

Photocatalytic–pervaporation using membranes based on organo-silica for wetland saline water desalination

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Photocatalytic–pervaporation using membranes based on organo-silica for wetland saline water desalination

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This study aims to analyse the morphological structure of organo-silica membranes and investigate their use and performance in a hybrid photocatalytic–pervaporation process for the desalination of wetland saline water. When applied to such raw water sources, this hybrid process – which is shown to have a number of benefits, including excellent water flux and salt rejection, and reduced membrane fouling – has the potential to produce clean water that is often needed in both developing and developed countries worldwide.

Water is essential for life on earth, but searching on a daily basis for sources of clean water can be a challenge.

Wetland saline water potentially could be considered as a usable water source as it is often abundantly available. However, it is essential that its high salt content is removed before it can be used as potable water.

One way of addressing this problem is to use a hybrid process of photocatalytic–pervaporation, based on organo-silica membranes. Researchers at Mangkurat University, South Kalimantan, Indonesia, are studying the morphological structure of organo-silica membranes and investigating their use and performance in a hybrid photocatalytic–pervaporation process for the desalination of wetland saline water, as this article explains.

Hybrid process

In the research carried out, organo-silica membranes were prepared by the sol–gel method, using tetraethyl orthosilicate (TEOS) and a carbon template from banana pectin, at a concentration of 0.5%, which was calcined at 300°C.

The hybrid process of photocatalytic–pervaporation was conducted under vacuum pressure, using an ultraviolet (UV) lamp for illumination and employed titanium dioxide (TiO₂) as a catalyst (at 0.02%, 0.05% and 0.08%) in the feed.

The morphological structures were studied and determined using scanning electron microscopy (SEM) and Brunauer–Emmett–

Teller (BET) analysis, whilst the performance of the membrane was examined by evaluating the water flux and salt rejection.

The results show that the organo-silica membrane has a mesoporous structure, with a top layer thickness of ~2 µm.

The photocatalytic process applied to the desalination of wetland saline resulted in excellent water flux – at 2.7 times higher than without the photocatalytic process (organo-silica 0.5% of pectin). Overall, the membrane possessed excellent water flux, at 9.93 kg·m⁻²·hr⁻¹, and extremely good salt rejection, in excess of 99%.

The combination of both photocatalytic and pervaporation, using an organo-silica membrane, is expected to inhibit the occurrence of fouling and enhance membrane performance during the desalination of wetland saline water.

Concerns worldwide

Nowadays, the distribution of clean water is a major concern for both developed and developing countries.

Wetlands can be used as a raw source of water to fulfil the need for clean water. However, as the tidal mouth or estuary of a large river meets the sea along the coastline, sea-water intrusion into wetland water makes this resource highly saline (3.5 wt%).^[1]

One of the ways in which this water can be used is to treat it by using desalination technology. This membrane-based process has a number of advantages in that it:

- takes place at room temperature;
- can be done continuously and has a varying nature;
- can be adjusted as needed; and
- is environment-friendly.^[2–4]

Organo-silica membranes

Organo-silica membranes, which are widely used for desalination, are made from silica and incorporate carbon in order to increase membrane strength.^[5–10]

According to Elma,^[11–13] the carbon–silica membrane has a large and stable flux for water. Fouling is one of the problems often faced by researchers who are investigating the use of membranes in water treatment. Fouling is caused by the presence of humic or organic material in wetland saline water, which eventually decreases membrane performance.^[14–16]

Photocatalysis

Photocatalysis can be used in purification processes. Because it degrades organic compounds,

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such as humic acid in peat water, it can be used in wastewater treatment.^[17, 18]

Photocatalysis, using TiO₂ as a catalyst, can degrade organic matter and reduce organic material content in peat water by 89.4%.^[19] This indicates the potential of this process in the decomposition of organic materials, found in swamp water, into simpler compounds.

In addition, the use of illumination using UV light can eliminate 99% of pathogenic bacteria and 99% of viruses,^[20] which further improves the quality of the results of wetland saline water treatment.

The aim of this research is to further improve results – enhancing water flux and salt rejection – by combining the photocatalytic use of UV light and the TiO₂ catalyst.

Enhancing membrane performance

The application of a photocatalytic process to a membrane-based treatment enhances membrane performance.^[21]

In this regard, TiO₂ is often used as a photocatalytic catalyst because it is environment-friendly.^[22] Also, using TiO₂ can decrease the activation energy required to accelerate the reaction and can oxidise organic pollutants into carbon dioxide and water.^[23]

The appropriate selection of membrane materials is also an important factor in reducing process costs without compromising the membrane's capability.^[24] Therefore, this research focuses on the performance of organo-silica membranes, in terms of water flux and salt rejection, in combination with UV radiation using photocatalysis with a variation (at 0.02%, 0.05% and 0.08%) in the amount of TiO₂ catalyst used, in order to treat wetland saline water.

Experimental work

Chemicals and materials

A silica-pectin membrane was prepared according to techniques discussed in previous work by Elma, *et al.*^[25]

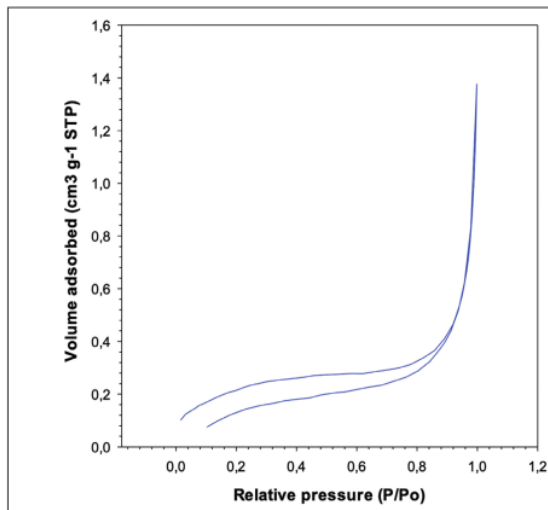


Figure 1. Plot of N₂ sorption isotherms for the calcined xerogels.

A NaCl solution of 3.5 wt% was used as the feed to represent “sea-water” and liquid nitrogen for cooling water vapour, condensing it to a liquid. In addition, an 18 W UV lamp was used in the photocatalytic process.

Photocatalytic-pervaporation

The process of desalination using the method of pervaporation was done with deionised water and wetland saline water.

An initial sample of wetland saline water was characterised by considering total dissolved solids (TDS), UV₂₅₄ absorbance and conductivity. A vaporisation test was then conducted in combination with photocatalysis – the photocatalytic-pervaporation process – using the membrane that had been developed and produced for treating the wetland saline water.

The water flux and salt rejection were measured to determine the photocatalytic effect on the desalination process of the wetland water.

subdistrict Beruntung Baru, Banjar Regency, South Kalimantan, Indonesia.

The pH of the wetland saline water used in this study is relatively neutral, with a pH of 7.82; dissolved organic carbon (DOC) of 16.52 mg.l⁻¹; TDS of 7260 mg.l⁻¹ and conductivity of 17.42 mS.cm⁻¹.

Based on the Indonesia Ministry of Health Regulation No 492 for 2010, setting out the quality requirements of drinking water, the TDS and conductivity should be lower than 700 mg.l⁻¹ and 2.5 mS.cm⁻¹, respectively. The DOC represents the organic content in the wetland saline water.

Xerogel characterisation and membrane morphology

The physical and textural properties of organo-silica xerogels were studied using nitrogen sorption BET analysis, which provides important qualitative data regarding the microstructure of the resulting molecular sieving membranes. A plot of the N₂ sorption isotherms for the calcined xerogels is shown in Figure 1.

According to the IUPAC classification, the isotherm belongs to type IV^[26] which indicates that the surface structure is mesoporous.^[27]

A rapid elevation in the volume of nitrogen adsorbed at high relative pressure and a

Results and discussion

Wetland saline water characteristics

The water sample was collected during the dry season from the village Muara Halayung,

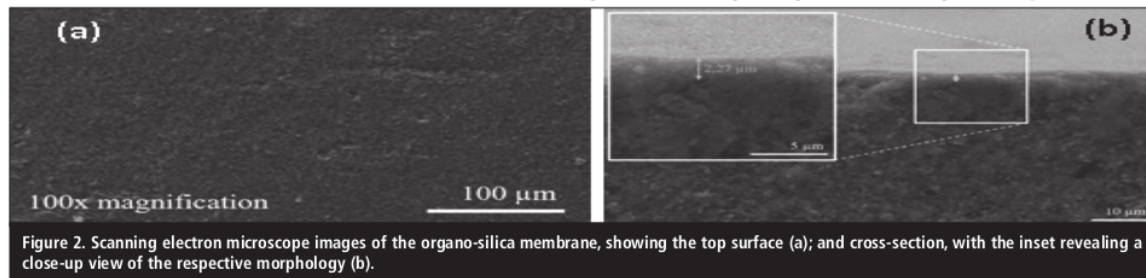


Figure 2. Scanning electron microscope images of the organo-silica membrane, showing the top surface (a); and cross-section, with the inset revealing a close-up view of the respective morphology (b).

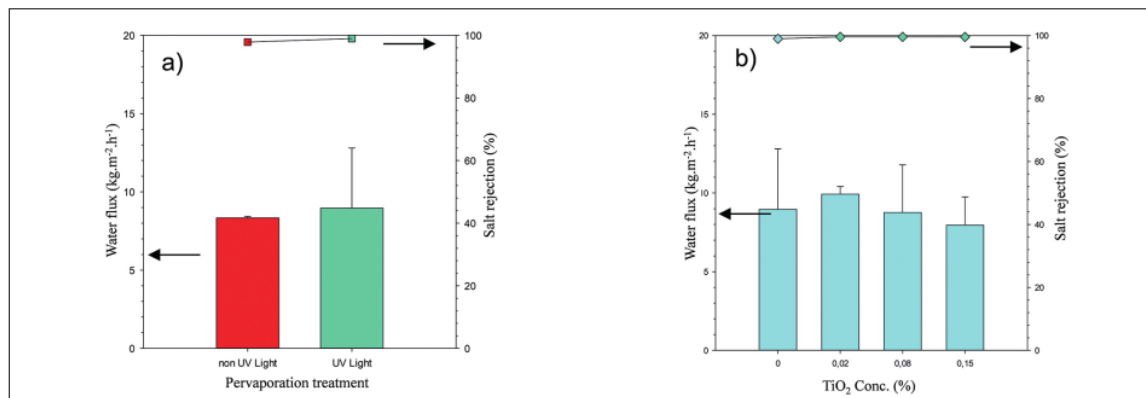


Figure 3. The performance of the organo-silica membrane (0.5% pectin concentration) without the TiO₂ catalyst (a); and photocatalytic-pervaporation membrane performance – with variation in the TiO₂ catalyst – for wetland saline water at 25°C.

hysteresis loop, as found in the present case, are characteristics of mesoporous materials. The prominent characteristic of type-IV isotherms is exhibited with a distinct hysteresis loop of H1 in the P/P₀ range of 0.4–1.0, which implied the presence of relatively large mesopores in the frameworks.^[28]

Figures 2a & b show representative scanning electron microscope (SEM) images of a synthesised organo-silica membrane and the underlying alumina substrate.

The top layer and the macroporous substrate layer, between the cross-section of the membrane, shows that there is a lack of clear distinction.

The SEM image in Figure 2a indicates that there are no cracks in the membrane surface. These would be caused by pore differences between the top layer and membrane support alumina which has a larger pore size of 100 nm.^[29, 30]

Membrane thickness

In addition, differences in both the calcination techniques and temperatures used, affect membrane thickness.

The thickness of the organo-silica membranes can be seen in Figure 2b. The SEM image shows an organo-silica membrane (0.5% pectin) calcined at 300°C and ~2 μm thickness.

Compared with the calcination technique that employs conventional thermal processing (CTP), resulting in a thickness in the region of 30–50 nm,^[31, 32] the thickness of this membrane, prepared by rapid thermal processing (RTP), is thicker.^[2, 3, 29, 33–36]

This is because the application of the RTP technique makes the solvent and water in the silica matrix evaporate completely, whilst they are still trapped in the pores in the CTP calcination process.^[37]

Even so, the RTP technique has the advantage of being able to reduce the costs and time associated with membrane fabrication. In addition, the RTP technique can produce larger clusters of siloxane and membrane pores, and potentially increase membrane hydro-stability and the resulting permeate flux.^[38–40]

Membrane performance

Water flux and salt rejection measurements are presented in Figure 3a.

Referring to Figure 3a, the water flux increases by 6.92%, with the application of UV light and without the addition of the TiO₂ catalyst, compared with the performance of the organo-silica membrane without UV light.

The flux of the non-illuminated membrane, without the reported UV lamp, amounted to 8.96 kg.m⁻².hr⁻¹, compared with flux of 8.34 kg.m⁻².hr⁻¹ achieved in previous studies when a less powerful UV lamp was used.

In addition, in this research, the results show that salt rejection also increased from 97.85% to 98.99%. This increase is caused by the absorption of photon energy through illumination using UV light, which results in an oxidation–reduction reaction. In these cases the radiation results in the process of photodegradation without the addition of the TiO₂ catalyst.^[37, 38]

The addition of the TiO₂ catalyst causes a decrease in the TDS in the feed water, which eventually results in better membrane performance compared with when just UV light is used.

This is achieved through a decrease in the TDS content in wetland saline water by 19.28% (with an initial measurement result of 5400 mg/l), which reduces the content of impurities in the water. These impurities affect the membrane greatly as they can reduce the

available volume in its pores, which results in a decrease in water flux.^[14]

TiO₂ catalyst

Figure 3b shows a comparison between photocatalysis with and without the use of the TiO₂ catalyst. It can be seen that when the catalyst is added there is slight increase and decrease in water flux.

With the addition of TiO₂, at a 0.02% concentration, the water flux increases by 9.77% of the yield without TiO₂ addition, to 8.96 kg.m⁻².hr⁻¹. This is because the TiO₂ catalyst is “charged” with UV light which occurs with the excitation of electrons and causes the interaction.

Interactions that occur form the hydroxyl radical because of the illumination. Such hydroxyl radicals interact with organic matter in the degradation process.^[23]

These radicals will be formed continuously whilst TiO₂ is illuminated with UV light. The radical will “attack” pollutants in the water so that they are degraded.

With the addition of TiO₂, at concentrations of 0.08% and 0.15%, the water flux dropped by a percentage consecutive decrease of 2.23% and 11.16%, respectively.

This is because the catalyst suspension of the solution is too large and blocks the UV light. Consequently, the production of hydroxyl radicals, which are active species in the photodegradation process, will also decrease.

Decreased photodegradation also can be caused by the deactivation of the catalyst's active site as it is covered by other catalytic sites that are in its underlying state (ground state). It also may be because of the sedimentation of catalysts during photodegradation – so that there is a catalyst surface that does not absorb light, or organic compounds in the wetland saline water. In this case the membrane becomes fouled and water flux declines.

In addition, the results show that salt rejection, which is reported in Figure 3, has increased as the concentration of TiO₂ weight increases.

Conclusion

This research has shown that the TiO₂ catalyst has an influence on water flux as well as salt rejection in the photocatalytic-pervaporation process.

The photocatalytic process results in excellent water flux for wetland saline water – at 2.7 times higher than without photocatalytic-pervaporation. The highest photocatalytic membrane performance achieved, with a water flux of 9.93 kg.m⁻².hr⁻¹ and salt rejection of 99.59%, was for the 0.02% addition of the TiO₂ catalyst.

The photocatalytic process reduces membrane fouling during the pervaporation process and enhances the performance of organo-silica membranes used in the desalination of wetland saline water.

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PATENTS

Cross-flow filtration device

Applicant: Sartorius Stedim Lab Ltd

A cross-flow filtration device that is designed to be applied to a pressurised feed liquid forms the subject of this patent. The device comprises a filter membrane and flow channel, for the pressurised feed liquid. The liquid flows in the channel that extends in a path over a retentate surface of the membrane, such that its direction of flow (in the channel) is tangential to the retentate surface, and filtrate derived from the feed liquid passes through the membrane leaving retentate liquid in the flow channel. Also included in this design is a collection chamber for the filtrate. This is formed on an opposite filtrate surface of the membrane. The cross-flow filter further comprises a sealed housing comprising a retentate side and a filtrate side which enclose the flow channel, filter membrane and collection chamber. The device also incorporates flow-channel guide walls that are on an inner surface of the retentate side of the housing. These walls are configured to form a fluid-tight seal with the filter membrane. In

doing so, they define the path of the flow channel over the retentate surface of the membrane.

Patent number: WO/2020/173762

Inventors: A. Gilbert, M. Purshouse, M. Bates, C. Biddell, A. Green and T. Davidson

Publication date: 3 September 2020

Pervaporation apparatus for glycerol concentration

Applicant: Nanjing Weixin Environmental Protection Equipment Technology Research Institute Co Ltd, China

This invention concerns a hollow-fibre pervaporation membrane-based method for concentrating glycerol. It involves a series of steps. In the first, ultrafiltration (UF) is performed on a glycerol crude liquid with a concentration of 20–50% to filter out total suspended solids (TSS), in order to obtain a filtered clear liquid. This clear liquid is heated, whilst its temperature is controlled between 60°C and 100°C. A hollow-fibre pervaporation membrane is then used to dehydrate and concentrate the filtered clear liquid so as to recover 92–95% of the glycerol. The hollow-fibre pervaporation mem-

brane is sequentially formed – from the inside to outside – of an inner support layer; a selected separation layer; and an outer protective layer. The selected separation layer is arranged between the inner support and outer protective layers. The outer protective layer is hydrophilic. Also discussed is an apparatus based on the hollow-fibre pervaporation membrane, which is used for glycerol concentration.

Patent number: WO/2020/173510

Inventors: Y. Tang, J. Liu and W. Wang

Publication date: 3 September 2020

System for performing RO with integrated pump storage

Applicant: Fluid Equipment Development Co LLC, USA

This disclosure relates generally to reverse osmosis (RO) systems, but more specifically to methods and systems for using an elevated reservoir for supplying feed for RO. An RO system is described that includes a first pretreatment unit and a fluid source located below the reservoir. A housing contains an RO membrane, and a first turbocharger includes a first pump and turbine portions. The first pump portion receives feed fluid from the first pre-

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