Recent Progress on Tailoring and Modification of Membranes for Membrane Distillation: A Review

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ABSTRACT

Membrane distillation (MD) has gained the interest of many researchers since it is a promising method for the separation and purification process. Membrane distillation (MD) is a non-isothermal separation process in which differential vapor pressure between porous hydrophobic membrane surfaces acts as a driving factor. A hydrophobic membrane is used in the application of MD, which permits only the passage of vapor produced on the feed side through its pores to the permeate side. One of the most significant obstacles to the commercialisation of the MD method is a lack of appropriate membranes for the process. On the other hand, conventional hydrophobic membranes are subjected to rapid wetting and severe fouling, mainly when low surface tension compounds are present in saline water, resulting in decreased MD performance. In recent decades, MD membranes have received exceptional scientific interest, with substantial progress being made in the design and production of MD membranes appropriate for use in many applications. This review gives a comprehensive overview of recent research developments in the tailoring morphological structure of hydrophobic membranes, emphasising advancements in the fabrication and modification of membranes towards exhibiting high efficiency in the MD process. In addition, the critical morphology characteristics, mainly surface roughness, wettability, and water contact angle, are analysed. Finally, the challenges faced and future research direction is highlighted.

Keywords: Membrane distillation, tailoring, modification, characterisation, recent progress, review

1.0 OVERVIEW ON MEMBRANE DISTILLATION

The rapid growth of metropolitan areas along with an increasing worldwide population are the driving factors that contribute to the demand for sufficient and potable water supplies [1]. According to the global population clock, the global population has surpassed 7 billion and is estimated to rise to 10 billion by 2050 [2–3]. Therefore, a key challenge for the emerging countries of the globe would be the availability of safe drinking water. To overcome the water scarcity issue, enhancing the efficiency and affordability of water treatment is a critical component that needs to be emphasised [4].

1.1 Fundamental on Membrane Distillation

Membrane technology has experienced a significant increase in popularity over the last few years as one of the water treatment methods. The membrane technology has great potential in water treatment due to its
substantial decrease in equipment size, energy consumption, and inexpensive capital cost [5–7]. Therefore, membrane distillation, which was first presented six decades ago, can be widely employed in commercial applications [8,9], although it is still in the research stages [8–9]. Membrane distillation is a thermally-driven separation process that comprises a porous hydrophobic membrane that allows the vapors produced by the feed liquid to flow through to the permeate side [9–11]. Figure 1 illustrates the membrane distillation process. The hydrophobic membrane is the critical component in the separation process. It prepares the medium for the vapor generated on the feed side to penetrate it and act as a barrier to prevent the liquid water from being transferred directly to the other side [12–14]. A variety of configurations with distinct thermal efficiency and mass transfer trade-offs can be a reference to assist membrane distillation and induce the vapor pressure difference across the membrane [11, 15]. The separation performance and cost of operation are greatly affected by the MD configuration. Numerous research teams have proposed innovative designs with increased energy efficiency, better permeate flux, or a reduced footprint [16–18]. These configurations include direct contact membrane distillation (DCMD), air gap membrane distillation (AGMD), vacuum membrane distillation (VMD), and sweep gas membrane distillation (SGMD). In DCMD, the permeate side of the membrane is in contact with the cooled draw aqueous solution, which causes the mass transfer to be enhanced, resulting in greater productivity [15, 19]. DCMD is the most simplified arrangement and has been extensively explored in the literature and laboratories for desalination and concentration of aqueous solutions [14,20–22].

For AGMD, a thin air gap is created between the membrane and the condensation surface. The membranes and the air gap allow the evaporated volatile molecules to flow through and then condense on the cool surface [25]. Typical AGMD systems have lower fluxes than DCMD or VMD configurations due to the high vapor transport resistance provided by the air gap [14, 24]. With VMD configuration, the permeate side of the membrane module is subjected to vacuum pressure [17]. The applied vacuum is lower than the saturation pressure of volatile molecules in the feed solution to produce the necessary driving force [25, 26]. In SGMD, a cold, inert gas, also known as a sweep gas, is drawn over the permeate channel, collecting vapour molecules accumulated on the membrane's surface as it moves along [27, 28]. Membranes with low mass transfer resistance and high heat transfer resistance are preferable for increasing the driving force in MD and, subsequently, achieving a high permeate flux [31]. The performance of the membrane varies greatly depending on the inherent characteristics of membrane materials [32]. The surface roughness, wettability, and water contact angle are several characteristics that describe the morphological structure of a membrane used in membrane distillation. The water contact angle and wettability of the membrane are known to be correlated with the roughness parameter. In addition, the measure of the static water contact angle is increased as the surface is roughened [31, 32].
1.2 Issues on Membrane Distillation

In membrane distillation, the membrane is susceptible to fouling and wetting issues, both of which can negatively influence the system's overall performance. Dissolving low surface tension compounds and organic components in feed water is the primary cause of wetting, resulting in decreased water output [36]. Therefore, numerous global scientific research has stressed the necessity for more selective membrane fabrication to overcome these constraints. Many efforts have been made to develop a superhydrophobic membrane by utilising low-surface energy materials and constructing hierarchical microstructures [37]. In long-term MD operation, membranes with a superhydrophobic surface tend to have better flux and lesser flux reduction than membranes with a hydrophobic surface. This is due to the fact that they have significantly stronger wetting resistance and self-cleaning characteristics [38]. Although the superhydrophobic membranes have increased the wetting resistance considerably, the membrane's intact oleophilic features still reduce the fouling resistance due to the hydrophobic-hydrophobic interaction between the membranes and organic foulants in the feed water [37, 38].

Apart from the superhydrophobic membrane, an omniphobic membrane characterised as water, and low-surface tension agent repellent has been developed and applied in the membrane distillation process. However, omniphobic membranes are still susceptible to fouling issues as a result of hydrophobic foulants such as oil droplets on the membrane surface [8, 17]. The research discovered that a membrane with a unique wettability feature was the most effective fouling and wetting resistance improvement [36, 40]. As a result, a membrane known as the Janus membrane with an asymmetric wettability has been developed and designed to address both membrane wetting and fouling problems concurrently. In addition, oleophobicity underwater may resist oil droplets and prevent oil fouling in the Janus membrane, which contains a thin hydrophilic layer that is hydrated with water. Thus, there is a lot of innovation has been made to improve membrane distillation performance. In this paper, we aim to review the recent development on the morphological structure of the membrane for efficient membrane distillation.
2.0 FABRICATION OF MEMBRANES FOR MEMBRANE DISTILLATION

For the past three decades, a commercially available membrane precisely optimised for MD operations has yet to be developed [19–21]. This is primarily because most of the early works focused solely on the theoretical analysis of the MD process. However, there have been significant advancements in fabrication technology in recent years, developing new membranes with enhanced surface wettability that may be used in the MD process. The fabrication of MD membranes may be carried out using a variety of operations such as sintering, phase inversion, stretching, template leaching, electrospinning, and track-etching. The chosen method for the fabrication of membrane depends on the materials and the design shape of the membrane [46].

2.1 Polymeric Membranes

A polymeric membrane is an organic membrane synthesised from an organic synthetic polymer that possesses hydrophobic surfaces such as polyvinylidene fluoride (PVDF), polytetrafluoroethylene (PTFE), and polypropylene (PP). This membrane is mainly utilised in the membrane distillation process as it offers a feature such as versatility and low thermal conductivity (0.1–0.5 W·m⁻¹·K⁻¹) [47]. For the fabrication of polymeric membranes, the most widely utilised methods include phase inversion, interfacial polymerisation, stretching, track-etching, and electrospinning [4, 33, 48–56]. Phase inversion is the most widely used method to prepare a membrane for the membrane distillation process. In this method, the mixing process converts the liquid phase homogenous solution to the solid phase. There are various types of phase inversion techniques, including Non-solvent induced phase separation (NIPS), Thermally induced phase separation (TIPS), and vapor induced phase separation (VIPS). However, due to the mild processing conditions and the easiness with which membrane shape can be controlled, NIPS is the most often used phase inversion technique for producing hydrophobic membranes [32]. Polymer, solvent, and non-solvent are the crucial components in the NIPS process [57]. The process starts with forming an initial homogenous solution by dissolving a polymer in its solvent. Then, the phase separation occurs where the solvent of the homogenous solution is exchanged into the non-solvent resulting in a polymer residue that forms the membrane [58]. Nejati et al. [59] synthesised a high-performance hydrophobic membrane using a PVDF dissolved in triethyl phosphate (TEP) as a casting solution. The development of a hydrophobic membrane with an asymmetric structure is the consequence of the non-solvent-induced phase separation of polyvinylidene fluoride (PVDF) in triethyl phosphate (TEP). The high surface porosity and roughness of the top surface attributed to the increase in the concentration of a soft coagulation agent (2-propanol) in the coagulation bath was observed since the roughness of the top surface of the membranes was significantly dependent on the amount of 2-propanol present. As a result of the considerable increase in surface roughness, the surface contact angle has also increased considerably. As a result, the performance of the MD membranes in terms of water vapor flow varied substantially depending on the orientation of the membranes.

It was discovered that the membrane porosity and thickness substantially impacted the water vapor
flow. When the membrane thickness was lowered while the porosity was increased, a rise in vapor flow was observed. Furthermore, due to the stabilisation of airgaps at the liquid-solid interface, hydrophobicity increases. Therefore, increasing the stability of the air gaps will improve MD performance by decreasing the likelihood of partial wetting of the membranes. Munirasu et al. [60] have synthesised a highly porous and intrinsically superhydrophobic PVDF membrane by utilising a phase inversion method with alcohols including methanol and ethanol as the non-solvent on non-woven support. From the characterisation of the membrane, the surface and cross-sectional SEM analyses of the PVDF membrane revealed that the morphology of the membrane is a sponge-like interconnected fibrous structure across the membrane with no selective top layer.

In comparison to the PVDF membrane on a support, the superhydrophobic PVDF membrane demonstrated comparatively high flux with good salt rejection, and the long-term MD measurements over a time span of more than a week revealed a steady flux after initial stabilisation. Additionally, the water contact angle of PVDF membranes produced using methanol as the non-solvent is around 164°. Because of the lack of a selective top layer and the presence of a sponge-like linked fibrous structure throughout, it is a promising material for studying surface wetting/nonwetting behavior. Another study by Thomas et al. [61] has fabricated PVDF membrane with sponge-like structures. The structure was achieved due to the delay in the rate of liquid-liquid de-mixing of the polymer, where the immersion time was strictly controlled during the coagulation step. The membrane was fabricated via the NIPS techniques without pore-forming additives for the MD application. Table 1 summarises the fabrication techniques for polymeric membranes.

<table>
<thead>
<tr>
<th>Polymer</th>
<th>Fabrication</th>
<th>Water contact angle (°)</th>
<th>Application and MD configuration</th>
<th>Ref.</th>
</tr>
</thead>
<tbody>
<tr>
<td>PVDF</td>
<td>NIPS</td>
<td>125 ± 2</td>
<td>MD/DCMD</td>
<td>[53]</td>
</tr>
<tr>
<td>PVDF</td>
<td>Phase inversion</td>
<td>164</td>
<td>MD/DCMD</td>
<td>[54]</td>
</tr>
<tr>
<td>PVDF</td>
<td>NIPS</td>
<td>100-127</td>
<td>MD</td>
<td>[55]</td>
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2.2 Ceramic Membranes

Since the beginning of the previous decade, ceramic membranes have received growing interest in various separation applications, including industrial wastewater and drinking water treatment, desalination, and gas separation [62]. Due to their superior mechanical, chemical, and thermal stability, the ceramic membrane is more resilient toward severe operating conditions than polymeric membranes [57, 58]. Phase inversion and sintering have recently been utilised to fabricate various ceramic membranes, including flat sheets [59, 60], hollow fiber [61–65], and tubular [62]. It has been demonstrated that the combined phase inversion and sintering method is a promising method for the production of ceramic membranes compared to traditional methods [72]. In this method, the asymmetric ceramic membranes are produced in a single fabrication step as only one heat
treatment is necessary [73]. This approach involves preparing a suspension of ceramic particles, an organic solvent, a polymer binder, and water. Due to the phase inversion caused by the exchange between solvent and non-solvent, the ceramic particles are immobilised through spinning or casting, depending on the shape. Finally, all organics are removed from the membrane precursors, and the membrane's mechanical properties are improved [74]. Currently, alumina is employed in the majority of commercial ceramic membranes. However, high sintering temperatures (about 1500°C) are required to provide appropriate porosity and mechanical strength in these materials. The requirement for pure alumina powder and a high sintering temperature leads to increased manufacturing costs, which is the most significant disadvantage of ceramic membrane technology. However, a great effort of research has been made on developing a low-cost hollow fiber ceramic membrane made of natural mineral powders and other inorganic materials [75].

A low-cost zeolite-based Horizontal Fuel Conditioning Module (HFCM) has been fabricated by Adam et al. [76] using a phase inversion-based sintering technique. Figure 2 depicts the impact of sintering temperature on the morphology behavior of the membrane. It has been shown that a highly porous membrane with a loosely packed fine structure was obtained at the lowest sintering temperatures compared to other sintering temperatures. As the temperature rises, the structure of HFCM becomes less porous, and the grains are densely packed. In terms of the contact angle, the contact angle was increased from less than 5° to more than 50° due to the dehydration of the ceramic membrane. The membrane surface shows superhydrophobic features at a low temperature which was attributed to the presence of OH groups [77]. As the temperature rises, the OH group is dehydrated resulted in more closely packed ceramic particles. In this condition, the ceramic membrane possesses a hydrophobic characteristic.

Figure 2 Cross-sectional SEM images of fibre 7 sintered at different temperature; (a) 900°C, (b) 950°C, (c) 1000°C, (d) 1050°C, (e) 1100°C, and (f) 1150°C; (1) overall cross-section, (2) zoom at the voids [70]
Another study of ceramic membrane fabrication has been accomplished by Bandar et al. [64]. This study fabricated a compelling novel membrane by integrating an inorganic substance (red clay) with liquid TEOS, ammonia, and sodium alginate powder to produce a porous distribution inside the red clay membrane. The average pore size of the constructed membrane was 33 nm, indicating that ultra-membrane distillation was accomplished without the need for a coating layer. Continuous water desalination experiments revealed that the membrane maintained its performance for four h, with the 16.5% flux decrement after 10 hours attributed to fouling caused by the presence of salts and suspended particles in the solution. The percentages of salt ions deposited on the membrane by well water were determined to be greater than 80%, and fouling deposits were easily eliminated by flushing with deionised water (DI) water. The membrane's initial performance (water flow of 30.13 kg/m²h) and water conductivity of 1 S/cm was recovered after a three h VMD test with DI water. In recent studies by Peng et al. [78], the ceramic membrane with anti-fouling PTFE coating was fabricated by vacuum filtration method. In this work, the outer surface of an alumina membrane substrate was disseminated, coated, and thermally sintered by using PTFE nanoparticles to form a resilient defect-free layer with hydrophobicity and a low friction coefficient. The PTFE coating acted as a protective barrier, reducing the adhesion of foulants due to their low surface energy, smooth surface, and low friction coefficient. Table 2 summarises the fabrication techniques for ceramic membranes.

<table>
<thead>
<tr>
<th>Polymer</th>
<th>Fabrication</th>
<th>Water contact angle (°)</th>
<th>Application and MD configuration</th>
<th>Ref.</th>
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</thead>
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<tr>
<td>Zeolite-based</td>
<td>Phase inversion-sintering</td>
<td>-</td>
<td>MD/DCMD</td>
<td>[70]</td>
</tr>
<tr>
<td>HFCM</td>
<td>Extrusion</td>
<td>-</td>
<td>MD/VMD</td>
<td>[58]</td>
</tr>
<tr>
<td>PVDF</td>
<td>Vacuum filtration</td>
<td>129 ± 1.5</td>
<td>-</td>
<td>[72]</td>
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</table>

### 2.3 Composite Membranes

Several hydrophobic surface modification techniques, including surface coating, grafting, mixing, and copolymerisation, can remove or decrease membrane wetting, a well-known MD process difficulty caused by a lack of hydrophobicity [79]. Among all of these techniques, coating the membrane surface with a hydrophobic layer is an efficient method of fabricating a composite membrane due to the simplicity of the procedure, low cost, and large volume of manufacturing. For example, Pan et al. [80] have successfully fabricated a composite membrane by using hydrophilic PVDF membrane as the supporting membrane. In this work, the PFPE membrane was coated on the supporting membrane by the dip-coating method. Based on the characterisation of the membrane, it has been shown that a coating layer has been applied to a substrate membrane and effectively cured without altering the original porosity structure or blocking the pores [81]. More importantly, the anti-wetting
characteristics were significantly improved due to the addition of the coating layer. Micrometer or nanoscale fibers can be created via electrospinning. This promising and flexible technique uses a strong electric field to spin a polymer or biopolymer liquid solution into fibers of micrometer or nanometer size. Some researchers have used electrospun fibrous membranes in the MD process, and they have found that they have excellent performance on permeate flux and salt rejection [76, 77]. The excellent performance of the electrospun fibrous membrane is due to its high hydrophobicity, high porosity, low thickness, and interconnected open structure [84]. Li et al. [85] have fabricated a nanofibrous composite membrane by electrospinning method. This research employed two types of spacers and two non-woven fabrics as supports in the electrospinning fabrication process. Layers of PVDF nanofibers were linked openly in three dimensions, resulting in a rough surface on the composite membranes. The membrane surface developed a multilayer re-entrant structure due to the overlapping nanofibers and the presence of beads-on-string, and all contact angles of the electrospun membranes were over 140°. The electrospun composite membranes were evaluated by DCMD utilising a 3.5% NaCl solution as feed, and the permeate flow reached a maximum of 49.3 kg/m²/hr for the highest permeate flux. The long-term and repetitive operation of MD desalination revealed that the manufactured membrane exhibited excellent scaling and wetting resistance and was easy to clean. Therefore, it is considered that the electrospun fibrous PVDF/spacer composite membrane has a great deal of promise for use in the medical device manufacturing process. The electrospinning method has been utilised by Li et al. [86] to fabricate a triple-layer membrane with outstanding anti-wetting performance. The membrane are consisted of hydrophobic PTFE-PVDF layer, PET support layer, and a chitosan-polyethylene oxide (CS-PEO) hydrophilic layer. Table 3 summarises the fabrication techniques for composite membrane.

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<tr>
<th>Polymer</th>
<th>Fabrication</th>
<th>Water contact angle (°)</th>
<th>Application and MD configuration</th>
<th>Ref.</th>
</tr>
</thead>
<tbody>
<tr>
<td>PVDF</td>
<td>Dip-coating</td>
<td>-</td>
<td>MD/VMD</td>
<td>[74]</td>
</tr>
<tr>
<td>PVDF</td>
<td>Electrospun</td>
<td>&gt;140°</td>
<td>MD/DCMD</td>
<td>[79]</td>
</tr>
<tr>
<td>PVDF-PTFE</td>
<td>Electrospinning</td>
<td>144</td>
<td>MD/DCMD</td>
<td>[80]</td>
</tr>
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</table>

### 3.0 MODIFICATIONS ON MEMBRANE STRUCTURES

A myriad of researches has been studied and concentrated on the membrane’s surface modification to enhance the membrane properties, especially on physicochemical and thermal properties of the membranes. The surface modification has conventionally been accomplished by inserting desired functional group into the surface that may then be used to modify the membrane surface characteristics, particularly in wettability, surface roughness, and water contact angle [87]. Membranes with specific wettability characteristics have been proven to be the most effective for improving fouling and
wetting resistance [12, 39, 40] when compared to other membranes. With inspiration from nature, the special wettability on MD membranes has been achieved by adjusting surface roughness [88, 89], and chemical compositions via surface engineering [90].

Phase separation is shown to be a cost-effective and easy method of increasing surface roughness without using low surface energy modifiers to improve the hydrophobicity of the membrane. Rabiei et al. [91] have successfully enhanced the membrane surface's hydrophobicity through the phase separation of the PC solution on the PVDF membrane. Figure 3 present the SEM images of the membranes coated with different concentrations of the PC solution. Based on this figure, increasing the concentration of PC solutions has gradually covered the membrane surface with polycarbonate deposits, resulting in the hierarchical roughness on the surface of the pristine membrane. In addition, the coated membrane showed an increase in the water contact angle from 82° to 130° with the highest concentration of PC solution. Thus, this study shows that manipulating the surface roughness through the phase separation method has successfully improved the hydrophobicity of the membrane.

![Figure 3](image_url) The SEM images of the surface and cross-section of the membranes coated with (A), 25 mg/mL; (B), 50 mg/mL; and (C), 100 mg/mL PC solution [85]

Nanoparticles and nanofibers are currently being used to modify MD membranes due to advancements in nanotechnological research. Nanoparticles have been included to improve the superhydrophobicity of the commercial membrane by enhancing the surface roughness and generating a multilayer structure on the membrane surface [35, 36]. Recent studies have demonstrated the modification of membranes by applying metallic nanoparticles. For example, Deka et al. [92] have successfully developed a modified PVDF-HFP membrane with zinc oxide, ZnO nanoparticles to improve the hydrophobic character of the membrane. In the presence of well-dispersed fluorinated ZnO NPs, hierarchical micro/nanoscale spherical structures were produced on the
micro/nanoscale (average diameter of 1.0 mm) morphology of the re-entrant membrane. Furthermore, the nanoparticles’ incorporation has formed the rough surface, which develops the air pocket on the membrane [93], allowing improper contact between water and the membrane surface. Due to the impact of combination re-entrant structures and the air pockets, the membrane attained superhydrophobicity while also exhibiting omniphobic characteristics.

Another study by Razmjao et al. [94] has utilised titanium dioxide, TiO$_2$, to modify the PVDF membrane. The TiO$_2$ has been coated on the PVDF membrane by a dip-coating method. Due to its simplicity and low overall cost of production [95], dip coating is a widely used coating technology compared to other alternative coating techniques such as spin coating, spray coating, or plasma treatments [96]. 1H, 1H, 2H, 2H-perfluoro-dodecyl trichlorosilane (FTCS) was employed to fluorinate the coated PVDF membrane. In this study, it has been observed that the modified FTCS–TiO$_2$–PVDF membrane is extremely hydrophobic due to the increase in water contact angle from the commercial PVDF membrane ($125\degree \pm 1\degree$) to FTCS–TiO$_2$–PVDF membrane ($163\degree \pm 3\degree$). The increase in water contact angle has resulted from the substantial reduction in surface free energy and increase in roughness reported for FTCS–PVDF membranes. Recent studies by Zhang et al. [97] presented the application of polydopamine (PDA) and poly(MPC-co-2-aminoethyl methacrylate hydrochloride) (MPC-co-AEMA) for the modification of PVDF hollow fibre membrane. The co-deposition of PDA/MPC-co-easy AEMA’s on the membrane’s outer surface dramatically changed the surface’s chemical and physical properties. The modification has been successfully shown to improve the membrane resistance towards fouling and scaling issues during the MD process. In this work, with increasing co-deposition time, the porosity, pore size, and roughness of the membrane surface decreased while the hydrophilicity of the membrane increased. The bulk structure of the membrane remained essentially the same throughout the experiment. The effect of the PTFE additive on the modification of the PVDF-CTFE layer has been investigated by Ray et al. [98]. The PVDF-CTFE layer was changed by using PTFE mixing and PFS polymerisation to improve its performance and hydrophobicity. It has been proven that adding 10 wt% PTFE to the PVDF-CTFE layer changed the pore structure. The hydrophobicity was increased by changing the PFS, as demonstrated by the contact angle and LEP values of the modified PFS. When PTFE and PFS are blended, it is possible to achieve a high flux by modifying the pore structure. The hydrophobicity of the membrane is increased by polymerisation of PFS, which allows the modified membrane to be employed for preventing wetting phenomena during the VMD process.

4.0 CHARACTERISATION AND PERFORMANCE OF MEMBRANE DISTILLATION

In applying the separation process by membrane distillation, membrane morphology is a crucial component as it is closely related to membrane performances. Therefore, the characterisation of such membranes is necessary to better understand their performance and build membranes designed explicitly for the MD process [99]. Many approaches are available for determining the morphological
characteristics of interest, starting with microscopy, such as scanning electron microscopy (SEM), atomic force microscopy (AFM), and water contact angle.

SEM is one of the most critical techniques for membrane characterisation since it provides information on the morphology and topography of the membrane. Furthermore, it also can be used to determine the pore size of a porous membrane [100]. In recent studies, M. Tian et al. [101] have utilised SEM to investigate the morphology structure of the isotropic PVDF membrane prepared by one step-fabrication. The porous PVDF membranes used in this study were created utilising a simple bottom-up technique that relied on water as a coagulation bath. The isotropic skinless PVDF structure was fabricated in the water bath by coating a non-woven substrate with the developing film and immersing it in the bath. Figures 4 and 5 illustrated the SEM images of the membrane morphologies from the cross-section and top surface of the membranes, respectively. The figures revealed that the membrane surface morphology via the bottom-up method differs from the typical phase inversion method. The membrane forms a rough top surface and sponge-like granular membranes consisting of spherulites.

Figure 4 The SEM image of the cross-section of the normal casting and bottom-up casting membrane [95]
In comparison to other microscopic technologies like SEM and AFM provides more detailed information on surface morphology and a three-dimensional (3D) perspective of membrane topography without the need for sample preparation [99]. Figure 6 depicts the AFM images of the PVDF membrane. The surface roughness of the PVDF-N membrane increases dramatically from 0.0718 μm to 0.375 μm when the bottom-up membrane is used with the same concentration of the PVDF dope solution as the PVDF-N membrane due to the formation of a particulate surface. The effective production of highly hydrophobic PVDF membranes employing a water bath for phase inversion is confirmed by a rapid increase in water contact angles from conventional casting membranes (71.7°) to the bottom-up membrane (144°). This simple, low-impact technique opens the door to mass-producing high-performance MD membranes on a massive scale.
The characterisation by AFM technique was conducted by Shirazi et al. [99] to characterise three hydrophobic membranes (PTFE, PP, and PVDF), which are often employed in various MD applications. Their pore size and pore size distribution, surface roughness, and hydrophobicity were measured to characterise the membranes. The fabrication technique can directly impact the pore size and pore size distribution and the surface topography of the material. The membranes with a size of 0.22 μm are generally explored as the optimum choice for different MD applications [102]. The SEM pictures of the membranes examined in this study observed a completely distinct surface topography and morphology and a non-circular pore structure in each of the membranes studied. With the help of the AFM technique, it was possible to determine the pore size and pore size distribution of these membranes, as indicated in Figure 7. From the AFM observation, the pore size of PVDF, PP, and PTFE membranes have values of 0.311 μm, 0.282 μm, and 0.278 μm, respectively.

Both the intrinsic hydrophobicity of the polymer and the surface roughness of the polymer are essential factors in determining hydrophobicity. Surface roughness can be used to indicate surface irregularities that occur during the production process of the material. Roughness statistics derived from experimental analysis such as AFM provide quantitative information about the shape of membranes [97, 98]. In general, the attainment of multi-scale surface structures increases the hydrophobicity of the membrane surfaces [105, 106]. This study investigated the inherent hydrophobicity of polymers such as PTFE, polypropylene, and PVDF by measuring the surface contact angle of their flat sheets. For PTFE, PP, and PVDF sheets, the obtained values were 120.7°, 103.2°, and 90.5°, respectively, with PTFE being the highest and PP being the lowest. AFM may also be used to measure the roughness of a practical surface directly. AFM analysis offers about 15 distinct roughness characteristics, which may be utilised for topographical investigations and surface energy measurements on various materials.

5.0 CONCLUSION AND FUTURE PERSPECTIVES

In this review, the membrane's fabrication, modification, characterisation, and morphological structure have all been addressed. It has been shown that the morphological characteristics of the membrane will affect the performance of the MD separation process. The surface roughness, wettability, and water contact angle are several characteristics that describe the morphological structure of a membrane used in membrane distillation that has been analysed in this review. Through the employment of various fabrication methods, including phase inversion, sintering, extrusion, and electrospinning accompanied with the modification of the surface, membranes with superhydrophilicity, improved MD performance, anti-wetting, and anti-fouling features have been developed and tested. In addition, a better in-depth understanding of the membrane materials best suited for the MD process can be observed through various characterisation techniques such as SEM, AFM, and water contact angle measurement. Tremendous efforts had been made to develop a suitable membrane for an effective membrane distillation process. However, despite the vast progress that has been in MD membrane research
and technology, there are still some significant obstacles that need to be addressed immediately. For example, there is a lack of membranes with high wetting resistance and permeation flux for treating wastewater containing high salinity and high concentration of low surface tension compounds such as surfactants or oil. To overcome these difficulties, an immense number of works is vital to enhance the fabrication of membranes with better characteristics and properties for efficient membrane distillation performances.

![AFM and SEM images of PTFE, PP, and PVDF membranes.](image)

**Figure 5** The AFM (right) and SEM (left) images of (A) PTFE, (B) PP, and (C) PVDF membranes with 0.2μm reported pore size[93]

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