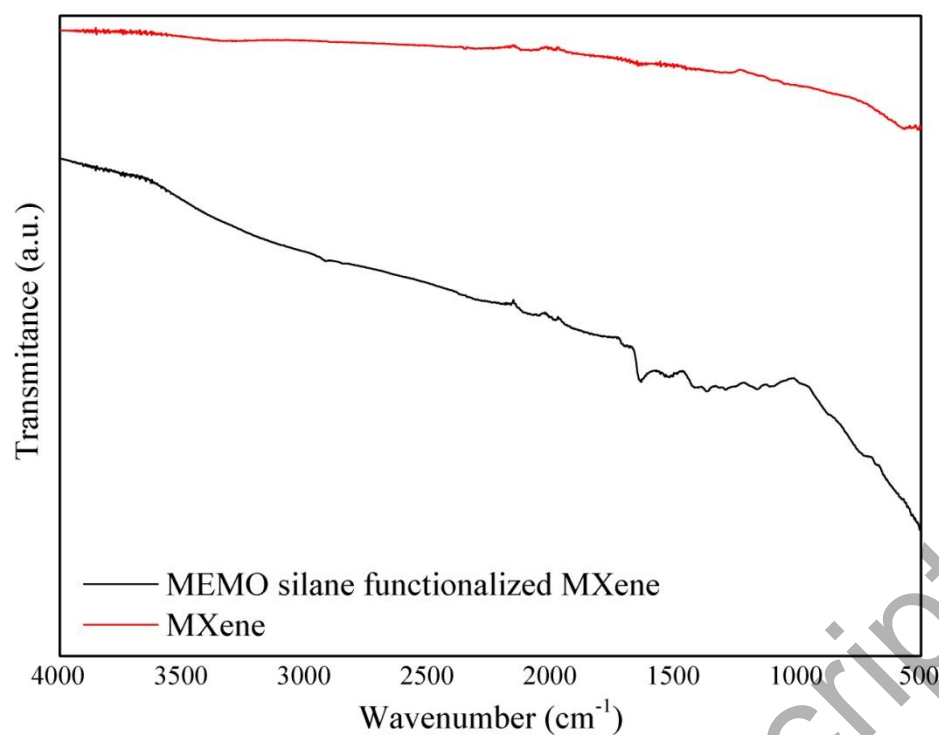


Figure 2. FTIR spectra of MEMO silane functionalized MXene and pure MXene.

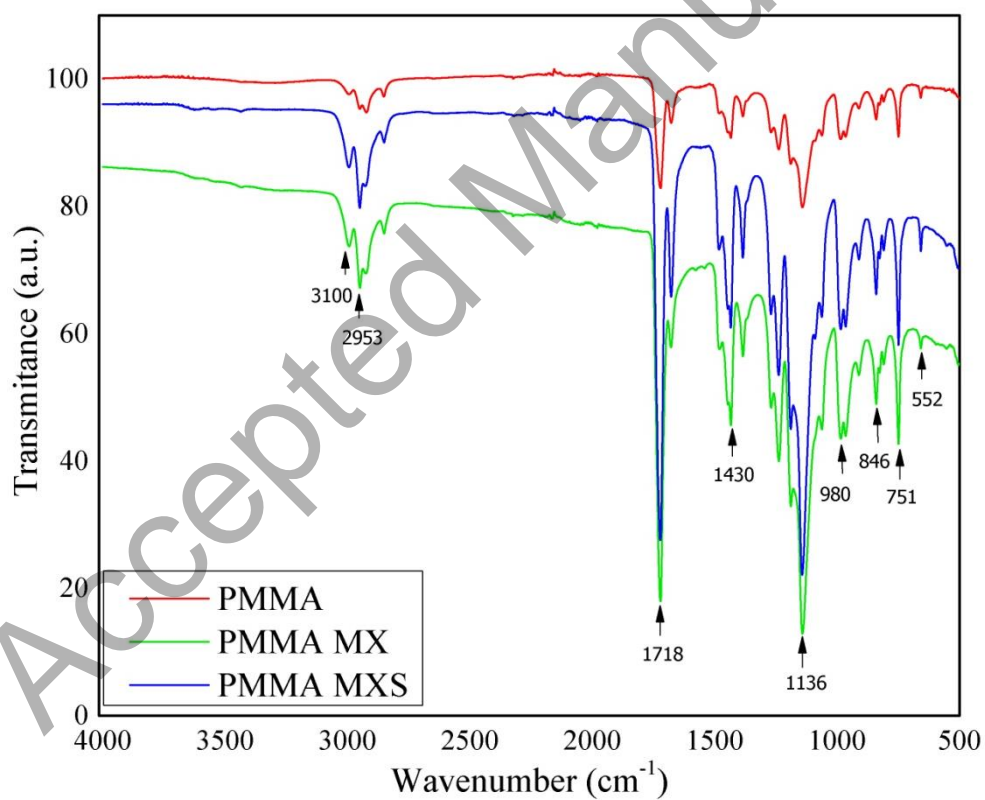


Figure 3. FTIR spectra of PMMA, PMMA-MX and PMMA-MXS.

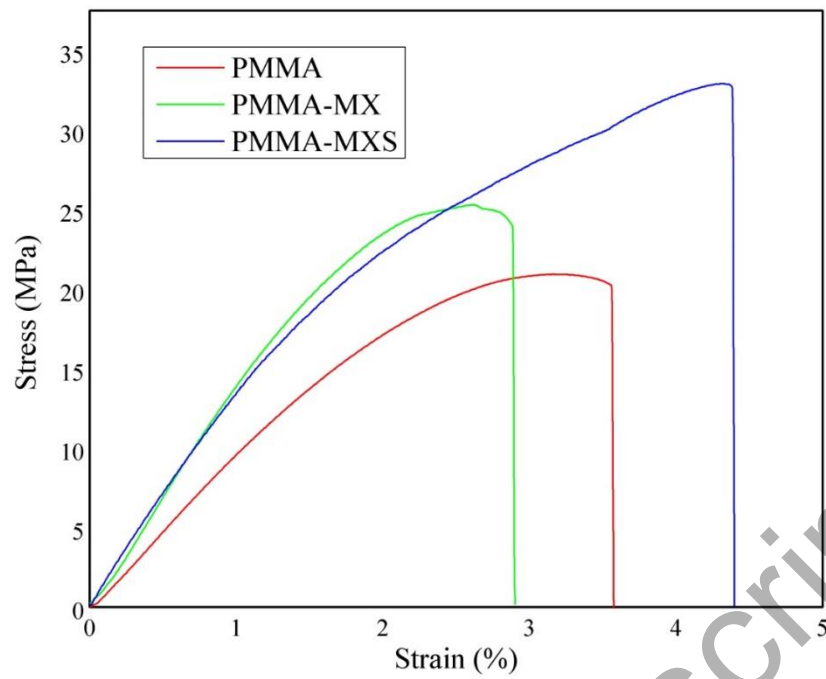


Figure 5. Stress vs. strain diagram for the PMMA, PMMA-MX, and PMMA-MXS.

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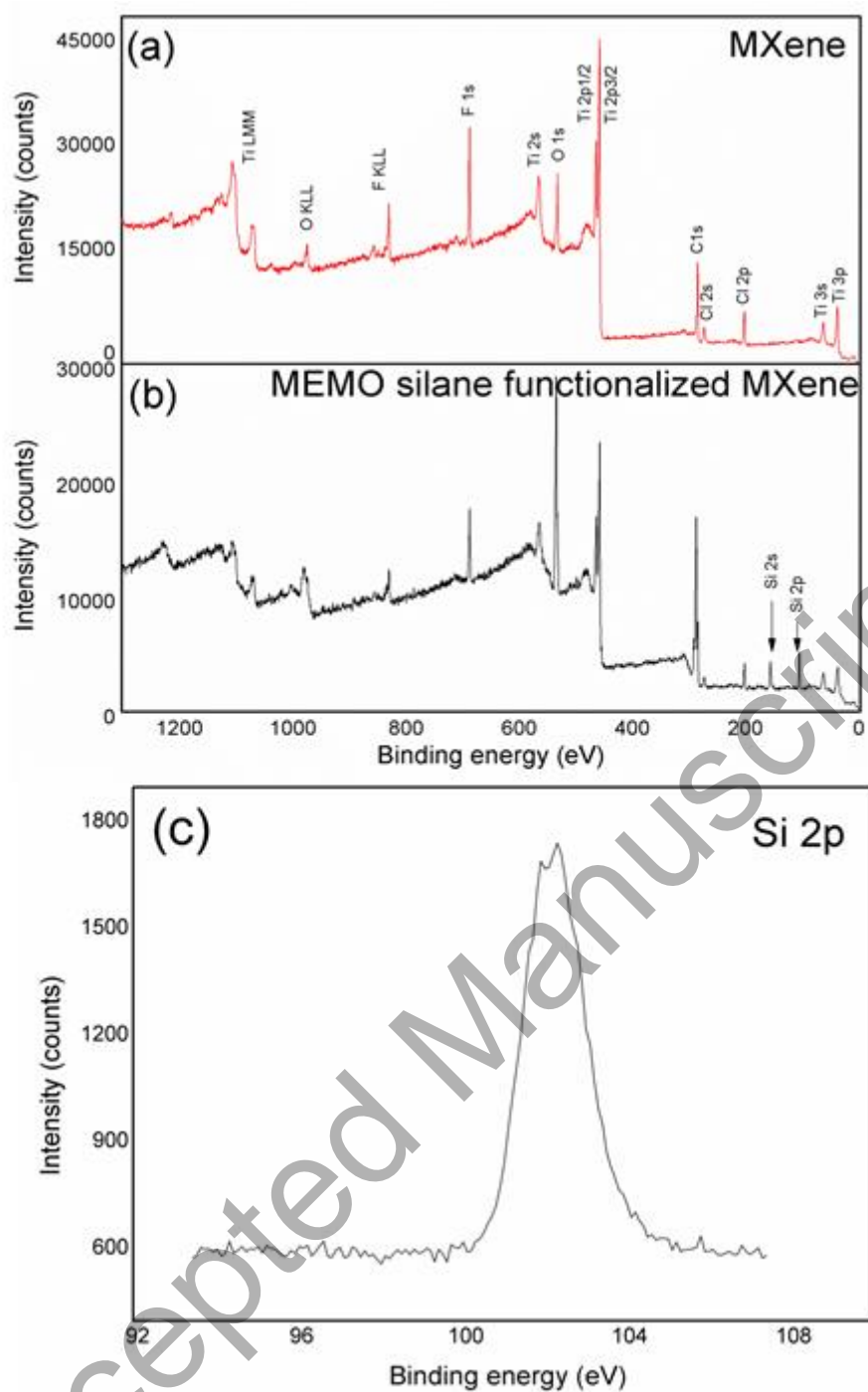


Figure 6. XPS survey spectra of a) MXene and b) MEMO silane functionalized MXene, and c) high-resolution XPS spectrum of Si 2p line of MEMO silane functionalized MXene.

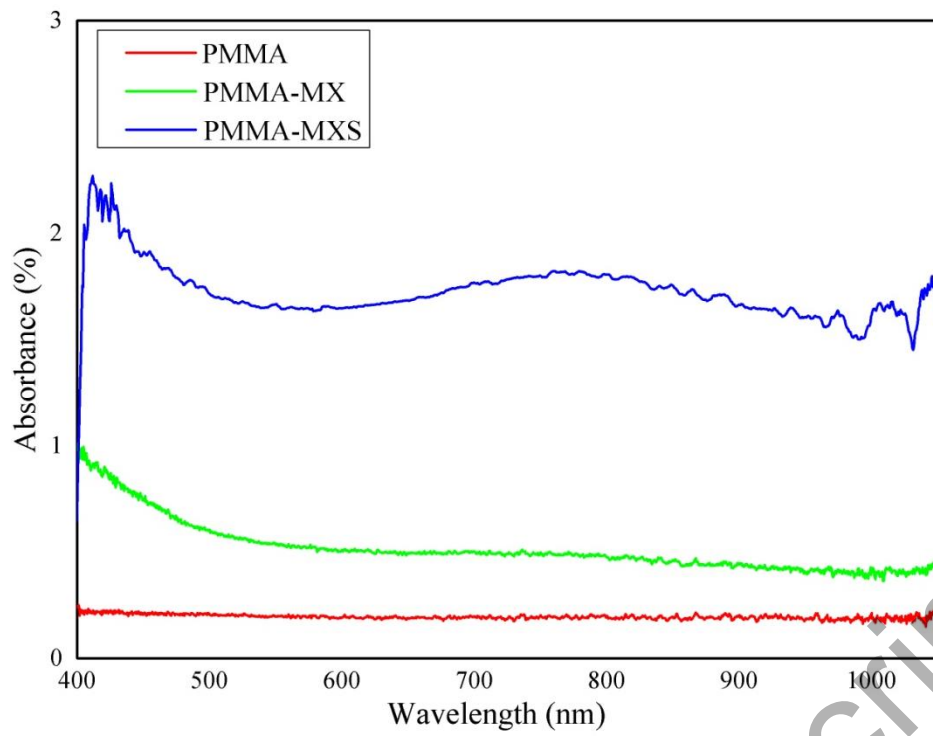


Figure 7. Absorbance spectra of PMMA, PMMA-MX, and PMMA-MXS.



Figure 8. The optical transparency of PMMA, PMMA-MX and PMMA-MXS samples.

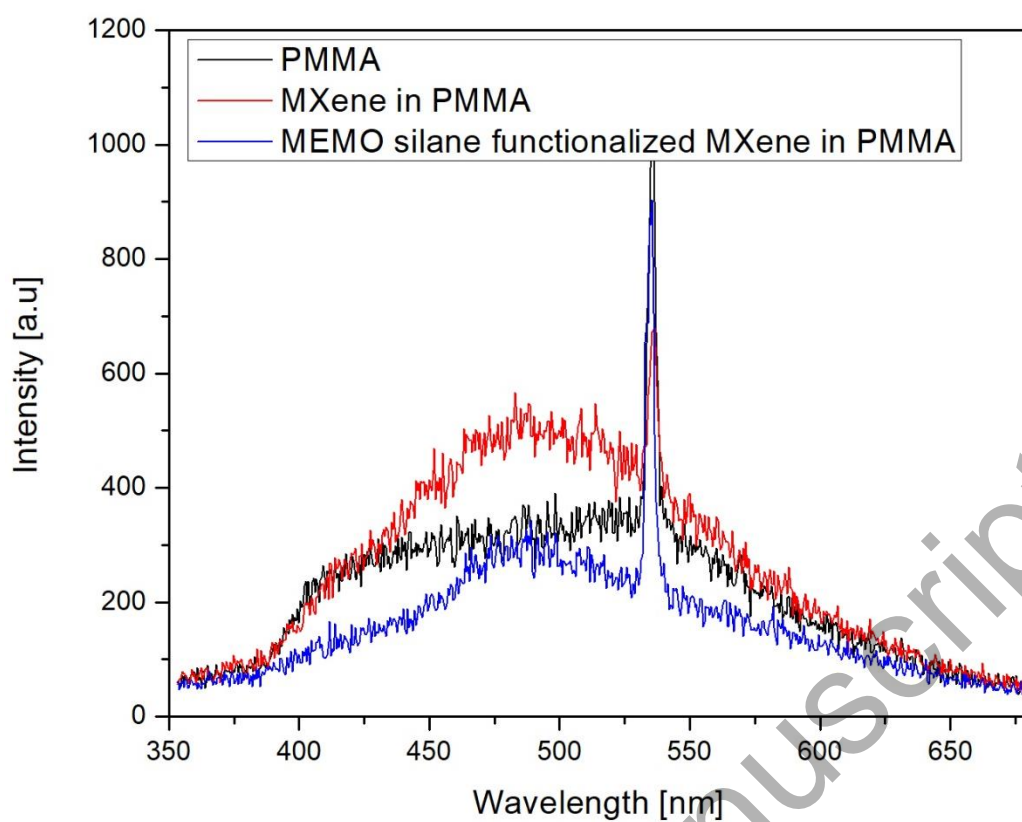


Figure 9. Emission intensity (integrated-in-time) of time-resolved optical spectra of PMMA, PMMA-MX, and PMMA-MXS.

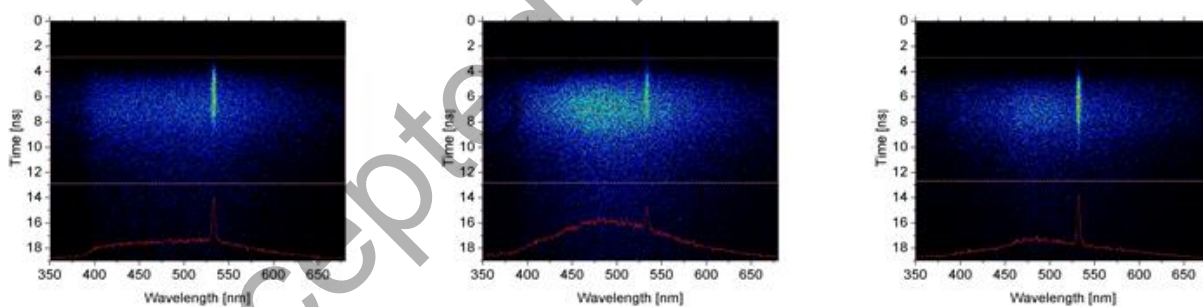


Figure 10. Streak images of time-resolved optical spectra of a) PMMA, b) PMMA-MX, and c) PMMA-MXS.

Table 1. Average values of Young's modulus, tensile strength and strain at break, with corresponding standard deviations (SD).

Series	Young's modulus, MPa	SD, MPa	Tensile Strength, MPa	SD, MPa	Strain at break, %	SD, %
PMMA	1015.22	182.29	21.98	1.48	3.63	0.67
PMMA-MX	1235.81	192.21	23.96	1.88	2.33	0.41
PMMA-MXS	1291.75	164.68	30.05	4.06	3.96	0.50

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