Radiation-grafted anion-exchange membranes: the switch from low- to high-density polyethylene leads to remarkably enhanced fuel cell performance

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Herein we detail the development of a new high-density polyethylene (HDPE)-based radiation-grafted anion-exchange membrane (RG-AEM) that achieves a surprisingly high peak power density and a low in situ degradation rate (with configurations tailored to each). We also show that this new AEM can be successfully paired with an exemplar non-Pt-group cathode.

It is critical that anion exchange membranes (AEM), developed for use in anion exchange membrane fuel cells (AEMFCs), can support both high power outputs and in situ durability. The literature shows a dearth of options that have an acceptable combination of both, which is mandatory to push AEMFCs closer to widespread implementation and commercialisation.

In recent years, significant progress has been made in the development of AEMs generally, 1 and RG-AEMs specifically. 2 RG-AEMs produced via a high dose rate electron-beam modification of low density polyethylene (LDPE) films have helped to advance the field, by both allowing for an enhanced understanding on how AEMs behave in AEMFCs, as well as their ability to support high H2 fuel cell performances (> 1.0 W cm−2 at > 60 °C with non-Pt cathodes).3,4 This performance was possible because such LDPE-based RG-AEMs have high conductivities, and kinetically fast water transport characteristics (that extends the region of mass transport power losses to higher current densities). However, a limiting aspect of thin LDPE-based RG-AEMs films are their modest mechanical properties under stress (Table 1), which needs to be improved for commercial applications. In this respect, the use of high-density polyethylene (HDPE) was hypothesised to be a promising avenue of research, especially as prior reports suggest that HDPE can be used to make viable RG-AEMs5 using low dose rate gamma ray methods (Sproll et al.6(a) report low-dose rates lead to a lower concentration of longer grafted chains, which leads to poorer performance characteristics).

We report for the first time a HDPE RG-AEM fabricated using a high dose rate electron-beaming method (amenable to large batch production), with comparable properties to a similarly fabricated LDPE RG-AEM, but with improved mechanical properties, fuel cell performance, and operational stability.

Fig. 1 Degree of grafting (dog) of the poly(VBC)-grafted membranes made from the e-beamed LDPE and HDPE films that were stored (pre-grafted) at -40 °C for increasing periods of time. The dog (%) was calculated as: dog = 100 × (m m) / m (where m was the mass of the e-beamed pre-grafted film and m was the mass of the (pre-aminated) VBC-grafted membrane).7

Broader context: A primary motivation for the development of anion-exchange membrane (AEM) fuel cells (AEMFCs) is the broader range of sustainable, non-precious-metal catalysts that are feasible; if costs are lowered enough, AEMFCs would be deployable in a range of stationary power sectors (e.g. back-up and off-grid). However, as the performance of AEMFCs typically drop when Pt-based electrodes are replaced with non-Pt types, it is essential that the highest performing polyelectrolytes are developed, both membranes and ionomers (the latter incorporated to impart ionic conductivity in the electrodes). The findings with the high conductivity AEM reported herein will also be of interest to developers of AEMs for metal-air and redox-flow batteries, electrolyzers (both H2O → H2 and CO2 → high-value chemicals and fuels), and salinity gradient power.

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The methods detailed in our previous publications. 2,4,7 In the RG-HDPE AEMs were fabricated and characterised using 6 h grafting at 40 °C for LDPE and 4 h at 50 °C for HDPE. After the further addition of 1 % vol. octyl-2-pyrrolidone dispersant (Aldrich product 338729, no removal of any inhibitors) by vinylbenzyl chloride (VBC, mixture of 3- and 4-isomers, Sigma-Weighted e-beam irradiated films were grafted with e-beam). The e-beam irradiated films were then stored in a freezer at -40 °C until use. Weighted e-beam irradiated films were grafted with vinylbenzyl chloride (VBC, mixture of 3- and 4-isomers, Sigma-Aldrich product 338729, no removal of any inhibitors) by immersion in a N2-purged aqueous solution of VBC (5 % vol. with the further addition of 1 % vol. octyl-2-pyrrolidone dispersant): 6 h grafting at 40 °C for LDPE and 4 h at 50 °C for HDPE. After thorough washing with toluene and drying at 16 h at room temperature, the grafted membranes were weighed before being submerged in aqueous trimethylamine (45 % wt.) for 24 h. After subsequent multiple room temperature washings with ultrapure water (UPW, 18.2 MΩ cm), heating in UPW for 1 h at 60 °C, ion-exchange by immersion in aqueous NaCl (1 mol dm⁻³, immersion for 1 h in 3 × fresh solutions), and further washing in UPW (until no traces of free Cl⁻ remained), the Cl⁻ anion forms hydrated thickness, water-uptake, and OH⁻ conductivity. Both RG-AEMs (OH⁻ form) also similarly retained their conductivities of a flowing 100 % relative humidity N₂ atmosphere, 4 based on the method first reported by Ziv and Dekel. 8 Tensile properties of the Cl⁻ form RG-AEMs (errors within 25 %).

The key properties of the two RG-AEMs are summarised in Table 1. The HDPE RG-AEM exhibited very similar properties to the LDPE RG-AEM: ion-exchange capacity (IEC), hydrated thickness, water-uptake, and OH⁻ conductivity. Both RG-AEMs (OH⁻ form) also similarly retained their conductivities in a flowing humidified N₂ atmosphere at 80 °C (Fig. 2); the loss of conductivity was only (8 ± 1) % after 500 h for both. Comparing the Raman spectra before and after 500 h, Fig. 3 shows only minor RG-HDPE degradation, in good agreement with the reduction in conductivity during that time. However, where the HDPE-AEM appears much more desirable compared to the LDPE-AEM was its tensile properties. The break stress of the HDPE-AEM was 52 % greater than the LDPE AEM and it also showed an enhanced ability to stretch without failing. This was accomplished without having to increase the thickness of the final hydrated AEMs (achieved just by using a different substrate). The use of thin membranes is desirable as they enable fast water transport, meaning that a higher current density can be supported since high AEMFC performance relies heavily on the back-diffusion of water from the anode to the cathode. 9,11
**Fig. 4** H₂/O₂ AEMFC comparison tests (80 °C) for the RG-AEMs. Full details on the test conditions used are given in ref. 4 with the following summary data (the only variable in these tests was the RG-AEM used): PtRu/C anodes (50 % wt. Pt and 25 % wt. Ru) with a Pt-loading of 0.4 mg cm⁻²; Pt/C cathode (40 % wt. Pt) with a Pt-loading of 0.4 mg cm⁻²; catalyst inks contained 20 % wt. radiation-grafted ETFE-based anion-exchange ionomer powder (IEC = 1.90 ± 0.06 mmol g⁻¹) and were sprayed directly onto Toray TGP-H-60 (PTFE-treated) gas diffusion substrate; the H₂/O₂ gases were supplied at 1 dm⁻³ min⁻¹ at RH = 92 % with no back-pressure applied. This galvanostatic data was collected using a Scribner 850E fuel cell tester.

Figure 4 shows the beginning-of-life AEMFC performances at 80 °C for cells operated with HDPE- and LDPE-AEMs using identical electrodes and benchmark fuel-cell-grade Pt-based catalysts (using the Surrey group’s standard fuel cell testing protocols, which have been detailed extensively in recent publications²,⁴ and allows us to maintain a local database of relative performances of different AEMs). Despite the LDPE-AEM and HDPE-AEM exhibiting similar ex situ thicknesses, water uptakes, and conductivities, the HDPE-AEM yielded a significantly higher H₂/O₂ AEMFC performance: the HDPE-AEM exhibited a peak power density of 2.55 W cm⁻² at 80 °C (cf. 2.01 W cm⁻² for the LDPE-AEM) with zero back-pressure gas feeds (with only ca. 0.2 bar pressure drops).

This is a dramatic demonstration of the findings by Sproll et al. related to the development of RG-proton-exchange membranes (RG-PEM),¹² who reported that conductivities, water uptakes, and resulting fuel cell (PEMFC) performances strongly depend on the micro-structure of the ETFE-base films used. The use of nominally identical base ETFE films (from two suppliers), differing only in microstructure, resulted in critical differences in the final membranes: larger crystalline sizes led to enhanced RG-PEM conductivities and durabilities of the resulting PEMFCs. The only significant variable in our comparison experiments (Fig. 4) was the starting base-material used. DSC experiments show that the supplied HDPE film had a crystallinity of 57 % (cf. 70 – 80 % in supplier literature, HDPE is defined as having a low degree of branching), while the LDPE film had a lower crystallinity of 47 % (ca. 50 % in the supplier literature, LDPE is defined as having a high degree of branching): these initial differences are clearly highly significant to the performances of the resulting AEMFCs. This also serves as a warning: a correctly selected combination of electron-beaming dose-rate and supplier (grade/additives) of base-material will be critical to any final, reproducible commercial production effort.

As there is no significant difference between the OH⁻ conductivity or thicknesses of the HDPE-AEM and LDPE-AEM, we hypothesise that the improved performance is due to the enhanced water transport from the anode and cathode in the operating fuel cells⁹⁻¹¹ and this must be due to a change in the nanomorphology or microstructure between the LDPE- and HDPE-AEM. This hypothesis needs to be rigorously tested and so we are in the process of planning a series of quasi-elastic neutron scattering (QENS) and small-angle neutron scattering (SANS) experiments¹³ to probe the ion-dynamics and nanomorphology in much more detail (comparing the new HDPE-based RG-AEM to both prior-art LDPE-²,⁴ and ETFE-based⁷ RG-AEMs). These results will be reported in a future, specific research paper.

These high performances were independently reproduced at the University of South Carolina with their own benchmarking parameters. The anode was PtRu(2:1)/C with a 0.7 mg cm⁻² PtRu loading; the cathode was Pt/C with 0.6 mg cm⁻² Pt loading; both catalyst layers contained 20 % wt. radiation-grafted ETFE-based anion-exchange ionomer powder (IEC = 1.24 ± 0.06 mmol g⁻¹) and were sprayed onto Toray TGP-H-60 carbon-paper GDEs containing 5 % wt. PTFE wet-proofing. The peak power density was 2.5 W cm⁻² with a cell at 80 °C operating with H₂/O₂ reacting gases. A 2.4 W cm⁻² peak power density was also achieved in the same cell at an advantageously lower temperature of 70 °C (gases supplied at 1 dm⁻³ min⁻¹ with an anode dew-point of 60 °C and cathode dew-point of 68 °C). The peak power density of this latter cell at 70 °C with CO₂-free air at the cathode was 1.1 mW cm⁻².

Given the above data, a HDPE-AEM membrane-electrode assembly (MEA) was tested over 440 h at 600 mA cm⁻² constant current discharge in a H₂/air(CO₂ free) AEMFC at 70 °C (Fig. 5, caption details the test parameters used). Both the cell voltage and area specific resistance (ASR) were recorded throughout.
The degradation rate of 790 ± 10 µV h⁻¹ for comparison (Fig. 5). This shows a much more rapid degradation rate of 790 ± 10 µV h⁻¹ (95% confidence intervals, n = 3530 data points, R² = 0.55), which is why we terminated the test early. The failure mechanism with the LDPE-AEM clearly involved an increase in in situ ASR (38 → 52 mΩ cm²).

With regards to commercialisation, catalyst cost and sustainability are important factors that must be considered when operating these devices. Due to the generally lower intrinsic activity of non-Pt catalysts for the oxygen reduction reaction in alkaline media, there is typically a 30 – 50% performance when they are applied to AEMFC cathodes. For most non-RG-AEMs, non-Pt-catalyst containing AEMFCs are routinely reported with power densities below 1 W cm⁻².14,15 We replaced the Pt/C cathode from Fig. 4 with an exemplar non-Pt-group cathode (BASF Ag/C, 40% wt. Ag, 0.85 mg cm⁻² Ag loading). As expected, the performance decreased when Ag/C was used (Fig. 6) with a 32% lower peak power density (1.72 W cm⁻²); this is a notable result given that the cost of Ag is currently only 2% of the cost of the Pt (£0.38 g⁻¹ vs £19.79 g⁻¹, respectively).16 The predominant causes of the lower performance were poorer electrode kinetics (V drop at low current density) and the earlier on-set of mass transport limitations (stemming from the thicker catalyst layer required).

These results represent a notable improvement over other recent important literature reports.4 For example, Maurya et al. tested a new polyfluorene quaternary ammonium ionomer with a 30 µM thick TPN AEM (a partially fluorinated polyphenylene-type with long alkyl side-chain),17 which achieved a H₂/O₂ AEMFC performance at 80 °C of 1.5 W cm⁻² using similar catalysts; this AEMFC type showed high stability for 250 h when discharged at 0.6 V at 80 °C, after which cell voltage degradation was observed. There has also been a recent conference report by Wang and Yan et al. of a poly(aryl piperidinium) (PAP) AEM that shows no degradation when immersed in aqueous KOH (1 mol dm⁻³) for 2000 h at 100 °C;18 this AEM yielded an AEMFC performance of 920 mW cm⁻² at 95 °C with low-Pt electrodes, which showed in situ durability over a period of 300 h when discharged at 500 mA cm⁻².19 This theme of producing AEMs that are chemically stable in extreme alkali environments follows on from the work by Holdcroft et al.,20 who showed that a poly(arylène-imidazolium) AEM showed minimal degradation in aqueous KOH (10 mol dm⁻³) at 100 °C. For a comprehensive review of AEMFC performances achieved in other studies published from before 2018, please refer to the review by Dekel.21 It is clear that there is a dearth of 500+ h in situ durability data in the literature, but with many promising alkali stable AEMs being reported, this situation is likely to soon change.

In summary, a new high-density polyethylene-based radiation-grafted anion-exchange membrane (HDPE-AEM) was developed. The switch from using low density polyethylene (LDPE) to using HDPE as a precursor film directly led to enhanced performance characteristics when the AEM was tested in a single-cell anion-exchange membrane fuel cell (AEMFC). The improved performance is hypothesized to be due to enhanced water transport characteristics, particularly the rapid water transport from the anode to the cathode, caused by the change in the nanomorphology/microstructure of the precursor films used to fabricate the AEMs. This phenomenon will be the focus of detailed future studies.

**Conflicts of interest**

There are no conflicts to declare.
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Footnotes
† Note an important initial observation that we have made. RG-AEM-based MEAs can currently be fabricated to produce high power densities or tailored for high in situ durabilities. It is evident that new MEA architectures will be required to produce a desirable sweet-spot that yield an optimal, acceptable balance of power output performance and in situ durability (such a lengthy materials engineering effort is beyond the scope of this initial materials development study).

References
1 (a) A. G. Wright, J. Fan, B. Britton, T. Weissbach, H.-F. Lee, E. A. Kitching, T. J. Peckham and S. Holdcroft, Energy Environ. Sci., 2018, 9, 2130; (b) Y. Yan, B. Xu, J. Wang and Y. Zhao, U.S. Patent Application 62/314,008 (2016); (c) J. S. Olsen, T. H. Pham and P. Jannasch, Adv. Funct. Mater., 2018, 28, 1702758; (d) H. Peng, Q. Li, M. Hu, L. Xiao, J. Lu and L. Zhuang, J. Power Sources, 2018, 390, 165; (e) S. Gottsfeld, D. R. Dekel, M. Page, C. Bae, Y. Yan, P. Zelenay and Y. S. Kim, J. Power Sources, 2018, 375, 170.
2 L. Wang, J. J. Brink, Y. Liu, A. M. Herring, J. Ponce-Gonzalez, D. K. Whelligan and J. R. Varcoe, Energy Environ. Sci., 2017, 10, 2154.
3 (a) X. Peng, T. J. Omasta, E. Magliocca, L. Wang, J. R. Varcoe, and W. E. Mustain, Angew. Chem. Int. Ed., 2019, 48, 1046; (b) X. Peng, V. Kashyap, B. Ng, S. Kurungot, L. Wang, J. R. Varcoe and W. E. Mustain, Catalysis, 2019, 9, 264.
4 L. Wang, M. Bellini, H. A. Miller, and J. R. Varcoe, J. Mater. Chem. A, 2018, 6, 15404.
5 (a) M. Mamliouk, J. A. Horsfall, C. Williams and K. Scott, Int. J. Hydrogen Energy, 2012, 37, 11912; (b) T. A. Sherazi, J. Y. Sohn, Y. M. Lee and M. D. Guiver, J. Membr. Sci., 2013, 441, 148; (c) R. Espiritu, B. T. Golden, K. Scott and M. Mamliouk, J. Mater. Chem. A, 2017, 5, 1248.
6 (a) V. Sroll, T. J. Schmidt and L. Gubler, Polym. Int., 2016, 65, 174; (b) R. Espiritu, B. T. Golden, K. Scott and M. Mamliouk, J. Power Sources, 2018, 375, 373; (c) F. Ranogajec, Radiat. Chem. Phys., 2007, 76, 1381.
7 L. Wang, E. Magliocca, E. L. Cunningham, W. E. Mustain, S. D. Poynton, R. Escudero-Cid, M. M. Nasef, J. Ponce-Gonzalez, R. Bance-Souahli, R. C. T. Slade, D. K. Whelligan, and J. R. Varcoe, Green Chem., 2017, 19, 831.
8 N. Ziv and D. R. Dekel, Electrochem. Commun., 2018, 88, 109.
9 Y. Zheng, U. Ash, R. P. Pandey, A. G. Ozioko, J. Ponce-González, M. Handl, T. Weissbach, J. R. Varcoe, S. Holdcroft, M. W. Liberatore, R. Hiesgen, and D. R. Dekel, Macromolecules, 2018, 51, 3264.
10 T. J. Omasta, L. Wang, J. M. LaManna, Y. Zhang, X. Peng, L. Wang, D. L. Jacobson, J. R. Varcoe, D. S. Hussey, B. S. Pivovar, and W. E. Mustain, Energy Environ. Sci., 2018, 11, 551.
11 T. J. Omasta, L. Wang, X. Peng, C. A. Lewis, J. R. Varcoe, and W. E. Mustain, J. Power Sources, 2018, 375, 205.

Graphical TOC entry:
Radiation-grafted HDPE-based anion-exchange membranes perform better than LDPE-based benchmarks despite exhibiting similar in situ properties.