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Functionalized Titanate Nanotube-Polyetherimide Nanocomposite Membrane for Improved Salt Rejection under Low Pressure Nanofiltration †

Anappara Sumisha a, Gangasalam Arthanareeswaran a,b, Ahmad Fauzi Ismail b, Dharani Praveen Kumar a, Muthukonda V. Shankar c

1. Introduction

Design and development of efficient materials for nanofiltration (NF) membrane is a great societal demand. NF membrane has been widely used for desalination of electrolyte solution and its performance can be tuned using appropriate type and density of charge carriers on the membrane surface. Mixed matrix membranes (MMMs) composed of nanoparticles blended with polymeric membrane has proved the enhancement of performance in terms of permeability, solute rejection and better anti-fouling properties. 1,2 In MMMs, mechanical mixing of inorganic nanomaterials into polymer solution is beneficial for homogeneous dispersion of nanoparticles in matrix. 3 To date, variety of nanosize metal oxides like TiO 2, 4,5 SiO 2, 6,7 Al 2O 3, 8 ZrO 2, 9,10 ZnO 11 and zeolites 12,13 were successfully incorporated into polymers as an additive to improve the performance such as hydrophilic and flux of membrane.

Functionalized titanate nanotubes were prepared using facile and eco-friendly method. Nanofiltration membranes were fabricated via simple phase inversion method. The neat and mixed matrix membrane (MMMs) was prepared using PEI as polymeric material and nanomaterials such as TiO 2 particles (TP), as-synthesized hydrogen trititanate nanotubes (pTNT), N-doped TiO 2 NT (N-TNT) and Cu-doped H 2 Ti 3O 7NT (Cu-TNT) were served as additives. The crystal phase characterization revealed the anatase phase for TP, trititanate phase for pTNT, anatase-rutile mixed phase for N-TNT, Cu-TNT materials and the similar observations were found with MMMs. The morphology analysis of neat PEI membrane exhibited a denser top layer and the beneath part of membrane is tighter. Different from neat PEI membrane nanocomposites of MMMs showed finger-like macrovoids towards the bottom of the membrane. The water uptake and hydrophilic character of membranes are found in the following order neat PEI > PEI/TP > PEI/pTNT > PEI/N-TNT > PEI/Cu-TNT. Interestingly, the salt rejection performance of monovalent (NaCl) and divalent (K 2SO 4 and CaCl 2) ions in the single salt mixture were found to increase in the same order. The salt rejection performance of PEI/Cu-TNT was found in the decreasing order K 2SO 4 (80%) < NaCl (75%) < CaCl 2 (45%). The high performance of PEI/Cu-TNT in salt rejection and antifouling properties are ascribed to tubular morphology, copper dopant results high hydrophilic character of the MMMs.

Nanomaterials exhibited superior physic-chemical properties, such as large surface area leads to vast majority of the active sites on the surface, improved catalytic, optical and electrical properties and homogeneous dispersion in solution. 14 The physico-chemical properties of TiO 2 material displayed its multifunctional application for photocatalysis, heterogeneous catalysis, dye-sensitized solar cells, lithium ion batteries, biological applications, ultrafiltration membrane and so on. 15-18 The purpose of polymer membrane modification with functionalized nanoparticles is to minimize membrane fouling due to adsorption or adhesion, or to introduce functional interactions to improve performance. 19 Nano-size TiO 2 particles has established much consideration to enhance membrane properties due to its high hydrophilicity, great chemical stability, antibacterial and abundant in nature. 20-22 Arrachart et al 23 successfully dispersed functionalized TiO 2 nanoparticles in PMMA matrix by phase separation method. The nanoparticles blended with polymer solution results strong hydrogen or covalent bond interaction leads to improved dispersion. 24

Polyetherimide (PEI) is the most preferred polymer for membrane applications as it displayed mechanical strength, thermal stability, film forming ability and chemical resistance under vigorous experimental conditions. 25 However, PEI membrane suffers with hydrophobic character. To overcome this, hydrophilic PEI/silica membranes fabricated by incorporating the fluorinated silica nano particles (fSiO 2). 26 The results revealed that incorporation of fSiO 2 layer greatly enhanced the hydrophilic...
property of membrane with improved mechanical strength. The
effect low TiO₂ concentration on the permeability and membrane
fouling of PES membranes was studied using polar solvent of 1-
ethyl-2-pyrolidone.²⁷ Authors concluded that TiO₂ nanoparticles
entrapped in membranes have more open structure, higher
hydrophilicity and permeability.

TiO₂ based nanotubes have attracted wide attention owing to
their potential for application in electrochemical separation
process.¹⁶,²¹ The free standing TiO₂ nanotubular membrane was
prepared by growth of a high aspect ratio TiO₂ nanotubular layer
on Ti, selective dissolution of the metallic substrate and opening of
the closed tube bottom by selective chemical etching.²⁸ Rajesh
Kumar et al²⁹ suggested that desalination application of TiO₂
nanotubes incorporated polysulphone (PSf) membranes. Further,
the detailed study about antifouling and permeation properties of
PSf/TiO₂ nanotubes membranes was investigated. The lower ratio
of TiO₂ concentrations provided a significant improvement
in the performance of blended membranes.³⁰ Addition of
functionalized nanomaterials into polymer membranes provide
the benefit of physical (porous structure) and chemical changes
(functional groups attachment) could lead to enhancement of
separation functions. Yang et al³¹ and Tedja, et al³² reported the
adsorption of serum proteins on to the surface of TiO₂
nanoparticles results in reduction of biological impact compared
to the non-protein treated TiO₂ particles. The literature survey on
TiO₂ based polymeric membranes confirms that the hydropoicity and antifouling properties were tuned using nano-
size particles and nanostructures as well. To the best of our
knowledge doped TiO₂ (copper or nitrogen) was not yet studied
as additives for membrane preparation.

For the first-time we have developed novel PEI nano-composite nanofiltration membranes by incorporating
functionalized TiO₂ nanotubes for enhanced desalination
performance. The effect of TiO₂ nanoparticles and functionalized
nanotubes such as hydrogen trititanate (H₂Ti₃O₇) nanotubes, N-
doped TiO₂ nanotubes and ion-exchanged Cu²⁺/H₂Ti₃O₇
nanotubes on PEI mixed matrix membranes were studied. The
physico-chemical properties of the TiO₂ based nanomaterials
were thoroughly characterized. The salt rejection performance
and anti-fouling property of modified PEI membranes were also
investigated.

2. Experimental Section

2.1. Synthesis and functionalization of H₂Ti₃O₇ nanotubes

The H₂Ti₃O₇ nanotubes were synthesized by hydrothermal
method as reported in our earlier publication.³³ In a typical
synthesis process, TiO₂ (LAB, Merck) powder (2.5 g) was
dispersed in 10 M NaOH aqueous solution (200 mL) and cooked
at 130 °C for 20 h in Teflon-lined autoclave (capacity 250 mL).
The white precipitate was subjected to washing (twice)
with distilled water followed by 0.1 M HCl, finally washed with
ethanol, and dried at 80 °C for 12 h. The obtained bright white
powder was functionalized through ion-exchange with copper
solution and N-doping using urea granules.

Ion-exchange method was employed to prepare modified Cu-
TNT using an appropriate amount of copper nitrate in aqueous
solution.³⁴ For instance, 0.5 gram of H₂Ti₃O₇ was suspended in
100 mL of 0.05M Cu(NO₃)₂ aqueous solution in 250 mL glass
beaker and stirred well. After 2 hours stirring, the suspension
was filtered under vacuum and washed thoroughly with 100 mL of
distilled water for 2 times. It was noticed that after ion-exchange
the filtrate solution exhibited low optical density than initial
solution, indirectly confirms the ion-exchange reaction with TNT
material. Thus obtained pale blue color precipitate Cu-TNT was
dried in hot air oven for 80°C for 12 h.

Wet-impregnation method was adopted for the preparation
of N-doped TiO₂ nanotubes.³³ First, 2 g of urea granules dissolved
in 10 mL of distilled water in 25 mL glass beaker. Into this
solution, 1 g of H₂Ti₃O₇ nanotubes was dispersed. Then contents
were allowed for magnetic stirring at room temperature for 30
min followed by heating at 100 ± 5 °C for 2 h till water
evaporation to dryness. Thus resulting powder mixture was dried
at 80 °C for 16 h before calcination at 350°C for 2 h. Pale yellow
powder was obtained.

2.2. Membrane preparation

Neat and functionalized titania/PEI Nanofiltration membranes
were prepared by phase inversion method using water as non-
solvent.³⁶ A neat PEI membrane at 17.5 wt% in NMP was used as
polymer matrix. The functionalized Ti-O based membranes were
prepared by dispersing 0.5 wt% of TP, pTNT, N-TNT and Cu-
TNT as additives. The casting solutions were prepared by adding
PEI and respective nanoparticles in 20 mL of NMP as solvent at
room temperature and sonicated for 2 h to disperse nanoparticles
uniformly in solution. It is then stirred continuously until clear
homogenous solutions were obtained. The PEI-additive solution
then cast onto the glass plate for the thickness of about 400 μm
with the help of a thin film applicator. Then, the glass plate
immediately immersed into a distilled water bath maintained at
20 °C. The synthesized pure and modified PEI membranes cut
into required area corresponding to dead-end experiment
employed in this study. The details of the nanomaterials used for
the functionalized membrane preparation are given below:

<table>
<thead>
<tr>
<th>Additive Material / Membrane</th>
<th>Sample ID</th>
</tr>
</thead>
<tbody>
<tr>
<td>TiO₂ (LAB, Merck, India)</td>
<td>TP</td>
</tr>
<tr>
<td>Pristine H₂Ti₃O₇ nanotubes</td>
<td>pTNT</td>
</tr>
<tr>
<td>Nitrogen doped TiO₂ nanorods</td>
<td>N-TNT</td>
</tr>
<tr>
<td>Copper ion-exchanged H₂Ti₃O₇</td>
<td>Cu-TNT</td>
</tr>
<tr>
<td>PEI blended with TP, pTNT, N-TNT and Cu-TNT</td>
<td>PEI/TP, PEI/pTNT, PEI/N-TNT and PEI/Cu-TNT</td>
</tr>
</tbody>
</table>

2.3. Characterization of nanoparticles and MMMs

The prepared and modified nanomaterials as well as membrane
were thoroughly examined using different characterization
techniques. Transmission electron microscopy (TEM) images
were collected using a FEI Tecnai F20ST electron microscope
operated at 200 keV and energy dispersive X-ray (EDX)
spectrometer. Powder X-ray diffraction (XRD) patterns of all the
materials were recorded using a D8 ADVANCE X-ray diffractometer (Bruker) equipped with Ni-filtered Cu Kα
(k = 1.5418 Å) radiation (30 kV, 50 mA). Scanning Electron
Microscopy (Model: TESCAN, SEM-5600) was utilized to study morphology of PEI and nanomaterials incorporated PEI membranes. Thermal properties of neat PEI and functionalized PEI membranes were quantified by thermo-gravimetric analysis (TGA). The measurements were taken using TGA thermal analyser (Model: DGT 2000, Perkin Elmer) that has been interfaced to a computer. Before loading 5 to 10 mg of membrane samples into the alumina crucible, the test samples were dried at 180 °C overnight in a vacuum oven to remove moisture. The thermogram was recorded at the temperature range of 10 to 600 °C and at the heating rate of 10 °C.min\(^{-1}\) under the supply of nitrogen gas. The thermal decomposition temperature range and the respective weight% loss can be studied and compared by this study.

2.4. Hydrophilicity
The hydrophilicity of the membranes was measured by an automated contact angle goniometer (Model: Ramehart, USA). Distilled water droplet of 0.5 μL was placed on the dry membrane surface and the equilibrium contact angle was calculated by determining the incident and receding angles. At least five different locations were selected arbitrarily on the membrane surface in order to yield an average value of equilibrium contact angle of the membranes.

2.5. Water uptake capacity and membrane porosity
Membrane samples were cut into desired size and soaked in water for 24 h and weighed immediately after blotting the free surface water (W\(_1\)). Wet membranes were dried for 24 h in vacuum oven and weighed (W\(_2\)). The percent water uptake was calculated using the following equation:

\[
\text{Water uptake capacity (\%) } = \frac{W_1 - W_2}{W_1} \times 100
\]

The porosity of membrane samples can be analysed by considering the weight of membranes at dry and wet states. It can be calculated by the following equation:

\[
\text{Porosity (\%) } = \frac{W_1 - W_2}{\rho_w \times A \times l} \times 100
\]

Where, \(\rho_w\) = density of water at room temperature (1 g/cm\(^3\)); \(A\) = area of membrane (cm\(^2\)); \(l\) = thickness of wet membrane.

2.6. Membrane performance
The performance of the membrane was studied by checking its pure water flux to understand the water transport behavior of the membrane. Each membrane was washed and cut into circular shape to be fitted into the stirred cell. It is initially pressurized with 500 kPa trans membrane pressure (TMP) for 30 min to reduce compaction effects. The pure water flux of the membrane was then measured at the same TMP using the following equation:

\[
J_w = \frac{V}{A \times \Delta t}
\]

Where \(J_w\) is the pure water flux in Lm\(^{-2}\)h\(^{-1}\), \(V\) is the volume of water permeated in liters, \(A\) is the area of membrane in m\(^2\) and \(\Delta t\) is the time for permeate collection in hours.

The salt rejection study of membrane was carried out by treating membranes with salt solution of K\(_2\)SO\(_4\), NaCl and CaCl\(_2\) with different concentration in dead end mode filtration. The synthetic salt solutions were prepared in four different concentrations i.e, 500, 1000, 1500 and 2000 mg.L\(^{-1}\). All the experiments were carried out at room temperature 30 ± 2 °C. The synthetic salt solutions then passed through five different fabricated NF membranes in dead end filtration cell at 500 TMP and flux was calculated by using the equation 3. The concentration of salts in feed and permeate was determined by using the conductivity of solution. The rejection of salt was calculated using the following equation:

\[
R (\%) = \left(1 - \frac{C_p}{C_f}\right) \times 100
\]

Where, \(C_f\) and \(C_p\) are feed and permeate concentrations of the salts, respectively.

The biofouling performance of the MMMs was evaluated based on the static adsorption study of protein solution. Typically, 100 mg.L\(^{-1}\) of egg albumin solution was prepared by dissolving 0.1 g of egg albumin powder in100 mL distilled water. Adding 1M NaOH or 1M HCl solutions adjusted the pH to 3, 5, 7, and 9. The membrane samples were cut into small pieces (2 × 2 cm), placed inside the vial containing 10 mL of egg albumin solution, and kept in shaker for 24 h to allow for albumin adsorption. The amount of egg albumin adsorbed on the membrane samples were calculated by using the initial and final solution of albumin concentrations by UV-visible spectrometer at a wavelength of 270 nm.

3. Results and discussion
3.1. Morphology analysis of H\(_2\)Ti\(_3\)O\(_7\) nanotubes
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Fig. 1 displays TEM image of H$_2$Ti$_3$O$_7$ nanotubes prepared via hydrothermal method. Fig. 1a shows (200 nm scale) the bundles of randomly oriented one dimensional nanotubes and overlapping of nanotubes results dark spots at different locations. The nanotubes at smaller scale (10 nm) are evident for multi-layered structure of nanotubes with hollow inside and open ended on the both edges. It is observed that, nanotubes having length several hundred of nanometer and internal diameter of 8-12 nm is observed (Fig. 1b). Fig. 1c displays selected area electron diffraction (SAED) pattern of the same material matches very well with XRD pattern (vide infra) confirms that nanotubes made-up of H$_2$Ti$_3$O$_7$. The improvement in properties of the functionalized PEI membrane depends on the available polymer–filler interfacial surface area, which is greatly influenced by both size and amount of nanomaterials used as filler. Based on the literature, we understand that optimum loading of filler material is essential to avoid aggregation we have restricted TiO$_2$ nanotubes loading 0.5 wt% used to improve the performance.

3.2. Crystal phase identification

X-ray diffraction patterns of TiO$_2$ precursor, synthesized and doped titanate nanotubes are shown in Fig. 2. The XRD pattern of pTNT material (inset) displays the weak, broad diffraction pattern and peaks centred at 20 = 10.2, 24.1, 28.3 and 48.2° are well indexed as the monoclinic structure of hydrogen tritiatanate, H$_2$Ti$_3$O$_7$ (JCPDS No.47-0561). Similar diffraction pattern observed for Cu1TNT material and it is identified as tritiatanate material. The XRD pattern of N1TNT is changed, indicating that the structural transformation had occurred during calcination process. The diffraction pattern of N1TNT depicted two characteristic peaks at 20 = 25.4 and 27.5° confirmed presence of tetragonal structure of anatase and rutile (JCPDS No.21-1272 & 21-1276). Similar observations on crystal phase change upon calcination were demonstrated in our earlier report. TiO$_2$ particles (TP) used as precursor for pTNT preparation showed a major diffraction peak at 20 = 25.4° corresponding to anatase phase (JCPDS No.21-1272). The diffraction pattern of pTNT and Cu-TNT exhibited semi-crystalline nature, whereas N-TNT and TP exhibited sharp peaks ascribed to well crystalline structure. The diffraction patterns of neat PEI and MMMs are displayed in Fig. 3. All the blended membranes show sharp or broad peaks corresponding to diffraction pattern of TP, N-TNT and pTNT respectively. The neat PEI showed minor diffraction peaks at 20 = 21 and 29° and matches well with earlier reports. The MMM samples displayed characteristic peaks of neat PEI and titania based materials TP, pTNT, N-TNT and Cu-TNT.

3.3. Morphology of PEI and MMM membranes

The membrane surface and cross-section morphologies of the neat PEI and functionalized PEI membranes were investigated by scanning electron microscopy (SEM) analysis. The influence of nanoparticles addition onto the membrane surface is demonstrated in Fig. 4. The neat PEI membrane shows a denser top layer and the beneath part of membrane is tighter. It was observed that a change in cross-sectional morphology of membranes with the addition of nanomaterials. This may be due to the interlinking of hydrophilic TiO$_2$ nanomaterials with hydrophobic PEI and which results in the fast exchange of solvent and non-solvent during phase inversion process and which in turn results in the increase of number of pores in functionalized PEI than neat PEI. Further, the fast exchange of solvent and non-solvent in the phase inversion process due to the hydrophilic nature of functionalized TNT and existing interactions between components in the casting solution and phase inversion kinetics.

It develops a broad finger like macro voids in the PEI/TP membrane than neat PEI membrane. In the third membrane, the finger-like macrovoids were gradually grown towards the bottom of the membrane and the shape of macrovoids became spongy...
like porous structure when pTNT nanotubes were added into PEI membrane. The similar morphological behavior by addition of silver nanoparticles was observed by our earlier work. This may be due to decrease of interfacial stresses existed between polymer and layered H$_2$Ti$_3$O$_7$ nanotubes, which were tensed by forming interfacial pores due to growth of the organic phase during the delayed demixing process. An extremely sponge like porous structure is observed in functionalized PEI membranes. Thus, morphology and surface properties of N1TNT and Cu1TNT nanomaterials were responsible for the large pore size and high water permeability of membranes. Celik et al also reported similar morphological behavior.

3.4. Thermal stability of PEI and MMM membranes

The thermal stability characteristics of neat PEI and MMMs samples were evaluated by thermogravimetric analysis (TGA) and results are presented in Table 1. The TGA results of neat PEI and mixed matrix membranes with titania materials viz., TP, pTNT, N-TNT and Cu-TNT nanotubes were illustrated in Fig. 5. The temperature range of decomposition of the membrane and their corresponding weight % losses determine the thermal stability of the membranes. The first step of degradation corresponded to a slight but steep drop in residual weight % due to small weight loss observed between 97.4 and 234 °C for all the membranes due to evaporation of water and the residual NMP solvent (its boiling point is 202 °C). The major thermal degradation of membranes mainly occurs in the temperature range of 407 to 504 °C. However, the thermal degradation of membrane samples decreases with the addition of nanoparticles. Among the membranes PEI/Cu-TNT shows lesser degradation and higher thermal stability. It shows a 7.8% weight loss at a temperature range of 409 to 506 °C. Whereas, the neat PEI membrane shows higher thermal degradation with 13.9% weight loss. The other membranes of PEI with N-TNT, pTNT and TiO$_2$ nanoparticles showed a weight% loss of 8.2%, 8.3% and 8.9% respectively. The TGA results also confirmed the presence of inorganic nanoparticles. During membrane fabrication, there is a chance for the nanoparticle to leach out at the time of phase inversion. The above results illustrates that the nanoparticles are still in the membrane and which enhances the thermal stability of mixed matrix membranes.

Table 1 Thermal stability of nanomaterials functionalized PEI membranes

<table>
<thead>
<tr>
<th>Sample</th>
<th>Decomposition stage</th>
<th>Temperature range (°C)</th>
<th>Weight loss(%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Neat PEI</td>
<td></td>
<td>442</td>
<td>13.9</td>
</tr>
<tr>
<td>PEI/TP</td>
<td></td>
<td>445</td>
<td>8.9</td>
</tr>
<tr>
<td>PEI/pTNT</td>
<td></td>
<td>458</td>
<td>8.3</td>
</tr>
<tr>
<td>PEI/N-TNT</td>
<td></td>
<td>456</td>
<td>8.2</td>
</tr>
<tr>
<td>PEI/Cu-TNT</td>
<td></td>
<td>456</td>
<td>7.8</td>
</tr>
</tbody>
</table>

Fig. 4 SEM images of top surface (left) and cross-section (right) of neat and MMM membranes.
3.5. Hydrophilicity and water permeation of nanomaterials and functionalized PEI membranes

The hydrophilicity of membrane influences both water permeation and desalination performance. Higher contact angle corresponds to a hydrophobic surface whereas smaller contact value corresponds to hydrophilic surface of membranes. Fig. 6 displays the surface contact angles of PEI membranes blended with TNT and functionalized TNT nanomaterials. The contact angle of PEI membrane is 79.8° further these properties improved for different membranes are 75.5, 69.2, 63.5, 60.3° for PEI/TP, PEI/pTNT, PEI/N1TNT and PEI/Cu1TNT respectively. It is observed that the contact angle of nano metal oxides incorporated PEI membranes decreases, especially with TNT than TP. A large amount of surface hydroxyl groups in pTNT is ascribed due to nanotubular morphology and layered structure as well. Rajesh Kumar et al reported that the contact angle of nascent polysulfone and PSf/chitosan membrane with 3.0% TiO2 NT are 73° and 58° respectively, that is contact angle of 20.5% decreased. Interestingly, in the present study PEI/Cu-TNT showed 20.1% decrease in contact angle without chitosan. Further decrease in contact angle of N-TNT is due to biphasic structure with oxygen defects in crystal structure. The Cu-TNT membrane showed lowest value among the nanotubes due to hydrophilic nature of copper species. Choi et al., reported that the hydrophilic functional groups into the surface of the carbon nanotubes (MWCNTs)/polysulfone blend membranes, thereby generating increased free volume and appropriate free volume cavity size. In general, the large amount of hydroxyl group on the metal oxide surface increases the hydrophilic character of the membrane that is hydrogen bonds between water and the membrane surface. In the case of nanomaterials functionalized membranes, the dimensions of membrane surface pores increase, more water is adsorbed by the membrane during the contact angle experiments. The above results are in-tune with earlier reports.

<table>
<thead>
<tr>
<th>Sl. No</th>
<th>Membranes</th>
<th>Water uptake capacity (%)</th>
<th>Porosity (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>PEI</td>
<td>62.3</td>
<td>14</td>
</tr>
<tr>
<td>2</td>
<td>PEI/TP</td>
<td>70.5</td>
<td>21</td>
</tr>
<tr>
<td>3</td>
<td>PEI/pTNT</td>
<td>74.0</td>
<td>25</td>
</tr>
<tr>
<td>4</td>
<td>PEI/N1TNT</td>
<td>79.0</td>
<td>29</td>
</tr>
<tr>
<td>5</td>
<td>PEI/Cu1TNT</td>
<td>82.2</td>
<td>33</td>
</tr>
</tbody>
</table>

Table 2 displays water permeation capacity and porosity of the membranes. Neat PEI membrane has the least water uptake of 62.3%. The water uptakes of the membrane increased to 70.5, 74.0, 79.0 and 82.2% for PEI membrane with nanoparticles TP, pTNT, N-TNT and Cu-TNT respectively. All the functionalized membranes showed increased in hydrophilic character with increase in porosity of the membranes. It reveals that, the membrane interconnectivity between the sub-layers increases with an increased number of surface pores of the skin layer, alternatively membrane hydraulic resistance decreases resulting in a higher water flux. The above results demonstrated the flux increases due to increased hydrophilicity, surface pore density and pore connectivity. An increase in water uptake with membrane functionalized with carbon nanotube and TNT has been reported.

3.6. Performance of mixed matrix membranes

3.6.1. Pure water flux

Fig. 7 represents pure water flux with time for different membranes. The membranes show a decline in water flux with respect to time when operated under high transmembrane pressure of 500 kPa and this loss in water flux is irreversible for membrane. Compaction is compression of the membrane structure under a high transmembrane pressure difference in which the walls of the pores became closer, denser and uniform.
resulting in reduction in pore size and flux during the compaction. After three hour operation the flux value reached to a steady state value which is higher for PEI membrane with Cu-TNT showed a 0.25 L/m²h steady state water flux. It was witnessed by water uptake study that the water uptake increases with nanoparticle doping and the same trend has been observed in the case of water flux too. Incorporation of nanoparticles into the membrane matrix results more hydrophilic properties, which facilitates the penetration of water into the membrane. As explained earlier for water uptake behavior, the same explanation will justify the water flux character of the membrane. The PEI/Cu-TNT membrane showed high initial and steady state water flux followed by PEI/N-TNT, PEI/pTNT, PEI/TP and neat PEI membranes. Incorporation of TNT nanomaterials into PEI membrane surface promoted the attachment of surface hydroxyl groups. The PEI surface and PEI network have strong interaction with water molecule, which led to attract water molecules by hydrophilicity of the membrane surface. This has beneficially contributed to the enhanced water permeability through the membrane. Zhao et al demonstrated excessive TiO₂ nanoparticles loadings decline hydrophilicity by decrease in specific surface area of aggregated nanoparticles. Hence, we chosen 0.5 wt % of TNT nanomaterial in PEI polymer matrix.

3.6.2. Permeate flux and rejection performance of salt solution

The permeate flux and rejection of salt solutions of K₂SO₄, NaCl and CaCl₂ were studied with concentrations 500, 1000, 1500 and 2000 mg/L at the transmembrane pressure of 500 kPa. The effect of concentration on critical flux of salt solutions is listed in Table 3, 4 and 5. From Tables, it is clear that flux is increasing with decrease in salt concentration for all the membranes. In addition, among the membranes, PEI/Cu-TNT showed higher permeate flux. The TiO₂ nanotube in polymer matrix opened tube entrance, thus induced transport of ions through-hole TiO₂ nanotube array membranes. In addition, MMMs prepared with N-TNT, pTNT and Cu-TNT are expected to enhance hydrophilicity in PEI polymer surface and matrix. The effect of novel functionalized nano-TiO₂ loading electrocatalytic membrane for oily wastewater treatment reported. The electron–hole pairs were generated from TiO₂ to promote the electron transfer, which was attributed to the special function of TiO₂ membrane. They suggested that developed functional membranes would provide a brand-new prospect for water purification.

<table>
<thead>
<tr>
<th>Concentration (mg/L)</th>
<th>Flux (L/m²·h⁻¹)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>PEI</td>
</tr>
<tr>
<td>500</td>
<td>0.111</td>
</tr>
<tr>
<td>1000</td>
<td>0.101</td>
</tr>
<tr>
<td>1500</td>
<td>0.051</td>
</tr>
<tr>
<td>2000</td>
<td>0.019</td>
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</table>

<table>
<thead>
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<th>Concentration (mg/L)</th>
<th>Flux (L/m²·h⁻¹)</th>
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<tbody>
<tr>
<td></td>
<td>PEI</td>
</tr>
<tr>
<td>500</td>
<td>0.111</td>
</tr>
<tr>
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<td>0.092</td>
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<td>1500</td>
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<table>
<thead>
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<th>Concentration (mg/L)</th>
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<tbody>
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<tr>
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<tr>
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<td>0.029</td>
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<tr>
<td>2000</td>
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</tbody>
</table>

The tables above indicate that, for all the salts with lower concentration i.e, 500 ppm shows higher permeate flux and high salt concentration i.e, 2000 ppm shows lesser permeate flux. The
reason for this can explain with the help of salt rejection, which is shown in Fig. 8, 9 and 10. The rejection data also shows that low concentration of salt solution giving higher performance than high salt concentration. This means that actual salt concentration in permeate is lower and therefore the higher rejection. This is because, as concentration increases, the sorption of salt into the membrane increases and ability to reject salt decreases. In addition, PEI/Cu-TNT membrane shows more rejection for all salts followed by PEI/N-TNT, PEI/pTNT, PEI/TP and PEI membranes. The incorporation of hydrophilic TNT inside the membrane matrix encourages the formation of loss structure with a more open pore and porosity, and with enhanced hydrophilicity. Both effects facilitate solute transport through the membrane structure, thus increasing the permeate flux. The higher salt rejections of PEI/Cu-TNT suggest that void size of the polymer when swollen is more important.

Among the salts, potassium sulphate showed more flux and rejection followed by sodium chloride and calcium chloride. This can be explained with the help of ionic size and charge density. The higher rejection of K⁺ ion upto 80% for the membrane PEI/Cu-TNT is because of the higher ionic radius of the K⁺ ion (157 pm) than Na⁺ (116 pm) and Ca²⁺ ions(112 pm). The performance of TNT incorporated membrane depends on permeate flux and salt rejection. It is found that copper doped TNT shows improved performance. The rejection of potassium salt water increased around 15% over the neat PEI membrane at 500 ppm concentration. The enhanced rejection performance of Cu-TNT/PEI membrane is mainly ascribed due to smooth and hydrophilic wall of nano-tubes, which facilitated attraction and rapid movement of water molecules in polymer and nanotube chains. In addition, SO₄²⁻ ion has net negative charge on their surface moiety. Furthermore, the higher radius of ions exhibit relatively better rejection, because these ions can restrictly pass through PEI/Cu-TNT membrane, suggesting that the introduced very small organics through narrow the spacing within Cu functionalized membranes to form new nano networks. In case of membranes with the addition of pristine TNT and functionalized TTNs acquires higher negative charge on membrane surface. This negative charge and the membrane surface negative charge will highly repel and give better sulfate rejection. The K⁺ ion permeate through the negatively charged membrane only with the conjugate SO₄²⁻ ion. In addition, the positive charge of copper doped TNT rejects the positively charged K⁺ ions. Hence, copper functionalized rejects more due to enhance ion selectivity. Similarly experiments with NaCl solution, PEI/Cu-TNT membranes holds higher rejection of up to 75%. The 21.70% of NaCl rejection at 1000 ppm was reported by Rajesh Kumar et al using 1 wt% of TiO₂ nanoparticles incorporated poly(amide-imide) asymmetric nanofiltration membranes. Interestingly, in our study, 3-fold (64%) higher rejection of similar concentration of NaCl solution was obtained using PEI/Cu-TNT membrane. The results are displayed in Fig. 8, 9 and 10. On contrary CaCl₂ rejection was lowered up to 45%, it is mainly because of the small ionic radius of Ca²⁺ ions and less negative charge of Cl⁻ ions. On whole, the membrane performance was ordered as a function of rejection as follows as PEI/Cu-TNT > PEI/N-TNT > PEI/pTNT > PEI/TP > PEI.

Fig. 8 Rejection of Potassium sulphate solution at different concentration

Fig. 9 Rejection of Sodium chloride solution at different concentration

Fig. 10 Rejection of Calcium chloride solution at different concentration
3.6.3. Protein adsorption
The effect of additive nanoparticles on membrane antifouling performance was also investigated through the static egg albumin adsorption test as shown in Fig. 11. It shows that change of egg albumin protein adsorbed onto the titanate nanotubes incorporated PEI membranes. Generally, the neat PEI membrane protein adsorption is higher because of its hydrophobic surface, so that the surface does not have significant ability to hinder the attachment of any kind of foulants. The nanoparticles incorporated membranes showed less egg albumin adsorption than a control membrane. The functionalized titanate nanotubes could be widened to attribute polymer onto a range of oxide type nanotubes. The affinity of PEI to functionalized titanate nanotubes in maintaining stable combinations of Cu1 TNT, NTNT and pTNT in aqueous solution egg albumin. Further, a study of the albumin adsorption on the membrane surface at different pH values was carried out. It can be seen that the highest adsorption occurred at pH 5 and shows a major aggregation of protein on the surface. The reason is that at pH 5 both protein and composite membranes are nonpolar; therefore the hydrophobic interaction causes the albumin to be more easily adsorbed on the surface. At lower pH, cationic species in solution resulted in chain relaxation leading to efficient solvent diffusion. At higher pH, there will be less amount of the protons in solution leads to protein on the surface. The reason is that at pH 5 both protein and composite membranes are nonpolar; therefore the hydrophobic interaction causes the albumin to be more easily adsorbed on the surface.

![Fig. 11 Effect of pH on protein adsorption of neat PEI and MMM](image_url)

4. Conclusions
Hydrogen trititanate nanotubes were synthesized using simple method. It was functionalized with doping process using nitrogen or copper precursors. Mixed matrix membranes (PEI-Titanate) was successfully prepared using titanate and functionalized titanate as additives. The nanotubular shape of pTNT was confirmed from TEM images. The crystal phase of both neat PEI and MMMs were analyzed using XRD pattern. The morphology analysis of MMMs showed macro voids formed after addition of titania based material compared to neat PEI membrane. The standard hydrophilic tests revealed that high hydrophilic nature of Cu-TNT among the membranes. The desalination experiments with different mono and divalent salt solution revealed that PEI/Cu-TNT showed high salt rejection besides enhanced flux and reduced fouling properties. In addition to that, the best membrane also exhibited sufficient stability against biofouling. The PEI/Cu-TNT displayed higher water uptake capacity, hydrophilicity, pure water flux and holds higher performance than neat and PEI/TP and pTNT respectively. Such enhanced salt rejection performance is ascribed due to increase in negative charges by addition of Cu-TNT to PEI, the resulting membrane showed improved charge densities and ions sieving performance.

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References