**Introduc**tion

The energy crisis, environmental pollution, climate change, and global warming are some of the crucial environmental problems in the modern world due to the emission of various hazardous byproducts into the environment. The combustion in the automobile engine produces carbon dioxide (CO₂) and carbon monoxide (CO) gases causing hazardous health problems to humans and, it is one of the major causes of air pollution. Many researchers are working on developing new techniques to produce clean energy which is challenging. Also, providing a realistic solution for the current energy demand is much needed. The use of hydrogen gas (H₂) as a clean energy source is one of the modern concepts as H₂ is clean as well as energy-efficient. One of the best ways to produce clean and eco-friendly hydrogen is through a process called photocatalytic water electrolysis or also known as an artificial photosynthesis.¹

In this process, sunlight is utilized in the presence of photocatalyst to decompose water into H₂ at the cathode and oxygen (O₂) at the anode. It was firstly reported by Fujishima and Honda in 1972, however, H₂ gas generating via green processes and high efficiency are still challenging.²

Water electrolysis through proton exchange membrane (PEM) is one of those green methods which can produce H₂ and is utilized in a wide range of applications. Production of H₂ occurs by the redox reaction that takes place in an electrolyzer cell. PEM water electrolysis is accrued by pumping of water to the anode side where water is splitted into oxygen (O₂), protons (H⁺) and electrons (e⁻). These protons transport via a proton conducting membrane to the cathode side. The PEM allows only the proton (H⁺) produced at the anode to transfer towards the cathode to produce H₂. Nafion, a conventional commercialized PEM, had been used extensively for the PEM water electrolysis, which is expensive with complicated synthesis steps and toxic to the environment due to the presence of fluorine. Considering the cost-effectiveness, and mechanical and chemical properties, polyether ether ketone (PEEK) is an alternative hydrocarbon-based polymer that can be used for the PEM water electrolysis by the sulfonation of polyether ether ketone. Cross-linking of the sulfonated polyether ether ketone (SPEEK) membrane using ethylene glycol (EG) as a crosslinker increased chemical and...
mechanical stabilities and decreased swelling ratio which made it suitable for the application as PEM. The proton conductivity of SPEEK would be reduced when diols are used as a cross-linker, therefore many researchers have incorporated hygroscopic inorganic metal oxide such as SiO₂, TiO₂, Al₂O₃, and ZrO₂ as fillers to improve physiochemical properties and conductivity of the membrane.

TiO₂ has useful properties among the above-mentioned metal oxides that can be used as a filler material. It is capable of becoming super hydrophilic when exposed to UV light which arises due to the increasing number of hydroxyl groups (–OH) of the TiO₂ surfaces during UV light irradiation. TiO₂ is a nontoxic, cost-effective, abundant material and possesses high chemical stability against the oxidative environment in photocatalytic water splitting. TiO₂ particles can be converted to nanowire via a hydrothermal reaction using a strong basic solution such as NaOH or KOH solution. Potassium titanate nanowire is one of the fascinating forms of TiO₂ that could be utilized as a reinforcing agent in it possesses good mechanical, chemical and thermal stability. Additionally, it was also used as a photocatalyst in water electrolysis. TiO₂/graphene nanocomposites have been studied as a filler in PEMs to enhance numerous properties of the membrane.

MXenes are a new versatile inorganic material of 2D transition metal carbides or nitrides, which were produced by selective chemical etching of MAX phases (M is an early transition metal, A is an A-group (mostly HIA and IVA element) and X is carbon and/or nitrogen). Mostly Ti₃C₂Tx layers are uniformly terminated by –OH and –O– and –F groups after the exfoliation, where T and x represent the terminating group and their numbers, respectively. These surface termination of MXene make them a prominent filler to increase the proton conductivity and physicochemical properties of SPEEK. MXene has been utilized as an electrocatalyst for H₂ evolution reaction (HER) in water splitting. When MXene was employed as a co-catalyst with TiO₂ in photocatalysis, higher H₂ evolution has been observed without the addition of any noble metal co-catalyst. Besides, the addition of co-catalyst with TiO₂ such as copper (Cu) could obtain a highly active H₂ production catalyst, Cu/TiO₂/MXene photocatalyst in which MXene acted as a hole mediator and Cu acted as an electron mediator and reduction co-catalyst. The coupled photocatalyst promoted H₂ evolution via dual-carrier-separation mechanism.

Thus, the present work focuses on the preparation of hybrid PEM, by utilizing sulfonated poly(ether ether ketone) (SPEEK) as a polymer matrix for the PEM water electrolysis. SPEEK with a high degree of sulfonation (DS) had been prepared via sulfonation and cross-linked with ethylene glycol (EG) to improve its dimensional stability. To further increase its proton conductivity, inorganic metal oxide; MXene-potassium titanate nanowire (MKT-NW) was embedded into the polymer matrix to prepare MKT-NW/C-SPEEK hybrid PEM. The PEM was prepared by a simple solution casting technique using ethanol and water (1:1) as a green solvent. The properties of hybrid composite PEM including photocatalytic activity under UV radiation were characterized.

### Experimental

#### Materials and chemicals

Polyether ketone (PEEK) powder (particle size ~ 80 μm) and lithium fluoride (LiF) were purchased from Sigma Aldrich. Absolute ethanol (99.9%), concentrated sulfuric acid (H₂SO₄, 98%), potassium hydroxide (KOH) pellets and concentrated hydrochloric acid (HCl, 37%) were obtained from QREC. Sodium hydroxide (NaOH) pellets and sodium chloride were bought from Merck. Titanium aluminum carbide (Ti₃AlC₂ or MAX phase) was supplied from Luoyang Tongrun Info Technology Co., Ltd., China, and Nafion-212 was bought from Wuhan Gaoshi Ruiilian Technology Co., Ltd., China. Titanium dioxide (anatase, particle size – 120 nm) was purchased from Ajax Fine Chemicals. All the chemicals were used without purification. Deionized (DI) water (conductivity = 0.067 μS cm⁻¹) was used throughout this work.

#### Preparation of hybrid composite membranes and characterizations

A hybrid composite membrane consisted of sulfonated PEEK (SPEEK) as a polymer matrix, ethylene glycol as a crosslinker with different amount of MXene–potassium titanate nanowire (MKT-NW) as an active filler. SPEEK was prepared via sulfonation of PEEK with conc. H₂SO₄ at 50 °C following the literature to obtain orange solid particle of SPEEK. The degree of sulfonation was characterized by 1H-NMR (Bruker AVANCE III HD 600 MHz NMR spectrometer).

For MKT-NW composite, an exfoliated MXene was firstly synthesized via an in situ production of hydrofluoric acid (HF) protocol using LiF and 37% HCl as an etching agent where the exfoliation was carried out at 50 °C for 48 h. Potassium titanate nanowire (KT-NW) was prepared by a hydrothermal treatment of TiO₂ anatase with 10 M KOH solution in a Teflon-lined stainless autoclave at 200 °C for 24 h as reported previously. MKT-NW potassium titanate nanowire (MKT-NW) was obtained by the hydrothermal reaction of MXene and KT-NW (weight ratio 1:1) in DI water/ethanol mixture (3:1) using a Teflon-lined autoclave at 120 °C for 3 h. The structure of MXene and MKT-NW was characterized by wide-angle X-ray diffractometry (XRD, Bruker, D8 ADVANCE) with Cu-Kz radiation (λ = 1.54 Å) in the range from 5° to 80 °C at the rate of 0.02°/0.5 s.

X-ray photoelectron spectroscopy (XPS) was recorded on JEOI JPS-9010MC with a Mg Kα source (1253.6 eV) at 12 kV and 25 mA. All XPS spectra were measured under a high vacuum pressure of 10–8 Pa at a room temperature. Software for running experiment is SpecSurf ver.1.9.3. The sample was fixed on carbon tape. The analyzed area was a cycle spot with a diameter of 6 mm. The survey scan spectra were measured with a pass energy of 50 eV, a binding energy range of 0–1100 eV and an electron-volt step of 1 eV. The narrow scan spectra were measured with a pass energy of 10 eV and an electron-volt step of 0.1 eV. The obtained spectra were evaluated using JEOI software to obtain the chemical state of the probing elements and elemental composition. All the binding energy values were referenced to the carbon peak C 1s at 284.70 eV.
In a typical procedure of the composite membrane preparation, SPEEK particles (5 wt%) were dissolved in ethanol/DI water mixture (1:1), then the amount of ethylene glycol was added as a cross-linker and stirred vigorously for 2 h. The MKT-NW suspension (MKT-NW in ethanol/water mixture) was added to the SPEEK solution, then the final mixture was sonicated and stirred vigorously for 24 h, to make the uniform dispersion. The solution was cast on a Petri dish, dried at room temperature, and cured at 120 °C for 24 h to obtain a cross-linking membrane. After that, it was cooled down to room temperature and the membrane was peeled off using DI water and protonated by the immersion in 1 M H₂SO₄ for 24 h. Then, the membrane was washed with DI water until neutral and dried at 60 °C for 12 h. The composite membranes were named as MKT-NW/C-SPEEK where x is the amount of MKT-NW (5 to 20 wt%). Pristine SPEEK membrane, as well as cross-linked SPEEK (C-SPEEK) membrane, were also prepared and used. The overall steps involved in the preparation of the composite PEM is described in Scheme 1.

The chemical structure of the membrane was characterized by attenuated total reflectance-Fourier transform infrared (ATR-FTIR) spectroscopy (PerkinElmer Spectrum 100 FT-IR spectrometer) in the wavenumber range of 4000–500 cm⁻¹. The morphologies of the synthesized inorganic powders and composite membranes were characterized by a field emission scanning electron microscope (FE-SEM, model Versa) with energy dispersive X-ray spectroscopy (EDS) mapping. Transmission electron microscope (TEM, JEOL, JEM-2100Plus) was carried out to study the structural change of pristine MXene and after deposition with KT-NW. The elemental distribution in the MKT-NW/C-SPEEK composite membrane was carried out using X-ray fluorescence microscopy (XRF, Horiba, XGT-7200). Dynamic mechanical analysis (DMA) of the membranes were carried out using Mettler Toledo DMA1 analyzer with a TCD detector and a 5A molecular sieve packed column using argon as carrier gas. Hybrid composite PEM was immersed in a 10% v/v aqueous methanol solution (120 ml) as a sacrificial reagent for hydrogen production. Before starting the experiment, the sacrificial reagent in the reaction system was purged with argon in the dark environment to remove oxygen completely. UV lamp was set closer to the membrane by exposing only 30% of it with the intensity of 27.1545 W cm⁻². Later, the light source was moved further from the membrane to expose 100% of it and the light intensity was recorded as 0.0375 W cm⁻². The workstation set up for testing the photocatalytic activity of MKT-NW/C-SPEEK at room temperature is depicted in Scheme 2.

Results and discussion

Structural and material characteristics

SPEEK formed by the electrophilic substitution reaction of PEEK with concentrated sulfuric acid.²⁴ From Fig. 1(a), the degree of the sulfonation of SPEEK was 73.10%. FTIR spectra showed...
characteristic peaks of sulfonyl group (\(-\text{SO}_3\text{H}\)) at 1250 cm\(^{-1}\) (O=S asym), 1075 cm\(^{-1}\) (O=S sym), 1010 cm\(^{-1}\) (S=O), and 707 cm\(^{-1}\) (S=O), which were introduced into the hydroquinone segment of the PEEK polymer chain (Fig. S1†).

From the FTIR spectra of all prepared MKT-NW/C-SPEEK composite membranes as depicted in Fig. 1(b), the hydroxyl group (\(-\text{OH}\)) at 3394 cm\(^{-1}\) had been shifted to lower wavenumber for C-SPEEK sample due to cross-linking between \(-\text{SO}_3\text{H}\) of SPEEK and \(-\text{OH}\) of ethylene glycol, the crosslinker.\(^{19}\) Besides, the characteristic peak of \(-\text{SO}_3\text{H}\) remained unchanged and no new peak observed which implied that there was no chemical reaction and only interfacial interaction between C-SPEEK and MKT-NW.\(^{11}\) However, the decrease in the peak intensity of \(-\text{SO}_3\text{H}\) was observed which may be due to physical interaction of MKT-NW composite with \(-\text{SO}_3\text{H}\) of SPEEK. For the incorporation of MKT-NW in C-SPEEK, the \(-\text{OH}\) functional group intensity slightly decreased which may be attributed to the extensive hydrogen bonding interaction between \(-\text{OH}\) terminated MKT-NW surface and \(-\text{SO}_3\text{H}\) of C-SPEEK.\(^{20}\)

In this work, KT-NW led to the formation of TiO\(_6\) octahedral by a zigzag pattern by connecting the edges and the potassium ion (K\(^+\)) are trapped into this structure. In the KT NW synthesis process, longer Ti\(-\text{O}\) bonds might be attacked by \(-\text{OH}\) ions of strong aqueous KOH solution and were broken down but shorter Ti\(-\text{O}\) bonds remained unchanged, leading to the formation of Ti\(-\text{O}\)-K bonds.\(^{21}\) Thus, KT-NW was successfully formed by a simple hydrothermal method at 200 °C. From XRD patterns in Fig. 2(a), the pristine TiO\(_2\) nanoparticles showed intense peaks at 2\(\theta\) of 25.37°, 37.87°, 48.16°, 54.00°, 55.21°, and 62.84° which are perfectly aligned to TiO\(_2\) anatase (COD ID 7206075) phase. The synthesized KT-NW corresponded to the crystal phase of K\(_2\)Ti\(_6\)O\(_{13}\) (AMCSD no: 0010602)\(^{22}\) and the intense peaks can be indexed as 200, 110, 310, 112, 402, 404, 020, and 422.

The starting material Ti\(_3\)AlC\(_2\) (MAX) shows the standard XRD pattern (attached in Fig. S2†) as reported in the literature.\(^{23}\) MXene or Ti\(_3\)C\(_2\)T\(_x\) was obtained chemically etching the MAX phase with HCl and LiF at 50 °C for 48 h, where HF was produced \emph{in situ} in the reaction. The characteristic peaks for the MAX phase were disappeared but several peaks remained even after selective chemical etching of Al by HCl/LiF as the mild etching agent.\(^{24}\) The intense peak at 002 (9.5°) of MAX phase has reduced in MXene while a new characteristic sharp peak appeared at a low angle (6.4°) confirming the delaminated layered MXene structure. The XRD pattern of the MXene is in accordance with the literatures.\(^{24,25}\)

The hydrothermal treatment at 120 °C led the formation of MKT-NW composite. The XRD pattern of MKT-NW composite ascribes the relevant peaks for MXene as well as KT-NW which confirmed the existence of the KT-NW in the MKT-NW composite. Furthermore, KT-NW would form less agglomeration compared to the TiO\(_2\) particle when introduced directly to MXene.\(^{26}\)

FE-SEM images illustrates the formation of a layer structure of MXene that has been formed due to the delamination and exfoliation by \emph{in situ} HF. Furthermore, EDS mapping has been
conducted to determine the amount of chemically etched Al from the MAX phase during its transition to form MXene (Fig. S3†). TiO$_2$ nanoparticles were completely transformed by a hydrothermal technique in the presence of concentrated KOH aqueous solution in the reaction system. Additionally, FE-SEM and TEM images show the nanowire morphology of KT-NW on the surfaces of MXene as illustrated in Fig. 2(c). The KT-NW has been compactly deposited on the surface of MXene as pleated nano wires by covering most of the surface area by developing a strong electrostatic interaction between them. The formation of MKT-NW had similar morphology with that of the work carried out to prepare MXene with TiO$_2$ nanofibers.\textsuperscript{27} In Fig. 3, the survey spectrum of MKT-NW shows the presence of C 1s, K 2p, Ti 2p, O 1s, and F 1s. All peaks precisely coincide with the binding energies given in the XPS database.\textsuperscript{28,29} The core level high resolution XPS analyses of MKT-NW powder shows the peaks of carbon (C 1s) is mainly found at 285.5, 286.8, and 289.7 eV which are assigned to sp$^3$ C–C, C–O, and C–F, respectively. The K 2p signals (K 2p$_{3/2}$ at 293.5 eV and K 2p$_{1/2}$ at 296.8 eV) and KF at 296.1 eV were revealed to formation of potassium titanates from hydrothermal of TiO$_2$ in alkaline media. The Ti 2p region shows two signals at Ti 2p$_{3/2}$ (ranging from 453 to 462 eV) and Ti 2p$_{1/2}$ (462 to 468 eV) with oxidation states (Ti$^{2+}$, Ti$^{3+}$ and Ti$^{4+}$) according to the formation of mixed oxides (TiO$_x$F$_y$)$_3$ and titanium oxy carbide (TiC$_x$O$_y$) which revealed non-uniform surface of MKT-NW. Deconvolution of O 1s also confirms the formation of Ti(IV) oxide and the presence of OH/O$_2$ groups. The F 1s spectrum exhibited a strong signal at 685.2 eV which assigned to fluorine binding to the Ti carbide (Ti–F), thereby creating a C–Ti–F bond. The F 1s signal at the slightly higher binding energy of 687.2 eV can be attributed to C–Ti–OF or a fluorine bridging atom (C–Ti–F–Ti–C) or even an Al–F in MAX phase.\textsuperscript{30} This result is consistent with EDS mapping (Fig. S3†), about 2.9% wt of Al (or 2.6% atom) leaching from the MAX phase.

The surface morphology of pristine C-SPEEK and MKT-NW/C-SPEEK composites film revealed smooth and rough surface characteristics, respectively (Fig. S4†). The XRF elemental mapping of 12%MKT-NW/C-SPEEK hybrid composite PEM (Fig. 4) exhibited mainly S, Ti, K and Al with a uniform spatial dispersion within the polymer matrix. The distribution of Ti element within the polymer matrix was observed which were derived from MXene and KT-NW, along with the trace amount of K and Al elements.

**Membranes properties**

The water molecule plays a vital role in the proton transfer mechanism as it can be used as a proton carrier to transfer protons from anode to cathode. The prepared SPEEK had high DS, thus the SPEEK membrane showed high water uptake and a swelling area which was later decreased after cross-linking with EG.\textsuperscript{31} But, it was observed that there was an increase in water uptake after introducing MKT-NW into the C-SPEEK membrane; which was due to the hydrophilic nature of MXene and KT-NW.\textsuperscript{32} However, the swelling area was decreased with the increase of MKT-NW which may be due to the physical interaction of –SO$_3$H of SPEEK and –OH of EG with MXene and KT-NW. The water uptake of 20% MKT NW/C-SPEEK was low and showed a high swelling area, which may be due to the accumulation of the MKT-NW.\textsuperscript{31} The water uptake and swelling area of all the prepared membranes are given in Table 1.

In the proton exchange membrane, ion exchange capacity is one of the important factors that insight the ionic sites of each PEM which relates to the proton conductivity. Here, C-SPEEK membrane had low IEC of 1.42 meq g$^{-1}$ due to the cross-linking of ethylene glycol with the –SO$_3$H group of SPEEK which caused the reduction of acidic sites.\textsuperscript{33} An IEC increased after MKT-NW was incorporated with C-SPEEK and further increased with an increasing in the amount of MKT-NW\textsuperscript{32} The

| Samples       | Water uptake (%) | Swelling area (%) | Proton conductivity (S cm$^{-1}$) |
|---------------|------------------|-------------------|-----------------------------------|
| SPEEK         | 23.40            | 21.01             | 0.0076                            |
| C-SPEEK       | 10.00            | 18.91             | 0.0034                            |
| 5%MKT-NW/C-SPEEK | 21.05           | 16.74             | 0.0038                            |
| 10%MKT-NW/C-SPEEK | 47.36           | 10.35             | 0.0061                            |
| 12%MKT-NW/C-SPEEK | 48.57           | 8.15              | 0.0097                            |
| 14%MKT-NW/C-SPEEK | 51.42           | 12.46             | 0.0078                            |
| 20%MKT-NW/C-SPEEK | 38.88           | 16.73             | 0.0066                            |
12%MKT-NW/C-SPEEK composite hybrid membrane performed highest IEC of 1.88 meq g⁻¹ according to the well distribution of MKT-NW within the membrane compared to the others. Moreover, a further decrease in the IEC was observed at the amount of MKT-NW more than 12% which may be due to the agglomeration of the inorganic filler.³¹ The IEC of all the prepared membrane is presented in Fig. 5(a).

The proton conductivity of all MKT-NW/C-SPEEK composite membranes at room temperature is shown in Table 1. The pristine SPEEK membrane exhibits high proton conductivity according to the presence of a large number of –SO₃H group in the hydrophilic domain part of the polymer. In addition, the casted membrane using ethanol and water (SPEEK) which have lower interaction as compared to the other casted SPEEK membranes using an organic solvent, leading to high proton conductivity of membranes.³² However, the conductivity thereby decreased as it was cross-linked with EG because the number of –SO₃H group would become lower as it will be cross-linked with –OH group of EG. Furthermore, after MKT-NW was incorporated with C-SPEEK, the proton conductivity increased as shown in Fig. 5(a). It was due to the hydrophilic nature of MXene as well as KT-NW which absorbed more water molecules that acts as the proton carrier in vehicular type mechanism. Nevertheless, the presence of –SO₃H from SPEEK and –OH from composite as well as EG would develop an extensive interconnected hydrogen bond network and shorten the distance for faster proton transfer pathway, through Grothuss mechanism, thereby increasing proton conductivity.³³,³⁴ The proton conductivity increased with an increasing in the amount of MKT-NW and the addition of MKT-NW more than 12% decreased the proton conductivity, which may be due agglomeration and non-uniform distribution of the composite. The 12%MKT-NW/C-SPEEK exhibits proton conductivity of 0.0097 S cm⁻¹ which was higher than that of the commercially available Naion-212 (0.0093 S cm⁻¹) due to the well distribution of composite in the polymer matrix.

Dynamic mechanical strength
To elucidate the effect of MKT-NW composite on C-SPEEK dynamic mechanical analysis, the composite membrane with 5%MKT-NW and 20%MKT-NW were considered as the minimum and maximum MKT-NW content, respectively. The storage modulus of C-SPEEK was increased when 5% MKT-NW was added whilst 20% MKT NW led to the further increment of storage modulus as shown in Fig. 5(b) and Table 2. The highest storage modulus of 20%MKT-NW/C-SPEEK was 5436 MPa which was 1.44 times higher than that of C-SPEEK (3762 MPa). Thus, the addition of MKT-NW improved the storage modulus of C-SPEEK, which was due to the strong interaction of highly hydrophilic MKT-NW developed with C-SPEEK. Hence, rendering improved mechanical strength to the MKT-NW/C-SPEEK hybrid composite PEM.

The glass transition temperature (Tg) of C-SPEEK, 5%MKT-NW/C-SPEEK, and 20%MKT-NW/C-SPEEK membranes were depicted using the plot of tan δ various with temperature. The Tg of C-SPEEK membranes was 221 °C while the addition of MKT-NW leads to the small increment of Tg membrane (~1 °C for 5%MKT-NW and ~2 °C for 20%MKT-NW) because of the formation of dense intermolecular hydrogen bonding within MKT-NW and C-SPEEK.

C-SPEEK had higher Tg value than SPEEK membranes as reported by X. Zhang et al. (181 °C)³⁷ after cross-linking with EG and the addition of MKT-NW does not bring noteworthy change in the Tg of C-SPEEK membranes. Although it is worth mentioning that MKT-NW had significantly affected to the mechanical strength of C-SPEEK membranes as observed from the storage modulus data at the different temperatures as summarized in Table 2. The C-SPEEK using EG as a cross linker had shown higher storage modulus as compared to the cross linked membrane using polyethylene glycol (PEG) or 1,4-cyclohexane dimethanol (CDM) as a cross linker.³⁸ Additionally, the MKT-NW/C-SPEEK had shown higher storage modulus than SPEEK composite membranes which was due to the cross-linking of SPEEK with EG as well as MKT-NW composite to form a strongly hydrogen-bonded within the hybrid composite membranes.³⁹

Photocatalytic activity of hydrogen production from water
The prepared MKT-NW/C-SPEEK hybrid composite membrane showed photocatalytic activity due to the presence of MKT-NW
in the polymer matrix. The optimized composite membrane (12%MKT-NW/C-SPEEK) was used to determine the effect of UV light intensity on photocatalytic H₂ production. Initially, the UV light was located close to the membrane in which ~30% of the total surface of the membrane was irradiated. And the light was moved further from the membrane to irradiate the total area of the membrane (~100%). The intensity of the light depends upon the distance of the light source and the exposed area of the membrane. The MKT-NW got more excited when the UV light intensity was higher (irradiated ~30% of the area), and this was compared with the membrane which had lower intensity of UV light (irradiated ~100% of the membrane). A significant effect on the photocatalytic H₂ production from water using the membrane had been noticed as the ~30% irradiated membrane produced 0.185066 µmol of H₂ within 5 h and ~100% irradiated membrane produced 0.096649 µmol H₂ within 5 h as shown in Fig. 6(a).

As the ~30% irradiated membrane with higher intensity could produce more H₂ gas, this irradiation was considered to study the effect of the MKT-NW composite in the photocatalytic activity of membranes. Two-hybrid composite membranes namely 12%MKT-NW/C-SPEEK and 5%MKT-NW/C-SPEEK had been selected to study the photocatalytic H₂ production. In order to know the effect of MKT-NW content to the photocatalytic activity, 5% and 12%MKT-NW/C-SPEEK were used in this study. It had been observed that both membranes could produce H₂ linearly with respect to time within 5 h. The H₂ production of 12%MKT-NW/C-SPEEK was slightly different from 5%MKT-NW/C-SPEEK. This might depend on the higher amount of MKT-NW in 12%MKT-NW/C-SPEEK composite PEM as shown in Fig. 6(b).

When UV light was irradiated the KT-NW in the polymer matrix, KT-NW would be excited to produced electron–hole pair (e⁻ and H⁺). Meanwhile, the valence band (VB) of KT-NW would remain with H⁺ since the e⁻ in the conduction band (CB) of KT-NW would swiftly move towards MXene and accumulated on the surface because of the superior metallic and electronic properties. Then, a Schottky barrier would develop on the interface of MKT-NW composite blocking the back transfer of e⁻ to KT-NW. And the H⁺ in the aqueous methanol solution would undergo reduction with the help of the photo-induced e⁻ on MXene producing H₂ (ref. 13–40) as presented in Scheme 3. Thus, making this hybrid composite PEM as a photocatalytic active membrane that produces H₂ from water without the addition of any noble metals as co-catalysts in the same way as that of the photocatalytic membranes prepared by immobilizing rGO/TiO₂ into Naion membranes.⁴¹

The membranes before and after H₂ evolution testing under UV irradiation were characterized by XPS wide scan and core level high resolution. Fig. 7 illustrates XPS spectra for MKT-NW and their composites with SPEEK. The spectra of MKT-NW (before and after H₂ evolution testing) shows the signals of F 1s, O 1s, Ti 2p, K 2p, and C 1s. For the MKT-NW/SPEEK composites, new peak of S 2p region ranging from 164 to 176 eV, which revealed the presence of sulfonyl entirely in the composite membranes, and the disappearance of the peak due to F 1s were observed. The depth of the XPS surface analysis is about 10 nm, which is the limitation of XPS characterization.
Table 3  The apparent atomic compositions of the initial MKT-NW and the composite samples as determined by XPS

| Materials         | C 1s  | O 1s  | Ti 2p | F 1s  | K 2p  | S 2p  |
|-------------------|-------|-------|-------|-------|-------|-------|
| MKT-NW            | 36.2  | 44.7  | 15.6  | 1.35  | 2.16  | 0.00  |
| 5%MKT-NW/C-SPEEK  |       |       |       |       |       |       |
| Before testing    | 77.4  | 19.4  | 0.28  | N/A   | 0.40  | 2.58  |
| After testing     | 71.5  | 27.1  | 0.15  | N/A   | 0.66  | 0.63  |
| 12%MKT-NW/C-SPEEK |       |       |       |       |       |       |
| Before testing    | 78.8  | 17.7  | 0.24  | N/A   | 0.72  | 2.51  |
| After testing     | 88.8  | 9.85  | 1.03  | N/A   | 0.15  | 0.22  |

Table 4  Comparison of present composites activity over the reported SPEEK-functionalized materials

| Membranes | Proton conductivity (S cm⁻¹) | Temp. (°C) | Ref. |
|-----------|-------------------------------|------------|------|
| 7.5%TiO₂–SO₃H/SPEEK | 0.0138 | 30 | 42 |
| 10%TiₓCₓTeₓ/SPEEK | 0.1470 | 40 | 11 |
| 4%CNx/C-SPEEK | 0.1850 | 95 | 20 |
| 5%Clay/C-SPEEK | 0.0197 | 80 | 43 |
| 8%SSi-GO/SPEEK | 0.1626 | 65 | 44 |
| 12%MKT-NW/C-SPEEK | 0.0097 | 25 | This work |

Conclusions

MKT-NW/C-SPEEK hybrid composite PEM had been successfully prepared. MKT-NW was prepared by a hydrothermal method and was used as the filler to improve the physicochemical properties of membranes. The MKT-NW led to improve the mechanical strength of the C-SPEEK membrane. The 12%MKT-NW/C-SPEEK had the highest proton conductivity of 0.0097 S cm⁻¹ with the IEC of 1.88 meq g⁻¹. Besides, it was a photocatalytic membrane which was able to produce 0.1850 μmol of H₂ in 5 h by photocatalysis of water. Consequently, the prepared MKT-NW/C-SPEEK hybrid composite PEM is a smart PEM with dual properties showing the photocatalytic activity of H₂ production from water, along with the proton conductivity improvement. The overall process utilized was green and sustainable as the mixture of ethanol and water was used as the solvent for PEM casting rather than using hazardous organic solvents. Thus, MKT-NW/C-SPEEK hybrid composite PEM could be utilized in many applications namely; PEM for PEM fuel cells, direct methanol fuel cells, microbial fuel cells and PEM water electrolysis or artificial photosynthesis, which may induce a useful impact on the field of clean and green energy.

Abbreviations

Mxene | Titanium carbide
Ti₆C₄Te₂ | Potassium titanate nanowire
KT-NW (K₂Ti₆O₁₃) | Sulfonated polyether ether ketone
SPEEK | Ethylene glycol
C-SPEEK | Cross-linked sulfonated polyether-ether ketone
MKT-NW/C-SPEEK | MXene-potassium titanate nanowire/cross-linked sulfonated polyether-ether ketone
DS | Degree of sulfonation
PEM | Proton exchange membrane
IEC | Ion exchange capacity
HER | Hydrogen evolution reaction

Conflicts of interest

There are no conflicts to declare.

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