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Challenges and mitigation strategies for general failure and degradation in polymer electrolyte membrane-based fuel cells and electrolysers

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Green hydrogen production and utilisation technologies employing the polymer electrolyte membrane (PEM) electrolytic cells (both fuel cells and electrolysers) are being deployed to decarbonise the various

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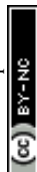
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Dr Malavika Arun is a polymer technologist working predominantly in the field of polymer degradation and combustion. Her recent research activities are centred on chemically modifying styrenic and acrylic thermoplastics with a view to increasing their flame retardance. In addition, recently she is also involved in the syntheses and characterization of functional polymeric resins which have potential applications as safer and high-performance membranes for electrochemical cells.

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Dr Sarbjit Giddey is a Senior Principal Research Scientist and currently Group Leader in the Energy Technologies Program at CSIRO. With over 20 years of research and development experience in hydrogen-related technologies, Dr Giddey's contributions have been instrumental in advancing renewable fuels production, utilization, and battery recycling technologies. As the leader of a dynamic group with diverse expertise, he drives ground-breaking R&D initiatives to enhance energy conversion efficiencies and promote the economic viability of emerging zero-emission technologies. Dr Giddey's pivotal role in the development of CSIRO's Electrolyser and fuel cell technologies has been instrumental in the establishment of the prestigious Centre for Hybrid Energy Systems (CHES) in Melbourne. His visionary approach to technology commercialization has led to the successful spin-off of two companies (Endua – https:// www.endua.com/ and Hadean Energy – https:// hadeanenergy.com.au/) in the electrolysis (hydrogen production) space, involving various industrial and commercial partners. Beyond his research endeavours, Dr Giddey is a recognized authority in the hydrogen and fuel cell sector and sits on the Standards Australia committee, contributing to the adaptation of industry standards in this domain. His immense knowledge and expertise have been shared with the scientific community through over 100 refereed publications and several book chapters, technical reports, patents, and provisional applications. The impact of his research is evident, with over 7600 citations to his name, underlining the far-reaching influence of his work.

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energy sectors. PEM fuel cells convert the green hydrogen to electrical energy with water as a byproduct, whereas PEM electrolysers produce green hydrogen by splitting the water molecules. However, at present, the PEM-based fuel cells and electrolysers possess some inherent issues stemming from the failures and degradation of the cell components. This review presents the challenges and mitigation strategies for general cell failure and the degradation mechanisms of perfluorinated PEM-based electrolytic cells. The review begins with an overview of the fundamental principles of PEM electrolytic cells, followed by a detailed discussion of fluorinated membranes, including their associated challenges and environmental concerns. The review then examines common cell failures, addressing catalytic system degradation, water management issues, ohmic losses, and other component degradation. A subsequent section focuses on PEM failures, analysing the mechanisms and pathways that lead to defects such as pinholes, hotspots, cracking, thinning, and other structural deteriorations. Building on this analysis, we propose several mitigation strategies to enhance the performance and durability of PEM fuel cell and electrolyser components. We then recommend these strategies to improve overall PEM fuel cell and electrolyser system performance and offer perspectives on future pathways for commercialising these technologies.

1. Introduction

Scientists and engineers, worldwide, are actively pursuing the generation and storage of renewable energy through environmentally benign routes with a view to addressing the depletion of and also the emissions from fossil fuels, which currently dominate power generation. Recognising the desirable attributes, such as environmental friendliness, sustainability, and abundance of renewable energy sources, researchers are focusing more on overcoming challenges related to their erratic nature. The storage of energy from renewable sources is a critical issue due to fluctuations in production that often do not go in hand with demand. One proposed solution is hydrogen production by water electrolysis, and its storage to counter the inherent intermittent nature of renewable sources like wind, wave, solar, etc. Developing efficient and cost-effective polymer electrolyte membranes (PEMs) for water electrolysis technologies[1] is crucial for harnessing the high energy density of hydrogen and ensuring reliable utilization of renewable sources to generate electricity to meet demand cycles. Advantages of PEM water electrolysers include enhanced hydrogen-production capacity, purity, and efficiency at higher current densities while operating at low temperatures (<100 °C). For higher-temperature applications in electrolysers, PEM cannot be used; instead, a solid oxide electrolyte is used in the range of 800 to 1000 °C.[2] Hydrogen fuel cells can also serve as emission-

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Dr Dattatray S. Dhawale is a Senior Research Scientist in the Energy Business Unit at the Commonwealth Scientific and Industrial Research Organization (CSIRO) Australia. Dr Dhawale is a prolific, well-cited, and multiple-fellowship-winning scientist. Dr Dhawale's achievements have been honored by several prestigious fellowships, such as Brain Korea-21 (Hanyang University, South Korea), MANA NIMS Japan, Lindau Nobel Laureate Meeting, Germany, etc. Dr Dhawale worked on several multidisciplinary projects at NIMS, Japan; University of Queensland, Australia; KAUST, Saudi Arabia; and QEERI, Qatar. His research interest encompasses the development of efficient nanomaterials for hydrogen and ammonia production by water electrolysis, fuel cells, solar energy conversion, energy storage, and sensors. Dr Dhawale has published over 80 peer-reviewed journal articles, which have attracted more than 6400 citations with an 'h'-index of 44 and an i10 index of 65, four patents, technical reports, and several book chapters. In addition, he mentors postdoctoral fellows and supervises PhD students through university and industry collaborations.

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free energy converters with immense potential, especially in addressing environmental implications, as they can be considered essentially carbon neutral during their operational cycles. Using renewable sources for hydrogen production can also make the entire process emission-free, thus offering a significant opportunity to reduce CO_2 emissions in various industrial processes.[3]

General Electric pioneered PEM water electrolysis in the 1960s as a replacement for alkaline electrolysis, with Grubb[4] conceptualising a solid polymeric material known as the proton exchange membrane (PEM) in electrolytic cells. The PEM facilitates proton transport, minimizing gas crossover and enabling high-pressure operation. These characteristics also contribute to lower operational costs and overall device expenses. Furthermore, the thin membrane reduces ohmic losses, improves protonic conductivity, and offers higher current density in PEM-mediated water electrolysis.[3] In addition, PEM-mediated electrolysis offers an environmentally friendly option for generating hydrogen and integrates seamlessly with renewable sources like wind and solar energy. In addition to the limited exploration in the past century due to low demand for electrolytically produced hydrogen, PEM electrolysis faces numerous unexplored challenges. During the 1980s, there was a rise in interest in producing hydrogen on a large scale through water electrolysis with PEM technology. However, the emphasis of this research was primarily on smaller units producing hydrogen, mainly for generating oxygen in space and underwater environments. The growing demand for sustainable energy has sparked renewed interest in PEM electrolysis in the past century.[5] Currently, hydrogen is derived from natural resources such as gas, oil, and coal through processes like steam reforming or gasification, leading to substantial carbon dioxide (CO_2) emissions. Water electrolysis can be considered as a convenient and adaptable alternative for hydrogen production to meet energy requirements. The PEM electrolysis technology is generally viable at lower temperatures (100–120 °C). It is also essential for the lifespan of PEM water electrolysers to fall within the 10^4–10^5 hour timeframe.[6] One key benefit of PEM cells is their ability to operate at relatively lower temperatures, often facilitated by a non-corrosive electrolyte. Furthermore, these cells exhibit a commendable tolerance towards CO_2 when utilizing atmospheric air. Some noteworthy advantages include the elimination of constraints associated with handling liquid phases coupled with relatively high voltage, current, and power density. PEM cells also excel in functioning at lower pressures, typically ranging from 1 to 2 bars, while maintaining resilience to pressure variations of the reactant gases. Furthermore, the compact and robust build of these cells, along with a simple mechanical design, add to their appeal, often complemented by stable building materials.
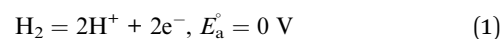
Despite the advantages mentioned above, the current state of cell development faces challenges in achieving a target operational lifetime exceeding 5000 hours (150 000 miles) for automotive and stationary applications for 10 000–40 000 hours.[7,8] The amount of energy needed to obtain hydrogen from water through electrolysis is four times greater than the energy required to produce it from methane. Hence, hydrogen energy production using water electrolysis is an expensive process.[3] Only deionized water can be used to feed the anode, as any impurities in the feed can affect the efficiency of the cell. Deionized water interacting with PEM is also corrosive to metals, which can cause corrosion of stainless-steel parts, releasing unwanted cations into the system and resulting in a loss of efficiency. Titanium metal parts can form an oxide layer on their surface, protecting them from harsh cell operating conditions. Therefore, self-protecting metals like titanium are generally used for cell parts of electrolysers, as they need to operate under harsh conditions. However, this oxide layer's growth on titanium can lead to increased cell resistance, which accelerates at higher temperatures. This, combined with membrane thinning, contributes to aging problems.[9,10] All of these factors can contribute to an increase in the production and operation costs of electrolysers (>2000€ per kW) and lower durability.[9,11]
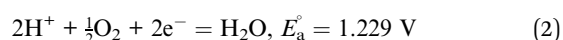
PEM fuel cells (PEMFCs), a crucial technology in this field, are still in their early stages, warranting further development for the successful commercialization of this promising technology.[12] A significant obstacle to implementing fuel cells in automobiles is their cost. Currently, the production cost is around USD 45 per kilowatt for volumes under 500 thousand units per year. A crucial prerequisite for commercialising PEMFCs is a reliable hydrogen fuelling station network. Furthermore, for these fuel cells to operate within a temperature range of −40 to 120 °C, they need expensive electro-catalysts, primarily platinum-based. Despite the cost, their desirable characteristics include low operating temperatures, high power density, and ease of scalability, positioning PEMFCs as a viable alternative for internal combustion engine (ICE) automobiles in moving forward.[13] Typically, a PEMFC (Fig. 1A) comprises several components: the membrane electrode assembly (MEA) (see Fig. 2 (ref. 14)), which includes a polymer electrolyte membrane (PEM), catalyst layers (CL), gas diffusion layers (GDL) with micro-porous layers (MPL), gas flow channels (GFC), and bipolar plates (BPP). Similarly, an electrolyser (Fig. 1B) also consists of a PEM, catalytic layers, porous transport layer (PTL), gaskets, titanium-based bipolar, and current collector plates.[15] The catalyst layers within these cells typically consist of a specialised catalyst layer comprising a catalyst (i.e., Pt/C, Ir/Ru) and an ionomer. The ionomer plays a critical role in binding the catalyst particles together and providing triple-phase boundaries as active sites for the reaction. Consequently, the composition of the ionomer ink is pivotal for managing ohmic losses and optimizing gas transport within the cell.[13]

Reactions taking place in fuel cells and electrolysers are shown in eqn (1)–(4) below, where the half-cell potentials (E_a°) for each reaction are shown below:

PEMFC anode:

$$H_2 = 2H^+ + 2e^-, \; E_a^{\circ} = 0 \text{ V} \tag{1}$$

PEMFC cathode:[16]

$$2H^+ + \tfrac{1}{2}O_2 + 2e^- = H_2O, \; E_a^{\circ} = 1.229 \text{ V} \tag{2}$$

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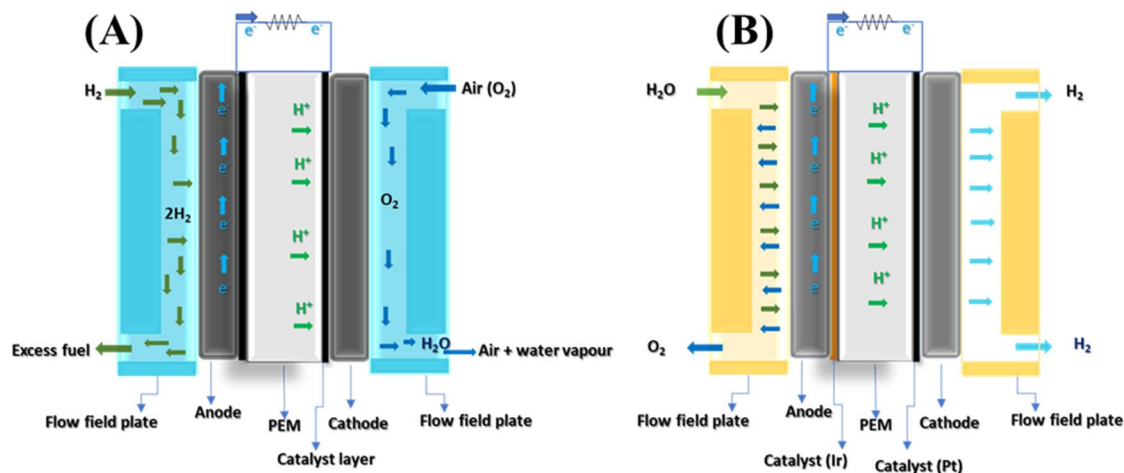
Fig. 1 Schematic representations of a PEM fuel cell (A) and PEM water electrolyser cell (B).
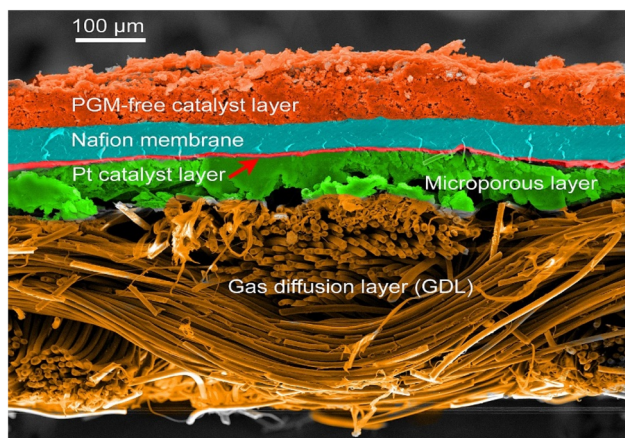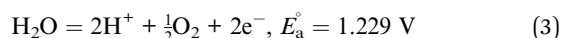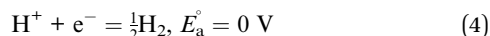

Fig. 2 SEM image of a typical membrane electrode assembly. Reproduced with permission from ref. 14, copyright© 2017 ECS – The Electrochemical Society.

PEMWE anode:[17]

$$H_2O = 2H^+ + \tfrac{1}{2}O_2 + 2e^-, \ E_a^\circ = 1.229 \text{ V} \qquad (3)$$

PEMWE cathode:

$$H^+ + e^- = \tfrac{1}{2}H_2, \ E_a^\circ = 0 \text{ V} \qquad (4)$$

Obviously, the electrolyte membrane, usually crafted from a highly specialised polymer, plays a crucial role in fuel cells. Generally, it acts as a barrier, preventing gas exchange between the hydrogen feed at the anode and air at the cathode while also serving as an insulator between the electrodes. Some of the important requirements of membranes used in PEM cells are: (i) they should have good mechanical strength and chemical resistance; (ii) they should have high proton conductivity (around 90–100 mS cm^{-1}) at low to medium temperature ranges (less than 100 °C). Generally, electrolytes used in electrolytic cells are typically classified into two types: liquid electrolytes

and solid electrolytes. Common liquid electrolytes include alkali-based solutions, molten carbonates, and phosphoric acid. Solid electrolytes, on the other hand, consist of polymer electrolyte membranes (PEMs), anion exchange membranes (AEMs) and solid oxide electrolytes (SOEs). Generally, PEMs offer several advantages over AEMs, SOEs, and liquid electrolytes. They often enable cells to operate efficiently at low temperatures (60–80 °C) while providing durability and requiring less maintenance. Additionally, PEM-based cells can achieve higher current and power densities by offering lower internal resistances compared to liquid and other solid electrolyte systems.[18,19] The PEMFC, originating from General Electric's use of an ion-exchange resin as an electrolyte in 1959 for space applications, subsequently has become a key player in portable power sources. The recent adoption of polymer membrane-based electrolytes has bolstered fuel cells and electrolysers, positioning them as leading candidates for automobiles and buildings and replacements for rechargeable batteries.

Generally, the performance factors for PEMFCs include operating temperature, pressure, and gas stream humidity. Wang et al.[13] found that increasing operating temperature and pressure, along with proper gas stream humidification, enhances the general performance of PEMFCs. However, inefficiencies, especially at low current densities, were observed at operating temperatures that are higher than the gas flow humidification temperatures. As mentioned earlier, the development of membranes for electrolytic cell applications traces back to 1959 when General Electric (GE) initiated tests with phenolic membranes. These early membranes generally had a short lifespan of up to 1000 hours, had low mechanical strength, and a power density of 0.05–0.1 kW m^{-2}. In the early 1960s, they enhanced the power density of their cells by introducing partially sulfonated polystyrene sulfonic acid membranes, reaching 0.4–0.6 kW m^{-2}, and these were employed application in NASA's Gemini flights. In the late 1960s, GE improved mechanical strengths and membrane lifespans with cross-linked polystyrene-divinylbenzene sulfonic

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acid membranes, achieving a power density of 0.75–0.8 kW m^{-2}. However, proton conductivities remained sub-optimum. In the 1970s, Du Pont's development of the perfluorosulfonic acid "Nafion®" marked a breakthrough in membrane technology, doubling specific conductivity and extending the membrane's durability by four orders of magnitude (10^4–10^5 hours).[20] Nafion® became a standard for PEMFC and continues to hold that status. Dow and Asahi Chemical Companies later developed advanced perfluorosulfonic acid (PFSA) membranes having shorter side chains with a higher ratio of ion conducting groups to the main chain.[21,22] Details regarding different types of membranes, their relevant properties, and their manufacturers are collated in the table (Table 1) below.

Generally, PFSA membranes can be modified through the incorporation of organic or inorganic additives such as $Zr(HPO_4)_2$ nH_2O, ZrO_2, SiO_2, SiO_2/TiO_2 blends, SiO_2/Al_2O_3 particles, resulting in the formation of composite membranes for better properties.[23] Alternative materials are also employed for membranes in addition to fluorinated ones, thus aiming to lower costs compared to exclusive use of fluorinated materials. Some of these partially fluorinated membranes have been found to yield enhanced stability and superior mechanical properties. While partially fluorinated membranes exhibit proton conductivity, their cost is at par with that of fully fluorinated counterparts. One of the widely accepted partially fluorinated membranes is a sulfonated copolymer that incorporates the α,β,β-trifluorostyrene monomer.[18] One of the initial commercial membranes, made of polystyrene-sulfonic acid (PSSA), can undergo modification by replacing tertiary hydrogens with methyl groups and fluorine. This modification can result in the creation of sulfonated poly(α,β,β-trifluorostyrene), thus enhancing stability and prolonging the membrane's lifespan. Polymer materials known for good chemical resistance, mechanical properties and thermal stability, such as tetrafluoroethylene, poly(tetrafluoroethylene-co-perfluoropropylvinylether), poly(tetrafluoroethylene-co-hexafluoropropylene), polyethylene, polyvinyl fluoride, polyvinylidene fluoride, poly(vinylidene fluoride-co-hexafluoropropylene), poly(ethylene-alt-tetrafluoroethylene), etc. possess promising structures for grafting sulfonic acid (SA) onto these membranes. Radiation-induced grafting stands out as a prevalent method for preparing such polymers, incorporating SA and modified SA segments. While PSSA-based membrane materials exhibit a significantly higher degree of hydration compared to Nafion membranes, their conductivity is relatively lower.[24]

Polybenzimidazole, an aromatic polymer, has been studied recently for membrane applications because of its favorable chemical resistance and thermal stability, along with cost-effectiveness compared to fluorinated counterparts. Nevertheless, its proton conductivity remains relatively modest (around 0.02 to 0.06 S cm^{-1}).[25,26] Another prospective class of membrane materials, polyarylene ethers, has demonstrated commendable mechanical and chemical attributes. To make these polymers more desirable for membrane applications, they can be functionalized using sulfonic acid groups.[27] Polyimide-based

Table 1 The PEMs can be categorized as fluorinated, partially fluorinated, non-fluorinated, acid-based, and other types.[18]

Sr. No.	Types	Examples	Manufacturers	Advantages
1	Fluorinated	• Perfluorosulfonic acid-based membranes	• DuPont • Asahi glass • Solvay • Solvay-Solexis • Aciplex-S	High stability, excellent chemical and mechanical strength, and high proton conductivity
2	Partially fluorinated	• Sulphonated polymer of α,β,β-trifluorostyrene • Crosslinked sulfonated polystyrene membranes	• Ballard advanced materials • Ionics Inc., Watertown	Better mechanical properties, higher stability and relatively high proton conductivity, and cheaper than perfluorinated counterparts
3	Non-fluorinated	• Sulfonated polyether ether ketone-(SPEEK) • Sulfonated polyarylene	• Produced by sulfonating commercial Victrex® and Gatone® PEEK • Asahi glass Company Ltd. and Asahi chemical industry Company Ltd., Japan	Cheaper, suitable for water purification and treatment, and more environmentally friendly
4	Acid-based blends	Polyethyleneimine (PEI)/SPEEK blend	—[a]	Lower vanadium ion permeability, higher water uptake, higher energy efficiency, lower charge capacity loss, good stability, and suitable for vanadium redox flow battery applications
5	Other types	Composite membranes	Can be prepared by combining Nafion (produced by DuPont, Aciplex, Asahi) with reinforcing fillers	Comparatively cheaper, high operating temperature range, can be recycled, higher water uptake, and more environmentally friendly

[a] Information not available.

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membranes possess noticeable thermal stability, albeit their manufacturing process has faced significant challenges associated with casting films from solutions.[28] Polyphosphazene-based membranes represent yet another category, displaying reasonable proton conductivity, complemented by favourable thermal and chemical properties. Moreover, these polymers offer facile functionalization capabilities, though they often exhibit poorer mechanical strengths.[29] Out of all the types of polymer electrolyte membranes available to date, perfluorinated membranes still stand out as the superior choice. This is primarily due to their exceptional physio-chemical properties, performance, and durability. However, they do have significant drawbacks, which will be discussed in the following section.

2. Fluorinated membranes

As already stated, perfluorinated membranes exhibit excellent chemical and mechanical properties, high ion conductivity, and prolonged performance. Therefore, they are the most commonly used solid electrolyte in both fuel cells and electrolysers. The carbon–fluorine bond situated in the main-chain backbone not only endows the material with inherent stability but also contributes to hydrophobicity and good thermal and electrical insulation properties.[30] However, a notable limitation is their operating temperature, as temperatures exceeding 90 °C can lead to membrane dehydration, resulting in diminished performance. The rather high hydrophobic nature of the fluorinated polytetrafluoroethylene backbone contrasts with the hydrophilic nature of sulfonic acid groups in the sidechains (see Fig. 3).

The frequently employed perfluoropolymer membranes include Nafion, produced by DuPont, Aciplex from Asahi Chemical, Flemion by Asahi Glass, and Gore-Select by Gore and Associates. These polymers are characterized by a long-side-chain structure that bears the ionic groups crucial for facilitating proton conductivity.[30] This unique structural feature of these polymers allows for efficient proton transport, making them integral components in various applications, particularly in electrolysers, fuel cells, and other such devices where proton exchange is a key factor. In 1964, DuPont identified the desirable properties of Nafion for use as a membrane separator in chlor-alkali cells, producing chlorine and caustic soda. Simultaneously, in the USA, GE was working on PEM fuel cells for the space program, initially using unstable polystyrene-based sulfonic acid polymer membranes. While DuPont was exploring potential applications, GE discovered that Nafion membranes had the necessary oxidative stability

and proton conductivity, marking the introduction of Nafion membranes in PEMFCs for space and military applications. Polymeric membranes gained acceptance in chlor-alkali cells due to environmental concerns and energy cost issues. In the 1990s, interest in PEMFC technology grew substantially, prompting DuPont to redirect efforts toward developing Nafion technology for the industry. The initially developed membranes, N-115 and N-117, were suitable for high catalyst loadings and low current density.[20,31] Nafion membranes also offer an extended operational lifespan lasting up to 57 000 hours.[30] The figure (Fig. 3) below shows the general structure of Nafion membranes.

Preparing perfluorinated membranes involves a complicated process of copolymerizing tetrafluoroethylene with fluorocarbon vinyl ether containing a sulfonic acid group. This process, while effective, presents challenges in terms of higher costs, thus limiting its commercialisation. To address these issues, Solvay Solexis developed a new membrane prepared by copolymerising perfluoro-3-oxa-pentene sulfonyl fluoride with tetrafluoroethylene, resulting in the product now known as Aquivion. Generally, Aquivion stands out for its short sulfonic acid sidechains, providing high crystallinity, a greater ion exchange capacity, and enhanced hydration. However, the relatively shorter side-chains lack the ability to form interconnected hydrophilic networks, as seen in long side-chain membranes like Nafion, thus compromising some of the other desirable properties.[32]

3. Limitations of fluorinated membranes

Nafion, while the dominant product in the market for PEM electrolyser and fuel cell (FC) design and production, has many drawbacks. Firstly, its proton conductivity relies on water-filled channels, making it unsuitable for temperatures significantly below 0 °C or above 100 °C. While stable against peroxide-type degradation, it can decompose when exposed to adventitious cations or peroxide radicals. In addition, contaminating ions can reduce the conductivity of the membrane. The degradation of PEM affects the performance of electrolytic cells. The main reasons are mechanical, chemical, and thermal stresses due to the operating conditions. These stresses can cause cracks, tears, punctures, and pinholes.[2,33] Such deformations can lead to short circuits, combustion from gas mixing (H_2/O_2), and other safety hazards. The release of hydrogen fluoride indicates membrane thinning and pinhole formation, as it is produced during the attack of $\cdot$OH radicals on the main chain. Moreover, Nafion generally exhibits poor mechanical properties and chemical stability, especially at elevated temperatures, often leading to noticeable degradation over multiple working cycles. According to a study conducted by Xiao-Zi Yuan et al.[34] on Nafion membranes, it was found that after 800 hours of operation, membranes underwent thinning and formed pinholes. In electrolysers, typically, a thicker PEM is used in comparison to a fuel cell. However, the degradation effect on the membranes is very similar to that found in fuel cells. This leads to higher
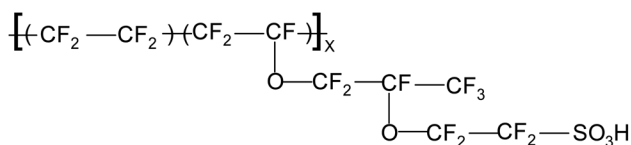

Fig. 3 A generic structure of Nafion.

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hydrogen crossover and, eventually, a decline in cell performance.[3]

As we know, the glass transition temperature (T_g) of a PEM is a significant aspect to consider for its end-use applications. Differential Scanning Calorimetric (DSC) studies have shown that Nafion exhibits two glass transition temperatures: a lower glass transition temperature (T_g) around 125 to 130 °C, attributed to the main chain mobility, and a higher T_g ranging from 195 to 240 °C, owing to the motion of the sidechains.[35] Water has a plasticizing effect on Nafion, which can slightly reduce the membrane's crystallinity. In a study by Huang et al.,[36] it was observed that relative humidity (RH) significantly affected the stress–strain behaviour of MEAs based on Nafion membranes at room temperature (see Fig. 4). As humidity increased, the elastic modulus decreased, indicating an inverse correlation between the two.

This observation confirmed that an increase in relative humidity has a plasticizing effect on the PEM, reducing its stiffness.

Additionally, operation at temperatures above the T_g of PEMs can cause a reduction in their dimensional stability. Therefore, generally, temperature and humidity are crucial factors for optimizing the performance of polymeric membranes in electrolyser and fuel cell applications.[37,38] Some of the major drawbacks of Nafion membranes are discussed in detail below:

(1) As discussed earlier, one of the most extensively researched materials for PEM electrolyser and fuel-cell membranes is Nafion® which is a copolymer composed of tetrafluoroethylene and perfluoro(4-methyl-3,6-dioxa-7-octene-1-sulfonic acid), developed by DuPont. According to the sources, the cost of Nafion 117 is approximately USD 711 for a piece of ca. 61×50 cm^2.[39,40] This high cost is attributed to the intensive process of fluorinating polyethylene into polytetrafluoroethylene and then functionalizing the resultant product with sulfonic acid groups. Therefore, producing this type of membrane on a commercial scale is quite challenging.
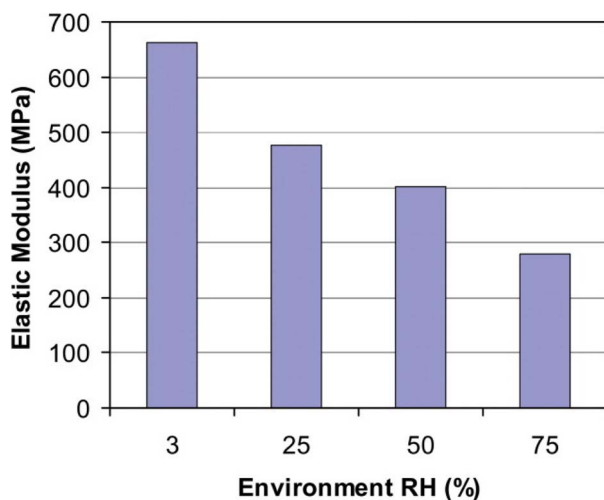

Fig. 4 A plot showing the relationship between elastic modulus and relative humidity. Reproduced with permission from ref. 36, copyright© 2006 Wiley Periodicals, Inc.

(2) Ionic clusters within fluorinated membranes, composed of sulfonic acid species, undergo dehydration when exposed to temperatures exceeding 80 °C. This characteristic often renders them unsuitable for high-temperature applications. Dehydrating these ionic clusters within the membrane can result in a significant loss of proton conductivity. The sulfonic acid species, crucial components forming the ion clusters in the fluorinated membranes, play a pivotal role in facilitating proton transport. However, as the temperature rises beyond the 80 °C threshold, the loss of water from the clusters disrupts the proton conduction pathways, thus adversely affecting the general performance of the membrane in high-temperature conditions.[41] This limitation underscores the necessity for alternative materials or design strategies to address the challenges posed by dehydration in order to enhance the suitability of fluorinated membranes for applications requiring elevated temperatures. The operational capabilities of Nafion membranes are also constrained by their limited performance under specific conditions, primarily stemming from poor ionic conductivities at low humidity and high temperatures. The challenge arises when the humidity levels decrease, impacting the ability of Nafion membranes to facilitate efficient ion transport. At lower humidities, the reduced water contents within the membrane hinder proton mobility, thereby diminishing overall ionic conductivity. Therefore, a combination of elevated temperatures and low humidity aggravates the challenges, as the membrane's ability to retain sufficient water content becomes severely compromised.[42]

(3) The high methanol permeability of Nafion membranes can pose a significant problem in the context of DMFCs. This trait of Nafion leads to a phenomenon known as 'methanol crossover'. This occurs when methanol molecules readily pass through the Nafion membrane, migrating from the anode to the cathode of the fuel cell. Methanol crossover is problematic because it results in the mixing of feed and oxidant flows, thus reducing the efficiency of the fuel cell. This effect, in turn, diminishes the overall performance of the fuel cell, leading to lower power output and decreased efficiency. Secondly, the methanol crossover contributes to mechanical instability. The interaction of methanol with the cathode can lead to detrimental side reactions, such as methanol oxidation, to form unwanted by-products. These by-products, in turn, can adversely affect the structural integrity of the fuel cell components, potentially causing material degradation, corrosion, or other related issues.

(4) The fluorine content in materials, particularly the stability of the carbon–fluorine (C–F) bond, contributes to environmental concerns. The C–F bond's remarkable stability makes it resistant to easy decomposition in the environment. Hence, this enhanced stability comes with a downside, as materials containing high levels of fluorine, such as per- and polyfluoroalkyl substances (PFASs), pose a risk of bioaccumulation.[43] PFAS is a general term used for materials containing fluorine atoms attached to an alkyl chain. Perfluorinated membrane materials and their degradation products fall under this category.
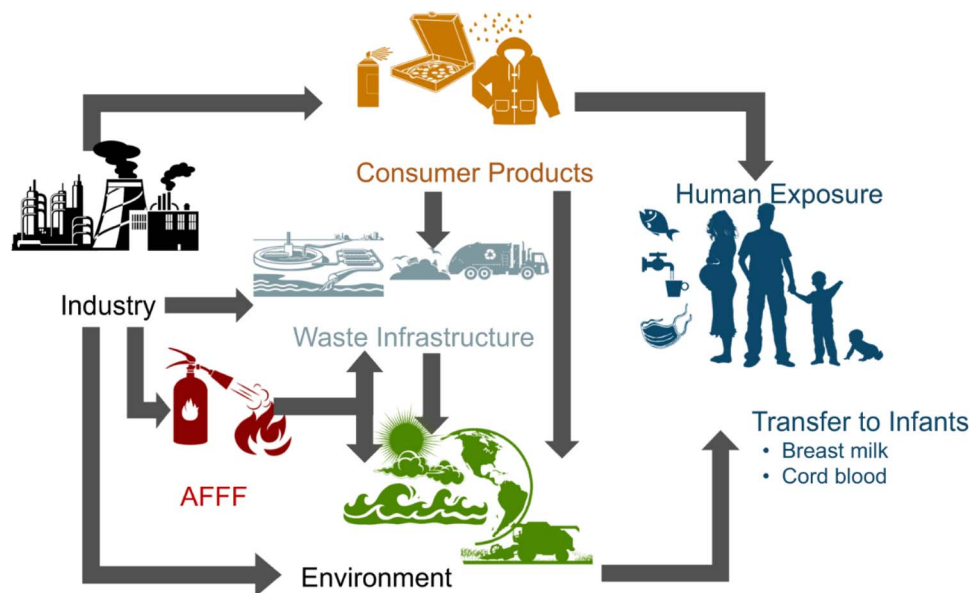
Fig. 5 Toxic effects of PFAS released into the environment from various sources, reproduced with permission from ref. 44, copyright 2019 Springer Nature LTD.

The high mobility of PFASs in water facilitates their leaching into water systems, causing environmental issues like soil and water contamination. This contamination, in turn, can potentially be a significant threat to both humans and ecosystems (Fig. 5 (ref. 44)). The persistence of PFASs in the environment allows them to accumulate over time, and their presence in water systems can lead to long-term exposure for aquatic organisms and, subsequently, organisms higher in the food chain, including humans.[38,45,46]

The bioaccumulation of PFASs raises concerns about their potential adverse effects on human health, as these substances can enter the food chain and accumulate in various tissues.[47,48] Additionally, the impact on ecosystems is substantial, as PFASs have been associated with various ecological disruptions, affecting aquatic life and potentially causing long-term harm to the balance of ecosystems. Addressing the environmental implications of fluorinated compounds, especially PFASs, necessitates a comprehensive approach, including developing environmentally friendly alternatives and stringent regulations to mitigate the release and accumulation of these substances in the environment. Perfluorocarboxylic acids (PFCAs), which are environmentally persistent and possibly toxic, have become widely distributed globally. Trace amounts, as low as nanograms per milliliter, have even been detected in human blood.[49] Toxicology research on laboratory animals has associated PFCAs with tumor formation, hormonal disruptions, developmental issues, and immune system toxicity. In humans, exposure to PFAS compounds has been linked to various health risks, such as thyroid disorders, ulcerative colitis, high cholesterol, liver disease, developmental delays, cancers, and immune suppression. Furthermore, Nafion by-product 2 has been identified in the blood of most study participants, including young children.

Moreover, the water-soluble nature of per- and polyfluoroalkyl substances (PFAS) allows them to traverse longer distances in both freshwater and marine environments and can lead to the contamination of drinking water. In addition, PFAS proves challenging to eliminate fully through water treatment processes, thus contributing to concerns about drinking water safety. Numerous studies have also documented elevated concentrations of PFAS in drinking water near industrial areas, emphasizing the wide-ranging consequences of these substances in water ecosystems.[49–52] Consequently, fluorine-containing membranes are being banned. This restricts the use of such membranes in PEM fuel cells and electrolysers, necessitating the development of non-fluorinated, non-toxic materials for membrane preparation.

When applied in marine settings, membrane-based fuel cells can encounter challenges due to harsh conditions impacting their performance. The presence of sodium, potassium, chromium, and magnesium ions in the marine environment poses a problem related to the efficient working of fluorinated membranes. These cations have the ability to replace protons in the sulfonic acid group sites, causing degradation of the electrolyte membrane and reducing its conductivity. In addition to the above-mentioned contaminants, NO_2 and SO_2 also emerge as major pollutants, with salt being a predominant factor in the marine environment. Mechanical degradation of PEMs plays a significant role in the loss of efficiency in electrolysers by reducing protonic conductivity and causing ohmic losses. The rate of degradation is strongly dependent on the cell's operating conditions.[33] Currently used Nafion membranes are believed to lose mechanical strength when they swell in water at higher temperatures. A study showed that polymer membranes cast at higher temperatures (200 °C) are more prone to degradation than those cast at lower temperatures.[53] High-

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temperature membrane casting could also lead to the chemical degradation of the ionomer. On the other hand, it has been observed that the membranes can shrink (thin) when subjected to compression and dehydration under a high vacuum inside the cell.

4. Failures of fuel cells and electrolysers

As mentioned in the previous sections, PEM fuel cells comprise multiple parts like catalysts, membranes, gas diffusion layers (GDLs), bipolar plates, and seals. During the service life, these parts can deteriorate or malfunction, resulting in the breakdown of the entire fuel cell system. Examples of degradation mechanisms encompass catalyst particle growth, preferential dissolution of alloys, corrosion of carbon supports, catalyst poisoning, membrane breakdown, loss of sulfonic acid groups, formation of surface films on bipolar plates, alterations in hydrophilicity, and decomposition of PTFE within catalyst layers and GDLs.[54,55] The following section overviews the prevalent and substantial failure issues linked to the currently used PEM-based electrolytic cells. Fig. 6 represents some of the general operating conditions that trigger cell failures.

4.1 General cell failures

In a typical PEMFC stack, the serial (bipolar) cell arrangement makes the stack susceptible to catastrophic failure if one cell fails. Meeting durability targets for automotive applications proves challenging due to changes in operating conditions, including cyclic power output and environmental factors such

as relative humidity and temperature. Current designs are not adequate to demonstrate stable performance beyond a few thousand hours under these demanding conditions. A significant risk for water electrolysers is the mixing of the resultant gases produced during the electrolysis process, namely H_2 and O_2, which are then stored inside the unit.[15] The gradual performance decline observed in electrolytic cells is often linked to electrode degradation, involving issues like platinum catalyst sintering, dissolution, ohmic losses, membrane degradation, and carbon corrosion. In this context, ensuring the durability of catalyst supports during start/stop cycling remains an ongoing challenge.[36]

4.1.1 Failure of the catalytic system. PEMFCs commonly rely on reformed fuel as the primary anode feed, containing CO, CO_2, NH_3, and traces of H_2S. However, poisoning platinum catalysts by pollutants in the feed, such as CO, poses a significant challenge to PEMFCs. In an electrolyser, platinum-based catalysts are mainly used at the cathode to facilitate the evolution of hydrogen gas. The high cost of platinum-based catalysts used in fuel cells and electrolysers add to the problem. Hydrogen impurities introduced during manufacturing can lead to performance degradation and even irreversible damage to MEAs in PEMFC stacks. Even a minor concentration of CO has the potential to cover the anode's platinum surface entirely. The extent of CO poisoning is closely linked to the CO concentration and the duration of exposure. Even trace amounts of sulfur-containing impurities can significantly poison the catalyst by allowing sulfur-based species to adsorb onto its active sites. To avoid anode catalyst poisoning, carbon monoxide-free hydrogen gas is typically required. Utilising ammonia as a hydrogen source in the feed can be considered a good choice to avoid catalyst poisoning from oxides of carbon-based impurities, but the concerns about trace ammonia's impact on the anode and membrane have limited its application.[56]

Electrochemical platinum dissolution is a concern in fuel cells and electrolysers, particularly under high cell voltage or open-circuit voltage (OCV) conditions.[57–59] Another important factor affecting the performance of both fuel cells and electrolysers is the migration of the platinum catalyst, as well as platinum agglomeration (see Fig. 7 (ref. 60)). Generally, platinum stability is compromised at potentials beyond 1.188 V in acidic environments. At potentials higher than 0.98 V, oxidation can occur, leading to platinum oxide formation. Anodic and cathodic dissolution is assumed to occur by forming metastable platinum complexes. The presence of chlorides accelerates catalyst dissolution by shifting the equilibrium between platinum oxidation and reduction. In fuel cell mode, higher potentials, typically ranging from 0.9 to 1 V, amplify this effect. Furthermore, platinum ions naturally tend to precipitate on larger particles, significantly reducing the catalyst's active surface area. This reduction in active area not only decreases catalytic efficiency but also increases the cathode potential, thereby leading to higher activation overpotential.

In water electrolysers, platinum catalysts are mainly used in the cathode, where the potential is usually lower. As a result, platinum catalysts are relatively stable at the cathode in
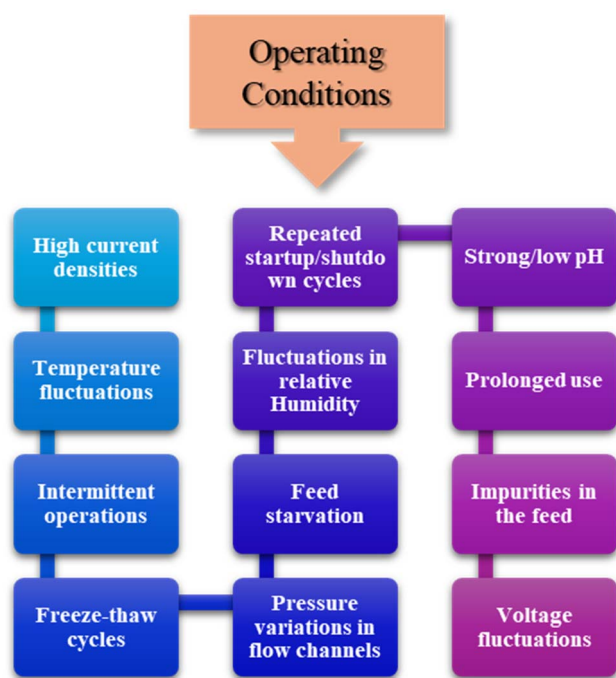

Fig. 6 A schematic representation of the general operating conditions that trigger cell failure in PEMFCs and PEMWEs.
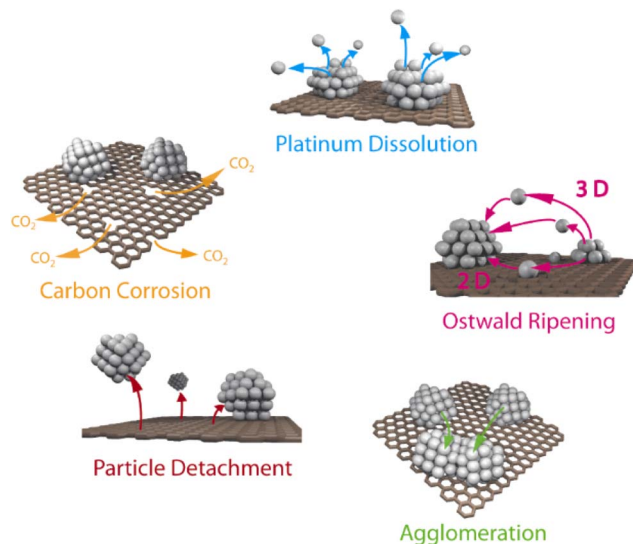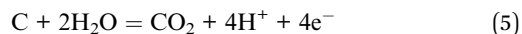
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Fig. 7 Platinum catalyst failure types.[60]

PEMWEs. However, they are prone to detachment from the carbon support due to the low potential. The carbon support can suffer from mechanical degradation after prolonged use, especially under harsh operating conditions in PEMWEs, such as high current density and temperature fluctuations. This degradation can manifest as cracks, structural weakening, or even material loss. Additionally, the presence of metallic impurities in the cathode can inhibit catalytic activity by forming agglomerates that block the active sites.[61] On the other hand, iridium is used as the catalyst in the anode. However, the potential at the anode is usually higher. Therefore, iridium can undergo electrochemical dissolution by forming soluble iridium oxides. In addition, higher chances of catalyst failures such as Ostwald ripening, migration, and agglomeration are in the anode. In addition, oxygen bubbles can nucleate in the anode where the oxygen evolution occurs. This can affect the stability of the catalytic layer. The formation and detachment of these bubbles can cause mechanical damage to the catalyst, potentially leading to catalyst detachment.[62] Corrosion due to carbon-mediated corrosion can be a significant issue in fuel cells, occurring during a fuel shortage or localised flooding. Generally, the catalytic layer in a fuel cell consists of a catalyst, catalyst support, and a polymer membrane. Carbon-supported platinum nanoparticles are used as catalysts due to their high surface area. However, carbon becomes unstable in PEMFC operating conditions, often leading to performance degradation as Pt detaches or agglomerates, owing to weakened Pt–carbon support interaction. The degradation is primarily caused by carbon's electrochemical oxidation,[57,63–65] which is generally accelerated at elevated temperatures and potentials. Although PEMFCs operate at low temperatures, certain conditions during transportation, such as high water content, high oxidation potential, low pH, *etc.*, can accelerate carbon-mediated corrosion. A typical electrochemical oxidation reaction of carbon is shown in eqn (5).

$$C + 2H_2O = CO_2 + 4H^+ + 4e^- \qquad (5)$$

The three main conditions causing carbon-mediated corrosion are high potential from feed starvation during flooding, repeated startup/shutdown cycles, and long-time effects during normal operation. During cyclic operations, it can be induced or exacerbated, leading to insufficient fuel and causing carbon-mediated corrosion to support the current flow. The standard carbon corrosion potential is 0.207 V. When the potential at the anode is below this value, the current is generated by utilising the fuel. When the value is beyond 0.027 V, corrosion of the anode catalyst layer occurs, which is driven by the degradation of the carbon support to facilitate the supply of protons. As a consequence, platinum agglomeration can also happen. A mitigation strategy to prevent this type of corrosion from occurring is operating the cells under fixed flowrates and current densities, which helps avoid these processes and reduces the extent of corrosion.[57,58] The platinum catalyst used in the cathode of electrolysers is also sensitive and requires careful handling.[33] Any impurities in the catalyst layer can alter its properties and lead to the formation of free radicals that degrade cell components. In a PEMWE, the porous transport layer (PTL) and the bipolar plates perform several critical functions.[63] They facilitate the distribution of electrical current and provide mechanical support for the MEA. Potential (voltage) and pH are two critical factors affecting the performance of the bipolar plates and PTL.

In PEMWEs and PEMFCs, a decoupling phenomenon has been observed between the applied current density and the potential at the PTL/CCM interface. The potentials at the anode and cathode remain constant, irrespective of the applied current density, and are independent of the PTL/CCM interface potential, which is considerably higher. Additionally, the pH at the interface is acidic, ranging from 0 to 3.[66] Due to the high electrochemical potential and the harsh environment at the anode, the catalyst layer is at high risk of corrosion during prolonged use. This is particularly critical since the anode is where water is oxidised to evolve oxygen. Therefore, titanium is preferred as the material for the PTL and BPP at the anode. However, the passive layer formed on titanium has a high interfacial contact resistance and is prone to corrosion after about 1000 hours of constant operation. To mitigate this, titanium is generally coated with noble metals such as gold, platinum, or iridium, resulting in 50 to 70% of the expense being devoted to making the bipolar plates and PTL robust for operation. Using other affordable metals like steel or copper to manufacture the bipolar plates and PTL parts would result in corrosion under the harsh conditions of the anode. This can lead to the release of unwanted metal ions, which can then attack the ionomers and PEMs, leading to their degradation, similar to the degradation observed in PEMFCs. Thinning of the polymer membrane through Fenton's reaction at the cathode, caused by ferric ion species, is also observed. The water used in a typical PEMWE is highly deionized, with a conductivity of approximately 1.0 μS cm^{-1}. This prevents membrane and ionomer poisoning due to foreign cations.[63,66]

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4.1.2 Water management issues. The main cause of limited oxygen transport in PEM fuel cells is the accumulation of liquid water generated at the cathode, primarily through the oxygen reduction reaction (ORR). If the water removal rate lags behind its generation rate, water flooding can occur, impeding oxygen transport and hindering cell performance. Moreover, in the context of channel flow, sufficient airflow rates can remove liquid water within the flow fields. However, pressure drops, especially along straight segments and around bends, may lead to water accumulation, potentially obstructing channels and causing cell shutdown. Critical physical attributes influencing these dynamics include the length and cross-sectional area of the pathway. Effective water management, which includes addressing flooding, membrane dehydration, and feed gas humidification, is crucial for optimising PEM fuel cell performance. Striking a balance between membrane hydration and preventing flooding is challenging, necessitating careful control of operating conditions. Failure to manage water effectively can thus result in unpredictable, unreliable, and inconsistent cell performance, impacting both transient and long-term functionality of the fuel cell.[57,58]

On the other hand, in a PEMWE, product gases can accumulate inside the cells, leading to bubbling and blockage of the flow channels. The problem occurs when gas generation at the cathode surpasses the flow channels' ability to remove it. Gas removal capacity depends on the water flow rate and the cross-sectional area of the channels. If bubbles persistently block the channels, localised water shortages can develop, leading to uneven current distribution and decreased efficiency. An increase in current densities has been observed to raise the risk of stagnant bubbles forming at the top end of the channels (as can be seen in Fig. 8).[62,67]

Water starvation is a significant problem in electrolysers. Overall starvation occurs when the total water supplied to the electrolyser is less than the water consumed, whereas local starvation results from a non-uniform supply of water, high current densities, and high pressure.[68] Water starvation leads to membrane drying, hotspots, and ohmic losses, as insufficient water flow causes uneven temperature distribution and impairs heat removal, leading to the degradation of the MEA. At high current densities (above 0.5 A cm^{-2}), membrane drying has been observed near the water outlet due to inadequate water flow, resulting in an increase in local ohmic resistance. Therefore, high current densities combined with insufficient water flow can significantly affect the voltage distribution.

4.1.3 Degradation of seals. Silicone rubber, a component used in fuel cells for sealing the fuel and air chambers, can decompose under operating conditions, with factors such as strong acidity, temperature, the presence of fluoride ions, and the degree of mechanical stress influencing the overall decomposition rate. The acidic environment and increased fluoride concentration can also accelerate silicone degradation. Severe degradation of gaskets, especially at the edges of the MEA, often poses risks of gas leakage, fuel or air shortage, and reduced fuel consumption and stack performance. Decomposition fragments can redeposit, particularly at the cathode side, increasing activation overvoltage and altering porosity and wetting behaviours. This may also lead to increased diffusion overvoltage, air or fuel starvation, and carbon corrosion. The acidic nature of the PEM, combined with thermal stress on the silicone seal, can lead to chemical degradation, often resulting in yellow discoloration. Although the silicone's mechanical functionality remains intact, its chemical breakdown can release products that move toward the cathode under the influence of the electric field. Perfluorosulfonic acid (PFSA) membranes generally prevent these products from permeating but can cause their accumulation in the catalyst layers.[57,58] In PEMWEs, degradation of the sealing gaskets can occur due to radicals, such as peroxides, attacking the components. This degradation can result in gas leaks, which not only reduce the overall efficiency of the system but also pose a safety risk. As the sealing gaskets degrade, fragments of the sealant can redeposit in the pores of the active layer or porous transport layer (PTL). This deposition can alter the porosity and wetting behavior of these layers, impacting the general performance and durability of the electrolyser. Additionally, metallic impurities or other fillers released from the degraded sealants can lead to inhibition of the ionomer and catalyst, which are critical for the electrolyser's operation. These impurities can cause a decline in the membrane's ionic conductivity and
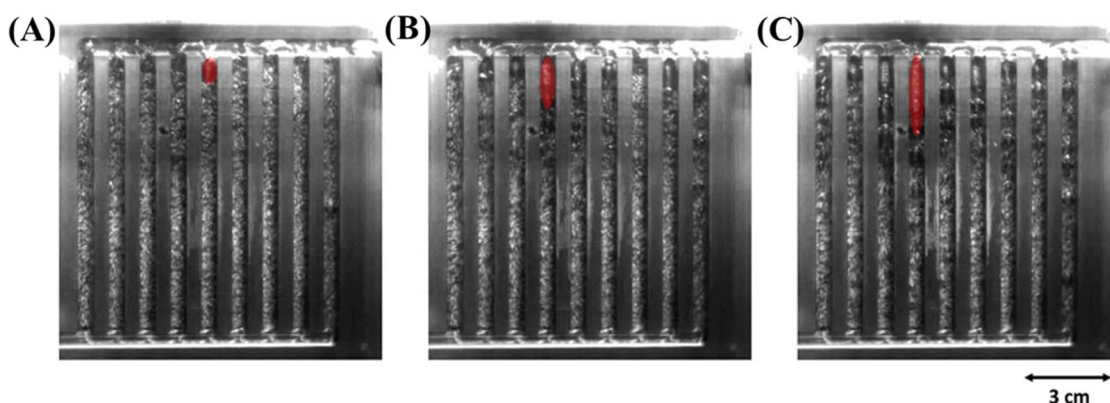

Fig. 8 Flow field of a PEMWE at different current densities (A) 0.3 A cm^{-2}, (B) 0.6 A cm^{-2}, and (C) 1.0 A cm^{-2}, showing stagnant bubbles at the top end of the channel.[67]

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interfere with the catalyst's activity, thereby reducing the efficiency of the PEMWE. To mitigate these issues, it is essential to use materials resistant to radical attacks and to implement regular maintenance and monitoring strategies to ensure the integrity of the sealing gaskets.[61]

4.1.4 Degradation of gas diffusion layers. The gas diffusion layer (GDL) is one of the critical components of a fuel cell. As the name suggests, the primary function of a GDL is to transport feed gases from the inlets to the catalyst layers. Additionally, it facilitates the flow of water from the catalyst layer to the flow channels. GDLs also conduct electrons and are typically made of carbon fibers. Therefore, the durability of GDLs is crucial.[69,70] Two factors affecting the durability of GDLs are mechanical degradation and chemical degradation. Mechanical degradation is primarily caused by compression during cell assembly, freeze–thaw cycles, and damage from water and gas flows. Furthermore, the carbon fibers in the GDL are susceptible to corrosion. Changes in the GDL can significantly impact the gas permeability and contact resistance of the cell, with compression pressure playing a crucial role. Open-circuit voltages (OCVs) are also greatly influenced by these changes. Additionally, low operating temperatures below zero can cause the GDL to stiffen, leading to mass transport losses.

4.1.5 Ohmic losses. In PEMFCs, overall Ohmic losses result from resistance to the transport of both electrons and protons. Electron transfer occurs in various elements, such as the carbon support in CL, the GDL, and the BPP. The transport of protons occurs in the PEM membrane and within the anode and cathode catalyst layers containing the membrane ionomer. The BPP, crucial for stack mechanical strength, also serves as a current collector, facilitates heat removal, and manages reactant supply and water removal through flow fields. However, BPPs must endure corrosive and humid conditions, posing challenges like corrosion-related electrical conductivity loss. Balancing high mechanical strength, corrosion resistance, and electrical conductivity is essential for BPPs.[57,58] It is to be noted here that the highest source of ohmic losses in a fuel cell

is the polymer membrane. Since the membrane conductivity depends on its hydration level, water management is crucial for the ideal operation of fuel cells. Higher current densities can sometimes lead to overheating of the electrodes, thereby causing the polymer electrolyte membrane to dry out quickly, leading to ohmic losses. Furthermore, poisoning of the membrane caused by contamination due to impurities contributes to an increase in ohmic losses. Another predominant cause is the resistance of the electrical elements of the fuel cell. Improper assembling of the cell stack and degradation of the components, in general, can also contribute to ohmic losses.[71]

Ohmic losses in a PEMWE can result from the ohmic resistances caused by the internal components. However, most losses are attributed to the resistance to proton transport within the membrane. Like a fuel cell, membrane thickness is an important aspect contributing to ohmic losses. Activation losses in an electrolyser depend on the rates of the OER and HER at the electrodes.[72] Insufficient feed at the anode can result in mass transport losses (see Fig. 9). This occurs when the reactant water at the anode is consumed quickly, leading to a shortage of reactant at the anode surface. Slow diffusion of product gases can also result in mass transport losses. Electrochemical impedance spectroscopy (EIS) can be employed to detect ohmic losses and resistance in fuel cells and electrolysers.

A durability study of MEA[73] was performed at 90 °C, ambient pressure, and an RH of 10% at the anode and 30% at the cathode on a PEMFC stack under OCV holding mode, with intervals of 24, 48, and 144 hours, for a total duration of about 250 hours. The cell voltage decreased gradually, first exponentially and then linearly, during OCV holding. After reactivation, the voltage drop was almost fully recovered. It was also found that hydrogen crossover current densities (HCCD) were increased owing to the irreversible membrane degradation, leading to a reduction in OCV. From Fig. 10A below, it can be seen that the HCCD rate steeply increased after the 144 hour interval, continuing until the test was finished. The membrane
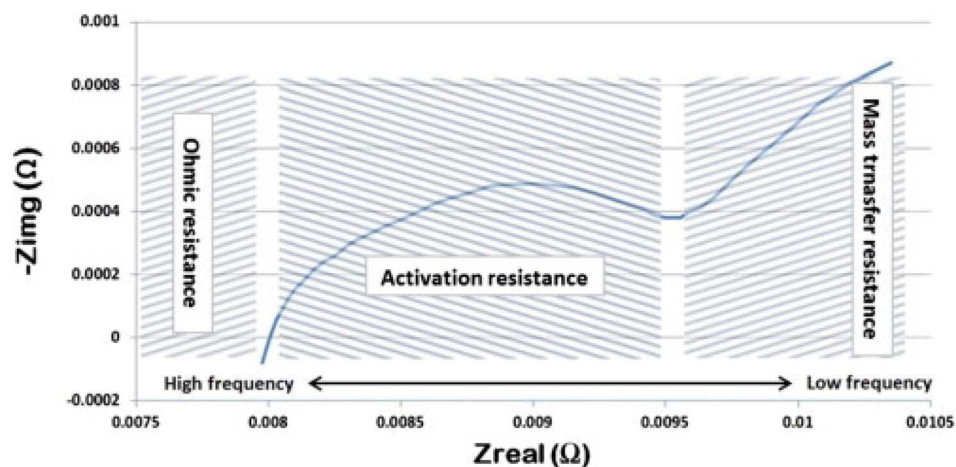

Fig. 9 An EIS plot showing the types of internal resistances causing losses in a PEMWE, reproduced with permission from ref. 72, copyright© 2014 Hydrogen Energy Publications, LLC. Published by Elsevier Ltd.

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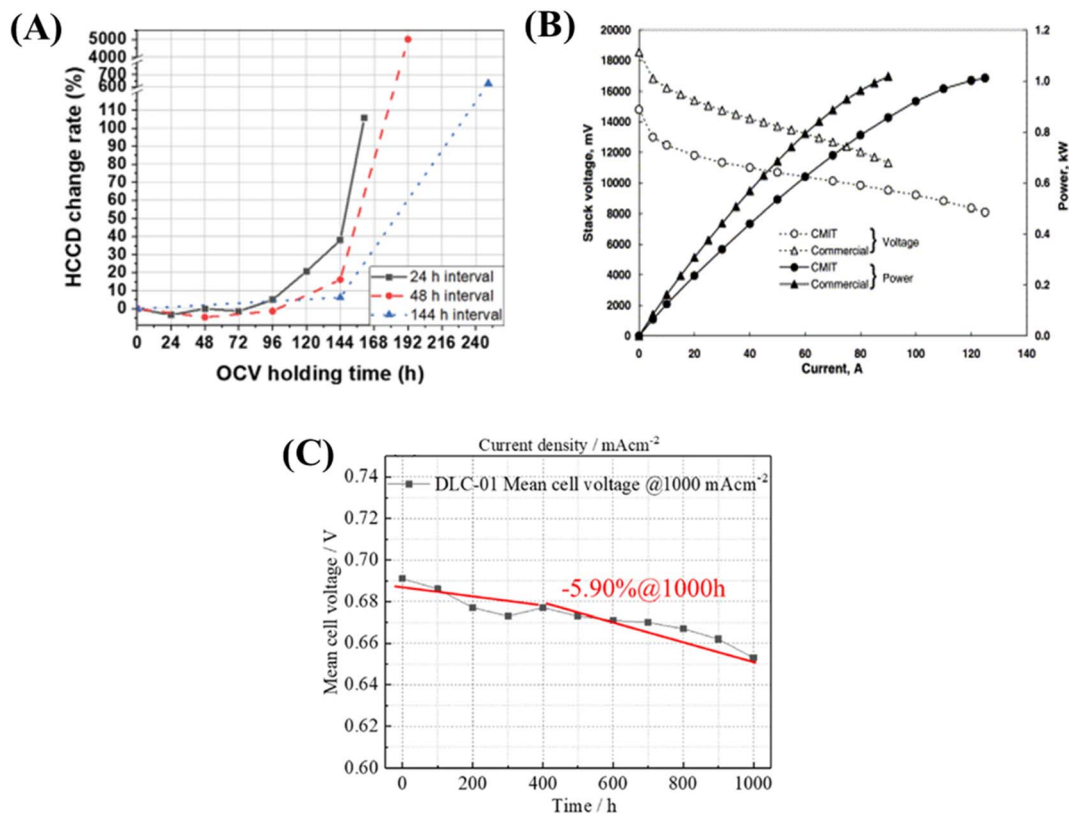J. Mater. Chem. A, 2025, **13**, 11236–11263 | **11247**

Fig. 10 (A) Plot showing an increase in hydrogen crossover overtime during the OCV holding tests on PEMFCs, reproduced with permission from ref. 73, copyright 2023 Springer Springer Nature SharedIt content; (B) graph comparing the IVP characteristics of two fuel cell stacks, reproduced with permission from ref. 74, copyright© 2003 Published by Elsevier B.V.; (C) graphs showing a drop in mean cell voltage overtime during the durability tests conducted on PEMFC stacks, reproduced with permission from ref. 75, copyright© 2021 Elsevier Ltd.

thickness also decreased from 25.6 microns to a minimum of 13.9 microns.

Fig. 10B shows the $I–V$ characteristics of a commercial fuel cell stack and a constructed 19-cell 'CMIT' stack discussed in a study by Giddey $et\ al.$[74] As we can see, with an increase in current, the voltage slowly drops for both stacks, possibly because of losses such as ohmic, activation, and mass transport losses, while the power increases until it reaches a peak before eventually decreasing. The commercial stack produced the highest output power of 1.02 kW when running at 90 A and 11.3 V at a temperature of 64 °C. In comparison, the CMIT stack produced 1.01 kW at a lower voltage of 8.1 V at a higher current of 125 A at 60 °C. This suggests that the commercial stack was relatively more efficient than the CMIT stack, and internal resistances within the cells and other components could influence cell performance.

In a 1000 hour durability study of 3-cell PEMFC stacks conducted by Chu $et\ al.$,[75] it was found that the cell voltage eventually dropped, indicating performance degradation, during two dynamic load cycles with varying operating conditions and load ranges. In the first 400 hours, the performance loss was slower but accelerated after 400 hours. By 1000 hours, the mean cell voltages of the stacks had dropped by 6% (Fig. 10C). After 500 hours, the hydrogen crossover rate was also rapidly increased in the cell stacks due to the failure of the MEA

components, which could primarily be attributed to pinholes or cracks formed in the membrane due to overheating. Higher operating temperatures have been found to improve the efficiency of the electrolysis process; however, higher temperatures may reduce the durability of electrolysers.[76] Fig. 11A shows the effect of operating temperatures on cell voltage and the resulting improvement in efficiency with an increase in temperature.

In the above study (see Fig. 11B), the performance of the MEA was assessed over 4000 hours. The changes in cell voltage were measured at different current densities (0.2 A cm^{-2}, 1.0 A cm^{-2}, and 2.0 A cm^{-2}). Initially, the cell voltage increased over time for all current densities until 1000 hours, after which it slightly decreased and then remained stable until 4000 hours of operation. The authors attributed this initial increase to the degradation of the MEA. However, the results varied depending on the types of catalysts and membranes used. This degradation can be mitigated to some extent by improving contact between the MEA components, which reduces the ohmic resistance within the cell. The choice of materials and robust design of the cell components play a crucial role in minimizing degradation, ensuring long-term efficiency and durability.[77]

Generally, nitrogen (N$_2$), nitrogen oxides (NO$_x$), and sulfur oxides (SO$_x$) can poison the catalyst and reduce reaction rates in the hydrogen cell. Carbon monoxide (CO) can also strongly bind to the catalyst, inhibiting the hydrogen oxidation reaction and

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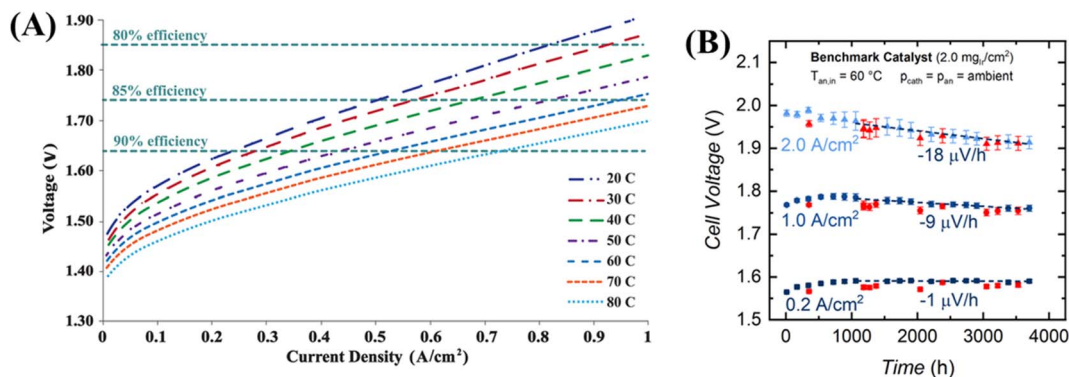This journal is © The Royal Society of Chemistry 2025

Fig. 11 (A) Graph showing the effect of temperature on voltage and current densities in a PEMWE, reproduced with permission from ref. 76, copyright© 2022 Hydrogen Energy Publications LLC. Published by Elsevier Ltd.; (B) graph showing the changes in voltage in 4000 hour durability test of a PEM electrolyser.[77]

impacting the cell's efficiency. Furthermore, hydrogen sulfide (H_2S), sulfur (S), metals, and ammonia (NH_3) can deposit and foul the membrane of the hydrogen cell, leading to decreased performance. Additionally, NH_3 can etch the membrane, potentially causing damage and reducing the cell's effectiveness. Oils, if present as contaminants, can clog flow channels within the cell, reducing gas transport and hindering overall performance. If these contaminants are not properly managed, they can also cause the degradation of the components of the cell over time, thus leading to decreased efficiency and potentially a shorter lifespan. Maintaining a high-purity hydrogen feed and preventing membrane fouling are crucial steps to optimize the performance and longevity of a hydrogen cell. Hydrogen crossover within PEM cells is another important problem that often accelerates the degradation of both the membrane and catalyst layer, thereby causing the development of flaws such as pinholes and cracks.[78] This, in turn, leads to the escalation of local hot spots and thermal degradation. Consequently, there is a loss of mass in both the membrane and catalyst layers, a reduction in activation surface area, catalyst sintering, and the generation of free radicals. These factors collectively diminish performance by lowering the open circuit voltage, voltage efficiency, fuel utilization efficiency, and overall efficiency of the PEMFC. Research indicates that fluctuations in relative humidity percentages can impact hydrogen crossover rates.[79] This phenomenon may arise from the maintenance of specific humidity levels within the membrane, which reduces its porosity and consequently lowers the rate of hydrogen crossover.[79]

4.2 Polymer electrolyte membrane failures

The failure of the membrane is thought to stem from a combination of thermal, chemical, and mechanical factors, often working in tandem, especially under harsh operating conditions.[80] Operational parameters, including humidity, temperature, freeze–thaw cycles, intermittent operation, and start-up/shut-down processes, significantly influence the deterioration of the membrane. The abrupt failure of a PEM cell or stack results from significant gas crossover occurring through cracks, pinholes, punctures, and damages in the membrane. Swelling and contracting caused by changes in humidity and temperature also influence the membrane's fatigue lifespan, with greater strain oscillations ultimately resulting in shorter fatigue life. Generally, the membrane's water content decreases as temperature rises, making it more susceptible to cracking. Basically, dry membranes are less mechanically stable, so elevated temperatures can compromise their integrity. Studies also indicate that the chemical structure of PFSA membranes starts to alter at temperatures exceeding 150 °C. However, this is not significant in the case of current fuel cells as they operate at a temperature below 100 °C. In addition, insufficient water feed may lead to the local drying of the membrane at the anode in electrolysers.[9,15] Metallic impurities in the feed water also hinder the smooth conductivity of protons, causing higher resistance. Some metal ion impurities can deposit on the catalyst layer, obstructing hydrogen evolution at the cathode. For instance, impurities like Ca^{2+} can form metallic hydroxides at the cathode site, resulting in additional ohmic losses.

Cyclic stresses, in addition, typically result in fatigue-related issues like pinholes and cracks, while continuous stress can induce membrane creep.[79] Although membranes can tolerate a certain degree of mechanical stress, such as pinholes and tears, in the long run, this can reduce the lifetime of PEM cells. Fluctuations in temperature and humidity cause stress in the MEA during cell operation, subjecting it to mechanical loading. Fig. 12A and B (ref. 55 and 81) shows the typical factors contributing to the degradation and failure of PEMs under the above-mentioned conditions.

Thinning of the membrane is an additional factor contributing to elevated hydrogen cross-over, resulting in declines in the overall performance. Consequently, the formation of hot spots may occur, and the impact of gas cross-permeation can intensify. Apart from thinning, chemical degradation also causes bubble formation on the surface and pores of the membrane. While chemical degradation alone doesn't generate cracks, it can accelerate crack development through two mechanisms: by merging bubbles on the surface of the membrane and by linking pores within the cross-sectional
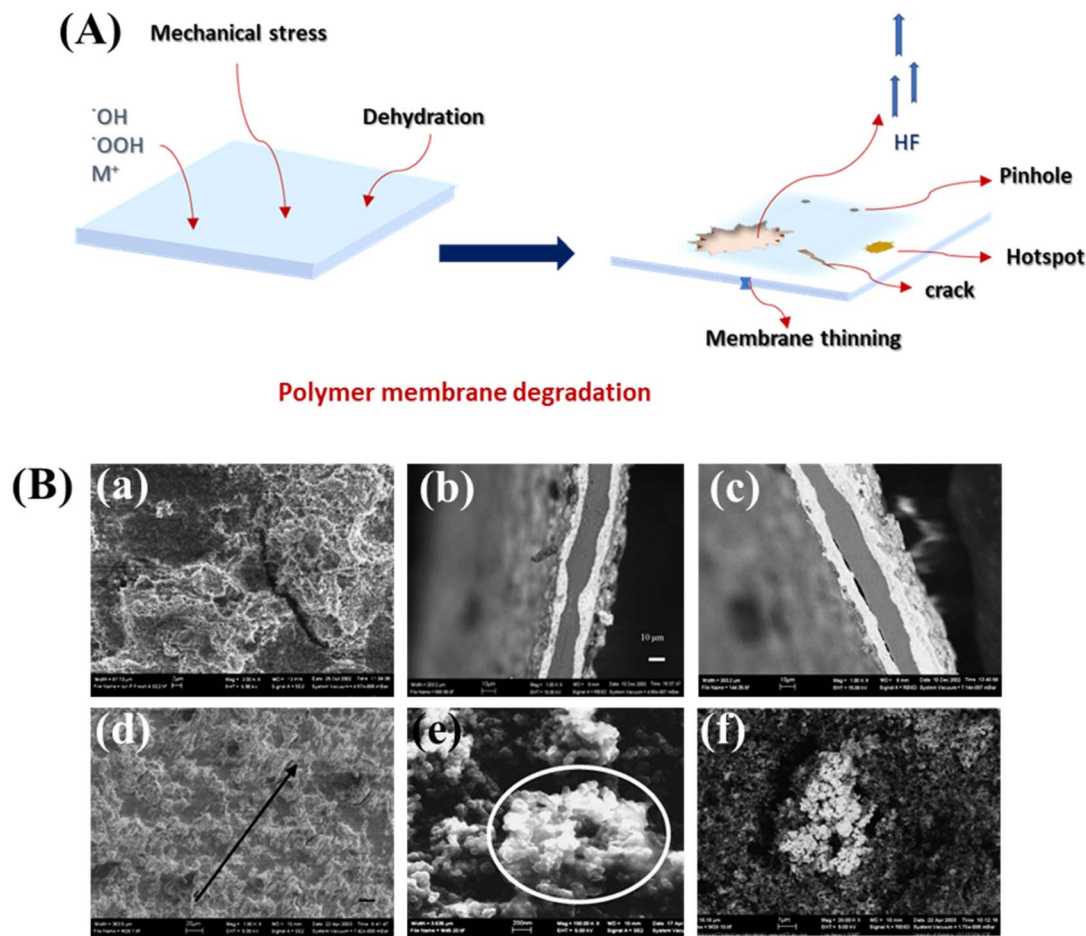
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Fig. 12 (A) Depiction of common factors contributing to the degradation and failure of polymer electrolyte membranes; and (B) SEM images of different types of membrane failures: (a) cracks (b) variations in thickness, (c) delamination, (d) catalyst orientation, (e) electrolyte clusters, and (f) platinum clusters, reproduced with permission from ref. 55 and 81, copyright© 2019 Elsevier Ltd.

area.[79] Nevertheless, this degradation phenomenon has been noted in electrolysis stacks only after approximately 15 000 hours of periodic operation at 1 A cm^{-2} and 80 °C. This implies that it is a relatively prolonged process, although operating at a higher current density could hasten the occurrence.[6] Research focusing on employing higher current densities (2 A cm^{-2} and above) observed increased hydrogen production. However, this was also found to accelerate the formation of hotspots in the case of electrolysers.[6] Higher current densities and operating temperatures also led to lower cell efficiency, with a voltage drop as the ohmic resistance and mass-transport resistance increased accordingly, which, in turn, raised the operational costs.[10,82–84]
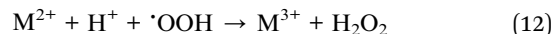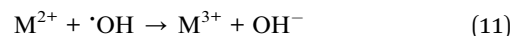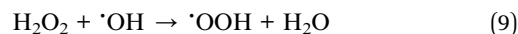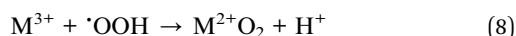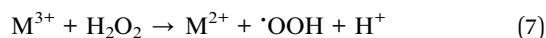
The chemical decomposition of PEMs is generally attributed to factors such as hydrogen peroxide formation, reactant gas crossover, platinum catalyst dissolution, recrystallization, and the presence of cationic contaminants. The onset of localized membrane degradation typically initiates at the cathode side and accelerates in dry and high-temperature conditions. A commonly used marker for identifying the degradation in fluorinated membranes is the presence of hydrogen fluoride and/or other chain fragments in the condensed discharge water.

In PEM fuel cells, the oxygen molecules can diffuse through the polymer membrane, especially if there are defects or if the membrane is not completely impermeable. As these oxygen molecules reach the anode, incomplete reduction can occur during the electrochemical reactions. This incomplete reduction process contributes to the formation of free radical species that are very reactive, such as HO$^{•}$ and HO$_2$$^{•}$. Incomplete reduction refers to the situation where oxygen molecules are not fully converted to water during the electrochemical reactions at the anode. The incomplete reduction often leads to the formation of active radicals, particularly HO$^{•}$ radicals, on the catalyst surface. Overall, the rate of membrane degradation is directly linked to the extent of gas cross-over, where more significant amounts of oxygen reach the anode. This incomplete reduction process is evidenced by detecting hydrogen peroxide (H$_2$O$_2$) in the product water. H$_2$O$_2$ is a by-product of ORR and hence indicates the presence of active radicals in the system. Increased gas cross-over intensifies the incomplete reduction process, resulting in higher concentrations of active radicals and accelerating membrane degradation.[85] The standard operating temperature of a fuel cell falls within the range of 60 to 80 °C.[12] Nevertheless, ongoing research is focused on

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developing PEMFCs that can be operated at temperatures surpassing 100 °C. However, operating at higher temperatures comes with certain drawbacks, such as increased component degradation, corrosion, and low humidity, which can lead to membrane degradation.

In fuel cell/electrolysis cell systems, stainless steel is often used for end plates due to its relatively high durability and corrosion resistance. However, over time, metal ions from stainless steel can leach into the system.[86,87] The presence of these metal ions, particularly in the form of contaminants, can speed up the degradation of PEM in cells. A proposed pathway for the degradation process involves a Fenton reaction, which is a type of chemical reaction involving the generation of hydroxyl radicals ($^{\bullet}$OH) from hydrogen peroxide (H_2O_2) in the presence of metal ions, typically transition metals like iron (Fe). These highly reactive hydroxyl radicals can initiate chain reactions that lead to the breakdown of the polymer chains in the membrane. The free radicals, particularly hydroxyl radicals, generated through the Fenton reaction are highly reactive species. In fuel cells, these radicals can initiate oxidative reactions that contribute to the deterioration of the polymer structure in the membrane. The eqn (6)–(12) below show the general pathways leading to the formation of free radicals from metal ions, where M is the metal ion.

$$M^{2+} + H_2O_2 \rightarrow M^{3+} + {}^{\bullet}OH + OH^- \tag{6}$$

$$M^{3+} + H_2O_2 \rightarrow M^{2+} + {}^{\bullet}OOH + H^+ \tag{7}$$

$$M^{3+} + {}^{\bullet}OOH \rightarrow M^{2+}O_2 + H^+ \tag{8}$$

$$H_2O_2 + {}^{\bullet}OH \rightarrow {}^{\bullet}OOH + H_2O \tag{9}$$

$$M^{3+} + e^- \rightarrow M^{2+} \tag{10}$$

$$M^{2+} + {}^{\bullet}OH \rightarrow M^{3+} + OH^- \tag{11}$$

$$M^{2+} + H^+ + {}^{\bullet}OOH \rightarrow M^{3+} + H_2O_2 \tag{12}$$

Another pathway for generating free radical species is the interaction between hydrogen and oxygen at the surfaces of a platinum catalyst, resulting in the formation of highly reactive species such as $^{\bullet}$OH, $^{\bullet}$H, and $^{\bullet}$OOH, which can contribute to membrane degradation. This process may occur on either the anode or cathode catalytic sites, depending on the current densities. The degradation of the main chain through the unzipping mechanism is considered one of the most common types of membrane degradation. During the polymerization of perfluorinated membrane materials, the end product acquires carboxylic acid end groups (–COOH) by using persulfate initiators. These end groups are prone to attack by any free radicals in the system. The $^{\bullet}$OH species can undergo a reaction with the –COOH group, resulting in the formation of carbon dioxide and water. This process leaves an active radical site on the main chain (see Scheme 1). The newly formed active chain component can react with additional $^{\bullet}$OH, initiating a series of chain-stripping reactions. These reactions culminate in substantially degrading the main-chain backbone.[88,89]

A predominant degradation pattern is found in the side chains, primarily due to more reactive linkages (see Scheme 2). The hydroxyl radical can attack the C–S bond in the sulfonic
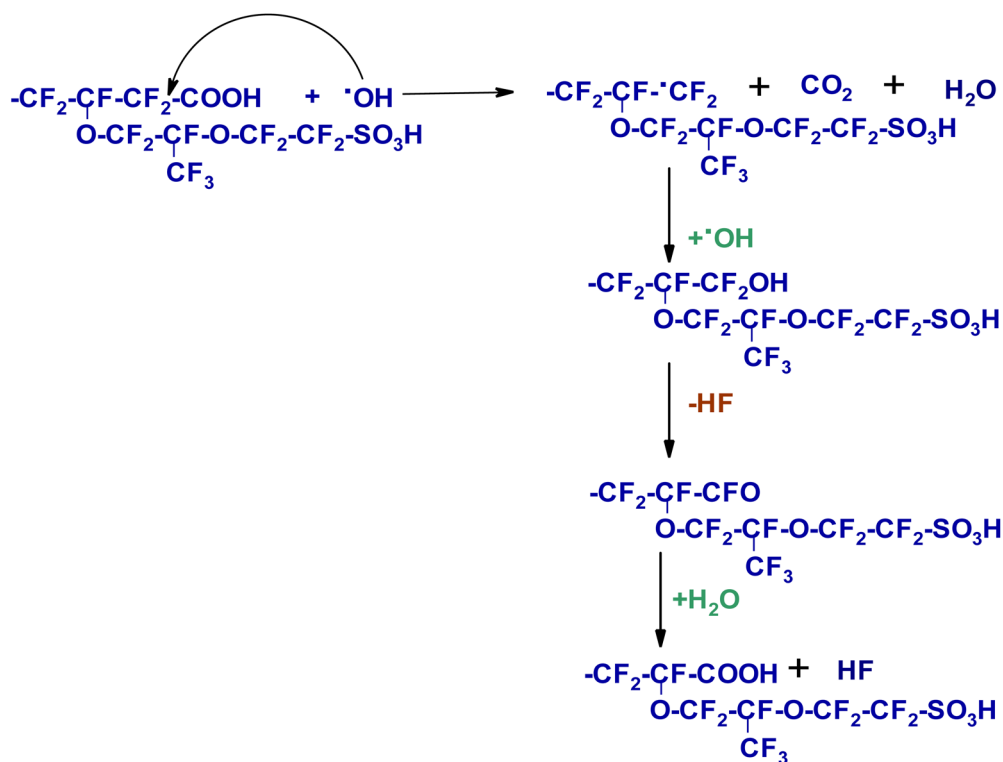
Scheme 1 The degradation mechanism in the main chain leads to the unzipping of the polymer molecules.
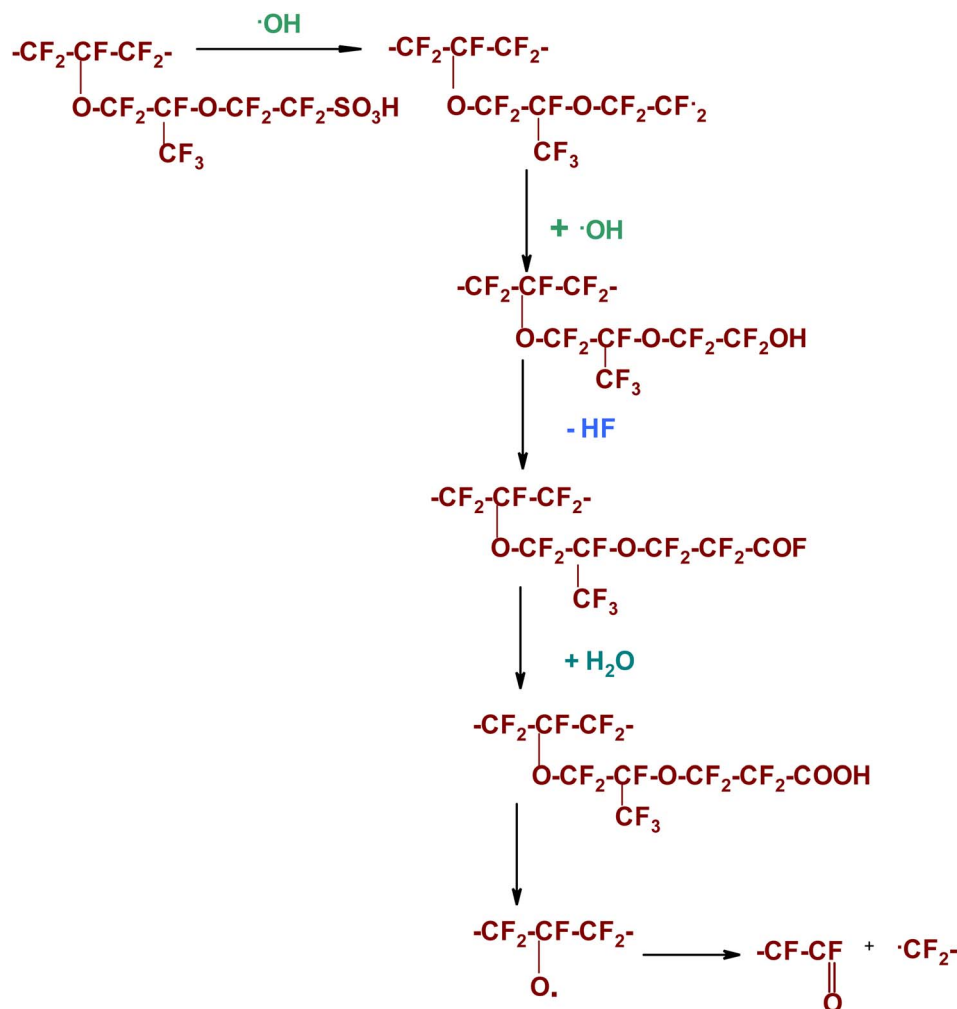
Scheme 2 The degradation mechanism in the side chain leads to unzipping of the chain.

acid end-group, leading to the generation of an active radical site in the side chain and the release of molecules such as water and sulfur trioxide. This active radical site formed in the side-chain can undergo further loss of molecules by reacting with more ˙OH, releasing hydrogen fluoride as a by-product. Another pathway involves free radicals attacking the C–F bond, either in the main chain or side chains.[58,89]

Temperature variations can also affect the degradation rate. When a Nafion membrane is immersed in water exceeding 110 °C, it undergoes a complete loss of mechanical integrity, leading to extensive disintegration. Furthermore, heightened membrane swelling can result in substantial dimensional alterations and the creation of internal constraints, diminishing its mechanical strength. These factors collectively contribute to the membrane's perforation. Research indicates that insufficient water supply to the anode causes localised membrane drying, exacerbating the issue. The challenge intensifies with increasing current density, making it increasingly challenging to provide adequate liquid water to the anode. Consequently, the electrical resistance in this region of the membrane rises, triggering a temperature increase due to the

Joule effect.[6] Another contributing factor to membrane breach is the mechanical stress caused by relative humidity (RH) cycling. To comprehensively address this issue, it is crucial to investigate the alterations in mechanical properties, particularly strength, arising from chemical aging and RH cycling.

5. Mitigation strategies

As discussed before, the performance of electrolytic cells influenced by factors like design, assembly, operational conditions, material degradation, and impurities. While performance degradation is inevitable as it is a natural consequence during the lifespan of a cell, an understanding of degradation mechanisms will help to minimize the rate, thereby enhancing the reliability, durability, and stability of these devices. Furthermore, ensuring the optimal performance of hydrogen cells requires a comprehensive approach to address potential contaminants. Purifying the hydrogen feedstock to fuel cells is a critical first step, involving advanced techniques such as pressure swing adsorption, membrane separation, or chemical scrubbing to guarantee high-purity hydrogen. In addition, to

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protect catalysts from poisoning by contaminants like CO, NO_x, and SO_x, it is essential to develop and employ more resistant catalysts, potentially utilizing alloyed catalysts or appropriate coatings that prevent the adsorption of these harmful substances. Improving membrane technology is equally vital, focusing on resistance to fouling and chemical attack by contaminants such as H_2S, NH_3, and metals. These may involve the development of new membrane materials or coatings with repellent properties against the contaminants. Regular maintenance and monitoring practices, including scheduled cleaning and replacement of affected parts, are also crucial for combating the impact of contaminants on the hydrogen cell. Table 2 below outlines the common challenges encountered during the operation of PEMFCs and PEMWEs, along with the corresponding mitigation strategies.

Additionally, implementing contaminant detection systems can provide real-time awareness of their presence in the hydrogen feed or within the cell. This also allows for prompt corrective actions to be taken before contaminants can cause significant damage. Design modifications, such as specialized flow channels or filters strategically placed to capture contaminants before they reach sensitive cell areas, can also minimise their impacts. By integrating these measures, the adverse effects of contaminants on electrolytic cell performance can be significantly reduced, ensuring more reliable and efficient operation over time.[90] A detailed discussion of the mitigation strategies mentioned in the table above is provided in the following sections.

5.1 Strategies for mitigating general failures in fuel cells and electrolysers

In the context of PEMFCs, it is crucial to maintain an optimal water flow within the system to ensure effective proton conductivity while mitigating ohmic losses. Adequate hydration of the membrane in a fuel cell stands as a fundamental goal for fuel cells utilising PFSA-based membranes. A promising approach to address this challenge involves the utilization of thin membranes with a thickness of fewer than 50 micrometers. Such thin membranes can facilitate a homogeneous water flow throughout the system, thereby optimising proton transport. It is critical to avoid operating at high current densities and abrupt, intermittent operating cycles, as rapid changes in load and voltage can significantly compromise the structural integrity of the MEA.[58,61] Another crucial approach involves reducing impedance within the cell. Reducing the resistance of the conducting elements within the cell system can be achieved through several strategies. This includes enhancing bulk electrical conductivity, minimising corrosion of components, and ensuring robust interface contacts by implementing gas diffusion layers between catalyst layers and bipolar plates. These measures collectively serve to minimise ohmic losses, thereby enhancing overall efficiency. Catalyst layers often utilise perfluoro sulfonic acid (PFSA) ionomers to enhance the triple-phase boundaries (active reaction sites). Applying a thin ionomer layer on the catalyst layer can enhance the proton transport to the PEM. However, this can affect the electronic conductivity and

mass transport, especially at lower catalyst loadings and high current densities. This leads to a reduction in O_2 concentration near the Pt surface (cathode). To address this, various strategies have been proposed to improve local O_2 transport. These include modifying the ionomer backbone to increase free volume, reducing sulfonic acid groups, and restricting their mobility by shortening the side chains. Other approaches involve replacing sulfonic acid groups with less adsorptive alternatives and tailoring these changes to the lower water content in catalyst-layer ionomers compared to bulk membranes.[13]

Furthermore, to limit platinum agglomeration in the cathode, it is beneficial to purge the anode channel with air. This proactively helps to maintain the integrity and performance of the cathode, thus contributing to the prolonged functionality of the fuel cell system. Furthermore, by employing a multifaceted approach involving membrane hydration, component optimization, and operational techniques, fuel cell systems can effectively manage proton conductivity while mitigating undesirable failure conditions, thus maximizing their performance and longevity. Running PEMFCs on hydrocarbon reformate may lead to the exposure of the anode catalyst layer to carbon monoxide (CO). To mitigate this, alloying the platinum catalyst can decrease CO coverage. In addition, maintaining the saturated vapor pressure of the feed during operation can be achieved by thinning the catalyst layer. Furthermore, raising the cell temperature can help in this regard. Introducing oxygen into the hydrogen feed is also another strategy to oxidize CO. Alternatively, modifying the gas diffusion layer with elements such as cobalt, iron, and copper can facilitate CO oxidation. These methods serve to enhance the performance and durability of PEFCs when operating with hydrocarbon reformation, ensuring efficient and reliable energy conversion. Furthermore, optimal catalytic supports can be carefully selected to reduce oxidation rates effectively. One proposed technique to enhance thermal stability involves graphitising carbon supports. Additionally, maintaining fixed current densities and flow rates during operation can help prevent the processes leading to carbon corrosion mentioned earlier. Introducing a microporous layer consisting of densely packed carbon particles held together by PTFE has been found to have several benefits. It reduces overall ohmic loss, mitigates electrode flooding, and enhances the mechanical stability of the MEA. Additionally, PTFE can be incorporated into gas diffusion layers (GDL) to impart hydrophobicity, thereby preventing cell flooding.[13] Fig. 13 summarises the important strategies that can be adopted for enhancing cell performance and minimising component failures of PEMFCs and PEMWEs.

To reduce the risk of flooding in PEM fuel cells, it is beneficial to steer clear of sharp corners in the flow-field path, as this can impede the passage of water. In this context, gas diffusion flow channels (GFCs) play a crucial role in fuel cells by effectively managing reactants and facilitating water removal. Extensive research has been conducted on various GFC designs, including parallel, serpentine, pin-type, and porous media flow fields, with channel dimensions typically ranging from 0.1 to 1 mm. By designing pathways with smooth curves instead of sharp angles, the likelihood of flooding decreases. Introducing

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Table 2 Challenges in electrolytic cells and their mitigation strategies

Sr no	General challenges	Mitigation strategies
1	Catalyst layer failures • Particle dissolution, detachment, agglomeration, Ostwald ripening • CO poisoning	• Avoiding high cell voltage or OCV, as well as frequent load and voltage changes • Maintaining optimal humidity and temperature suitable for operation, depending on the type of membrane used • Preventing chloride impurities from entering the system • Optimizing and homogenizing particle size to achieve a uniform catalyst layer distribution • Preventing carbon corrosion can mitigate particle detachment • Lowering water content in the CL can reduce Pt leaching, prevent carbon corrosion, and also enhance efficiency • Modification of the CL morphology by introducing additives like nanotubes, nanofilms, *etc.*, can enhance the structural reinforcement • Traditional Pt/C catalysts can be replaced with de-alloyed Pt–Cu core–shell type catalysts, which enhances the durability Using carbides, nitrides, or oxides of metals such as titanium, molybdenum, and tungsten as the support may enhance durability and resistance to corrosion • Oxidizing CO by passing O_2 in the feed, also by introducing H_2O_2 in the anode gas supply can recover catalyst from CO poisoning • A higher cell temperature can reduce CO adsorption on Pt catalysts; however, the upper limit depends on the temperature range suitable for the specific type of PEM used
2	Corrosion of the metallic parts of the cell assembly (PTL, BPPs, carbon support)	• Avoiding high cell voltage or OCV, as well as frequent load and voltage changes • Avoiding thermal and mechanical stresses • Using top-grade materials: graphite, alloyed stainless steel, *etc.* that are less corrosive • Platinization of the PTL and nitridation of BPPs ensure the formation of a protective layer on the surface, which protects against corrosive triggers. • Adding inert oxides like SnO_2, TiO_2, Ta_2O_5, and Nb_2O_5 to catalyst support enhances its stability in oxidative environments • In PEMFCs, hydrogen starvation must be avoided to prevent extremely high cathodic potential, which accelerates the corrosion of metallic components
3	Water flow problems (flooding/starvation/*etc.*)	• Avoid over-humidifying the membranes and operating at high current densities. • Operating at higher temperatures, suited for the type of PEM used, ensures a higher evaporation rate of liquid water, which can help lower flooding • Use of thinner membranes lowers water retention, facilitating faster water transport from CLs to GDLs • Ensuring homogenous water flows through the system by arranging the inlet and outlet channels in an interdigitated pattern, where the pathways end at the CL • Optimise reactant flow channel designs to avoid obstructions such as sharp angles and longer pathways • Anode water removal by forcing liquid water through membrane from cathode to anode reduces cathode flooding. • Adding a microporous layer (MPL) to the GDL improves the contact between the CL and GDL, which in turn improves water management by enhancing water removal from the catalyst layer • Avoiding operation at high current densities and pressures can reduce water starvation in electrolysers
4	Bubble formation	• Optimizing the ionomer content in the catalyst layer (CL) affects its porous structure, thickness, and distribution of triple-phase sites, thereby enhancing bubble evolution and transport efficiency • Use of superaerophobic CL facilitates formation of smaller bubbles that can be released much easily • Using porous PTLs can manage bubble transport more effectively • Optimising flow channel designs by adopting serpentine and parallel flow field types, with shorter paths can enhance bubble movement and removal. • Application of external physical fields, such as magnetic, ultrasonic, and gravitational fields, has been shown to enhance bubble evolution and transport
5	Degradation of GDLs	• Optimising compression pressure can reduce degradation due to mechanical stress • Coating the GDL with a PTFE layer enhances its mechanical and chemical stability while also making it hydrophobic, reducing excess water buildup • Doing corrosion-resistance treatments

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Table 2 (*Contd.*)

Sr no	General challenges	Mitigation strategies
6	Ohmic losses	• Minimizing ionomer degradation can prevent the release of fluoride ions, which contribute to issues such as GDL degradation, corrosion, and damage to other components • Minimising corrosion of the components can enhance electrical conductivity • Robust interface contacts ensure less contact resistance and strong interfacial bonding of the MEA • Enhancing bulk electrical conductivity by employing metallic parts that are highly conductive • Adding a microporous layer also enhances proper contact between CLs and GDLs • Optimal humidity and temperature specific to the PEM are also important for improving electrical conductivity
7	Degradation of PEM	• Minimising local drying can prevent the formation of hot spots • Ensuring uniform membrane thickness can reduce mechanical failures, thereby controlling hydrogen crossover issues • Ensuring high purity of the reactants prevents free radicals and other species from entering the membrane, which can trigger degradation • Ensuring uniform temperature and humidity control measures is vital in preventing membrane dehydration, shrinkage and rupture • Reducing voltage fluctuations is critical in preventing degradation and formation of pinholes • Hydrogen starvation must be avoided in the case of FCs to prevent damage • Reducing input feed pressure is critical in some cases of PEMWEs, however, for optimal performance, a certain level of pressure is typically necessary to ensure high gas production rates and effective electrolysis. • Optimising ionomer content is crucial in preventing hydrogen crossover and maintaining adequate proton conductivity of the membrane
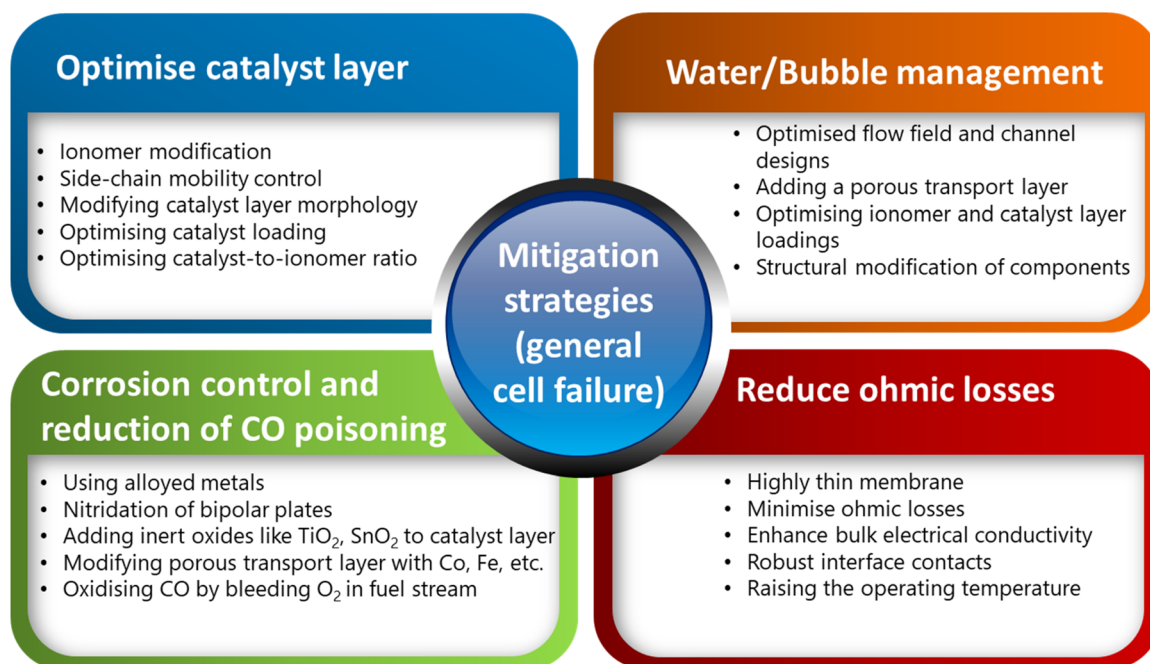
Fig. 13 An analysis chart shows some common mitigation strategies for enhancing the overall efficiency of PEMFCs and PEMWEs.

shorter pathways can also alleviate pressure drops in the flow of liquid water. This adjustment, in turn, ensures that water flows more efficiently through the system without encountering significant resistance. Among these designs, porous media flow fields have attracted increasing attention and are utilised in some recent hydrogen-operated electric vehicles. These

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channels enhance heat removal and electric conductance, especially by employing materials such as metal and carbon foams, which are renowned for their high porosity, corrosion resistance, and conductivity.[13] Water generated on the cathode side can also play a crucial role in keeping the membrane adequately hydrated. When managed correctly, this water can naturally diffuse and pass through the membrane by convection, thus assisting in maintaining its hydration levels. Therefore, proper redirection of water formed on the cathode side can contribute to some degree to preserving the state of hydration of the membranes within the fuel cell. As mentioned in the previous paragraph, incorporating a microporous layer into the interface of the gas diffusion layer can additionally reduce the likelihood of flooding. Generally, the accumulation of ice in cells can pose a problem, especially when there's an excess of water present. To prevent issues, it is essential to keep the cell's temperature above freezing. One solution is to remove excess water before starting cell operation through gas purging. Generally, optimising the compression pressure during cell assembly can help reduce mechanical stress on GDLs to some extent.[69] Furthermore, increasing the hydrophobicity of GDL materials is an effective way to ensure proper water management, which is also critical for preventing the dissolution of GDLs in water. Carbon corrosion in GDLs can be mitigated by applying PTFE coatings and performing corrosion-resistant treatments.

Reducing the corrosion of bipolar plates can often be achieved by employing top-grade materials with excellent corrosion resistance, high electrical conductivity, and strong mechanical properties. Materials such as graphite and stainless steel are viable options. Graphite, being a standard choice, generally offers high conductivity and corrosion resistance, although it is prone to mechanical failures and can be expensive. On the contrary, stainless steel possesses strong corrosion resistance and mechanical properties. However, its electrical conductivity is hindered by oxide layers forming on its surfaces. These characteristics of stainless steel can be adjusted by altering the chromium content in various grades of steel alloys.

PEMWEs typically operate under harsher conditions than PEMFCs, such as higher pressures, current densities, temperatures, and power load cycles. These factors accelerate component degradation. However, selecting materials with high chemical, mechanical, and thermal stability can mitigate this degradation to some extent. SPEEK and PBI membranes, for example, demonstrate excellent mechanical and chemical properties. Additionally, nitriding bipolar plates (BPPs) have been found to minimise corrosion, protecting the gas diffusion layer (GDL) and preventing pinhole formation.[91] Adding inert oxides like SnO_2, TiO_2, Ta_2O_5, and Nb_2O_5 to the catalyst support reduces corrosion and lowers costs by decreasing noble metal loading.[92] Studies also suggest that increasing catalyst loading can reduce the MEA degradation rate. Modifying catalyst morphology can address other challenges; for instance, platinum nanotubes are less prone to dissolution, and nano-thin films help prevent agglomeration.[3]

Bubble management in PEMWEs can be enhanced by optimizing the catalyst layer structure, including thickness, porosity, and triple-phase boundary distribution. Controlling ionomer content is crucial for regulating pore size. Structural modifications such as super wetting electrode designs, 3D ordered layers, or magnetized catalysts improve bubble management. Flow field designs, particularly parallel configurations, also aid in the efficient removal of bubbles.[62]

5.2 Strategies for mitigating membrane degradation in fuel cells and electrolysers

Firstly, mechanical failures of membranes can be significantly reduced by minimising local drying. Carefully designing the flow field structure is also essential in this case. Additionally, reducing voltage fluctuations, such as overshoots and undershoots, can, in turn, minimise membrane degradation and pinhole formations.[93] Effective heat transfer and temperature control play a crucial role in preventing membrane damage. A notable issue with fluorinated membranes is that, at temperatures above 100 °C, water evaporation can occur, leading to membrane dehydration. This dehydration can cause various problems, such as cracks, pinholes, hotspots, and degradation of the conductive side chains. These issues can be mitigated to some extent by modifying the polymer structure through the incorporation of fillers such as metal oxides or carbon nanotubes,[94,95] using non-fluorinated membranes, or employing polymers with shorter side chains.[96]

Other critical factors must be considered, including reducing the feed input pressure and maintaining uniform moderate temperature and humidity conditions throughout the operation. In operational applications requiring high feed pressure, it is essential to balance power requirements with input pressure. During load demand variations, hydrogen starvation in fuel cells must be avoided to prevent damage to the membrane and associated catalyst layers. Pressure management is equally critical for PEMWEs.[97] Temperature control is vital for protecting electrolyte membranes from dehydration, shrinkage, and rupture.[98] Therefore, implementing a thermal management subsystem is crucial to ensure uniform temperature distribution throughout the system. Another important aspect is maintaining optimal membrane humidity. In the case of PEMFCs, this can be partially achieved by passing the feed gases through external humidifiers set to the required temperature.[97] It is also essential to maintain uniform membrane thickness to effectively manage mechanical failures. Secondly, addressing issues related to corrosion is of paramount importance. The impurities can originate from BPPs, membranes (specifically Nafion®), and sealants, thus highlighting the diverse range of impurity sources and associated potential contaminants. Here, the mitigating steps involve minimizing the corrosion of metallic parts, such as BPP, in the system and preventing the degradation of cell parts like silicone seals. To further enhance the system's longevity, it is also crucial to avoid the intake of metal ion impurities and air pollutants in the cell. It is noted here that air contains contaminants such as N_2, NO_x, SO_x, NH_3, and ozone. In addition, reformate hydrogen, another source, includes CO, CO_2, H_2S, ammonia, and methane. By carefully managing these factors, one can ensure the optimal

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Fig. 14   An analysis chart shows some common mitigation strategies for minimizing membrane degradation in electrolyser and fuel cells.

performance and durability of the membrane system.[58] Fig. 14 shows some common strategies that can be used to reduce membrane degradation. Therefore, ensuring the durability of associated components, such as catalyst layers, GDLs, sealants, BPPs, and other components, is crucial for maintaining the integrity of the PEM. The degradation of these components can release free radicals into the system, which may attack the membrane, leading to its degradation and eventual failure. The feed gases or water must also be highly purified to prevent impurities from entering the system. Impurities can cause corrosion and degradation of cell components, and the resulting degradation of products can, in turn, contribute to the deterioration of the membrane assembly.

Employing proper GDLs can help mitigate pinhole formation in the membrane and associated hydrogen crossover by effectively managing the heat generated by electrolytic reactions within the cells.[79] Reducing hydrogen crossover also lowers the degree of chemical degradation of the membranes by decreasing the probability of free radicals, which trigger chemical degradation, being carried through the membranes.[99] Ensuring close contact and even assembling of the MEA can significantly reduce these issues. Hydrogen crossover also

depends on membrane thickness, surface morphology, and ionomer concentration. Optimizing these factors can again reduce the crossover rate.[91]

Efforts to improve Nafion's properties generally focus on enhancing mechanical stability, typically through controlling membrane swelling, temperature, and reinforcement with inert matrices. Composite membranes, combining Nafion with inorganic fillers, are also employed to enhance chemical and temperature stabilities. However, due to matrix substitution, these enhancements may reduce proton conductivity and water vapor permeability.

To address Nafion's high methanol crossover, fine inorganic particles can be added to the polymer matrix, or other polymers with different properties can be introduced within the void space. Two approaches currently exist for this modification: semi-interpenetrating polymer networks (semi-IPN) and interpenetrating polymer networks (IPN). However, high methanol permeability remains an issue, especially in proton exchange membranes in direct methanol fuel cells (PEM/DMFCs) that use Nafion membranes. Here, cost is also a significant concern due to the complex manufacturing process of Nafion membranes. Therefore, alternative approaches to enhance performance

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while reducing costs are actively pursued in the field of hydrogen cell technology.[24]

A recent study[100] examined the challenges of commercialising large-scale high-pressure PEM water electrolysis and highlighted that issues such as gas cross-permeation, membrane degradation, and hydrogen embrittlement of metal components are underexplored. The study found that composite membranes, prepared through various methods, effectively reduce gas permeation. Chemical degradation can be mitigated by introducing free radical scavengers to eliminate free radicals. Additionally, anti-corrosion coatings with high chemical stability and good electrical conductivity can partially inhibit hydrogen embrittlement. Several recent studies have also utilised various modelling techniques to analyse the challenges of commercializing fuel cells and electrolysers, as well as to predict methods for mitigating these challenges.[97,101–104]

A recent review[105] of fuel cell systems explored the challenges of implementing this technology in electric vehicles, identifying hydrogen storage as a significant obstacle. The review proposed that fuel cell/electrolyser hybrid technology could offer a potential solution. Additionally, storing hydrogen in the form of hydrogen-based compounds, such as methanol, was highlighted as a way to simplify storage. The review also discussed strategies to enhance fuel cell performance, including increasing cell temperature, raising reagent pressure, using highly conductive electrodes, employing thinner membranes, and optimizing humidity levels.[105–107]

6. Conclusions and future perspectives

Hydrogen has emerged as a beacon of promise in the realm of alternative fuels, driven by its unique characteristic of producing water as the sole by-product when oxidised to produce power. This feature positions hydrogen as a standout candidate in the pursuit of carbon-neutral energy sources, as its use does not contribute to greenhouse gas emissions or environmental pollution. Embracing hydrogen holds the potential to revolutionize energy across multiple sectors, paving the way for a cleaner and more sustainable future. The versatility of hydrogen extends beyond its role as a fuel to incorporate applications in fuel cells for electricity generation. Its abundant potential to be produced using electrolysers powered by natural sources, such as solar and wind, further strengthens its sustainability credentials. However, fully realising hydrogen's potential faces challenges related to commercialising polymer electrolytic fuel cells and hydrogen-based technologies.

The durability of PEMFCs and PEMWEs is a crucial factor hindering the commercialisation of automotive and stationary applications, along with concerns regarding cost and hydrogen storage. Substantial improvements are needed to make electrolytic cells more durable for automobile applications. For example, PEMFC systems must be upgraded to increase operating time to 30 000 hours in the coming years.[108] Therefore, commercialisation of hydrogen-based cells is still a major challenge owing to the issue of failure and degradation associated with various components within the cell system. The primary accelerators of PEM degradation include elevated partial pressure of reactants, reduced RH, increased temperature, and heightened voltage. Consequently, addressing these factors is crucial in mitigating degradation and enhancing the longevity of fuel cell systems. In PEMFC stacks, the serial cell arrangement makes them vulnerable to catastrophic failure if a single cell fails, and durability is difficult to achieve due to varying operating conditions. Similarly, in water electrolysers, performance declines over time due to electrode degradation, including catalyst sintering, membrane degradation, and ohmic losses, with the risk of gas mixing posing additional safety concerns. Contaminants like nitrogen, sulfur oxides, carbon monoxide, and ammonia can poison the catalyst and foul the membrane in hydrogen cells, leading to reduced performance and efficiency. Additionally, hydrogen crossover within PEM cells accelerates degradation, causing issues like membrane and catalyst layer damage, reduced efficiency, and increased risk of thermal degradation. Maintaining high-purity feed and controlling humidity levels are crucial to optimizing performance and longevity.

The selection of materials for the polymer electrolyte membrane, a critical component in PEM cells, holds immense importance in addressing these challenges. Currently, Nafion stands out as the predominant choice due to its outstanding conductivity and favourable chemical and mechanical properties. However, despite its favourable characteristics, widespread adoption of Nafion faces obstacles due to its high production costs and environmental implications. The stability of the carbon–fluorine (C–F) bond in fluorine-containing materials, such as PFASs, which form the basic structure of Nafion and other perfluorinated membranes, makes them resistant to decomposition in the environment, leading to significant environmental concerns. PFASs are water-soluble and can easily contaminate land and water sources, causing risks to human health and ecosystems due to their persistence and potential for bioaccumulation. The presence of PFASs in water systems can cause long-term exposure for aquatic life and humans, leading to ecological disruptions and health risks. Additionally, PFASs are difficult to remove through water treatment processes, contributing to the contamination of drinking water, particularly near industrial areas. Addressing these issues requires developing environmentally friendly alternatives and implementing strict regulations to mitigate the impact of PFASs. Perfluorinated polymeric membranes, which are currently used in PEM cells, face two primary challenges. Firstly, they struggle to maintain high protonic conductivity when water content is low, particularly under conditions of high temperature and low humidity. Secondly, ensuring durability under these conditions remains a significant obstacle, as existing sulfonic acid-based materials exhibit low conductivity under low hydration conditions.[109]

To enhance electrolytic cell performance, it is crucial to develop novel polymer-based membranes that operate effectively at higher temperatures, require less hydration, and are made from environmentally friendly materials. Improving proton conductivity at lower hydration levels and elevated temperatures can be achieved by incorporating highly acidic

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functional groups or designing new polymer backbones. Polymers similar to Nafion, but without halogenated components, are being explored for their potential in fuel cells and electrolysers, focusing on reducing costs and environmental impact. Customising membranes for specific applications, such as varying degrees of hydration tolerance or resistance to contaminants, can further optimise performance for different types of fuel cells and electrolysers. In addition, developing membranes that work synergistically with other cell components, such as electrodes and catalysts, can improve overall system efficiency. Inorganic/organic hybrid materials such as polymer/clay nanocomposite hybrids are one of the promising candidates due to their excellent mechanical and thermal stability, along with low gas cross-permeation effects. Reinforcing polymers with fibres or nanoparticles can enhance their mechanical strength and durability, helping membranes withstand operation stresses, such as swelling and shrinking, due to hydration cycles. Reducing gas crossover, particularly of hydrogen and oxygen, is critical for both safety and efficiency, with nanocomposite membranes incorporating impermeable fillers like clays or graphene to achieve this without compromising proton conductivity. Chemical stability is also essential, as the membrane must resist degradation under oxidative and acidic conditions, extending its lifespan and maintaining efficiency over time. Cost-effective materials are vital for large-scale deployment, potentially through the use of cheaper raw materials or simplified fabrication processes. Using non-toxic, halogen-free materials can reduce the environmental footprint and make membranes safer for disposal and recycling. A proposed model connects PEM electrolysers and fuel cells into a single integrated system. In this setup, hydrogen produced by the electrolyser from feed water is supplied to the fuel cell. The water produced by the fuel cell during operation is then purified and cycled back into the electrolyser. This arrangement aims to create a self-sustaining power generator. However, further research is needed to confirm the feasibility of this system.

To address these challenges, there is a pressing need to develop novel proton-conducting membranes/components and structures capable of meeting the demands of higher-temperature systems. Researchers and industry stakeholders are actively exploring other similar or advanced materials to replace Nafion but with reduced costs and environmental footprint. This ongoing research and development effort is critical in advancing the commercial sustainability of hydrogen-based fuel cells and electrolysers, ultimately contributing to a cleaner and more sustainable energy future.

Abbreviations

BPP	Bipolar plates
CL	Catalyst layers
DMFC	Direct methanol fuel cell
DSC	Differential scanning calorimetric
GDL	Gas diffusion layers
GE	General electric
GFC	Gas flow channels
HER	Hydrogen evolution reaction
HDV	Heavy-duty vehicle
ICE	Internal combustion engine
IPN	Interpenetrating polymer networks
MEA	Membrane electrode assembly
MPL	Micro-porous layers
NO_x	Nitrogen oxides
OCV	Open-circuit voltage
OER	Oxygen evolution reaction
ORR	Oxygen reduction reaction
PEI	Polyethyleneimine
PEM	Proton exchange membranes
PEMFCs	PEM fuel cells
PEMWEs	PEM water electrolysers
PFAS	Polyfluoroalkyl substances
PFCA	Perfluorocarboxylic acids
PFSA	Perfluorosulfonic acid
PSSA	Polystyrene-sulfonic acid
PTFE	Polytetrafluoroethylene
PTL	Porous transport layer
RH	Relative humidity
SA	Sulfonic acid
SO_x	Sulfur oxides
SPEEK	Sulfonated polyether ether ketone
T_g	Glass transition temperature

Data availability

Data availability is not applicable to this article as no new data were created or analyzed in this study.

Author contributions

Malavika Arun: data collection, data curation, writing-original draft; Sarbjit Giddey: funding acquisition, supervision and conceptualisation, review, and editing; Paul Joseph: supervision, review, and editing; Dattatray S. Dhawale: funding acquisition, project administration, supervision, conceptualisation, data curation, review, and editing.

Conflicts of interest

There are no conflicts to declare.

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