

The reaction between Nafion sulfonyl fluoride precursor membrane and 1,4-dimethylpiperazine does not yield reliable anion-exchange membranest

Cite this: *J. Mater. Chem. A*, 2013, **1**, 1018

Received 2nd November 2012
Accepted 23rd November 2012

DOI: 10.1039/c2ta00955b

www.rsc.org/MaterialsA

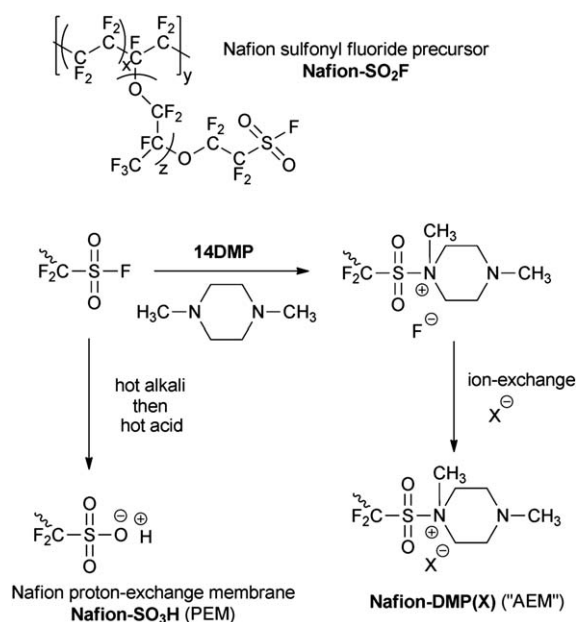
Recent reports that the reaction between Nafion sulfonyl fluoride precursor and the cyclic diamine 1,4-dimethylpiperazine yields stable anion-exchange membranes appear to be premature. On aqueous work up, membranes with high cation-exchange capacities and zero anion-exchange capacities are produced.

There is currently a significant level of research involving the development of alkaline polymer electrolyte fuel cells (APEFC) containing alkaline anion-exchange membranes (AAEM).¹ Prior AAEM studies involve either the investigation of different polymer backbones² or different functional anion-exchange head-groups.³ Studies into soluble alkaline ionomers for membrane electrode assembly fabrication are also intensifying.⁴

There have been two recent reports of the development of anion-exchange membranes (AEM) based on Nafion perfluorinated membranes.^{5,6} Both of these prior studies involve the reaction of Nafion® sulfonyl fluoride ($-\text{SO}_2\text{F}$) precursor membrane (Nafion- SO_2F) with the diamine 1,4-dimethylpiperazine (14DMP) (Scheme 1). The first communication on this system was in 2011 by Ramani *et al.*⁵ where Nafion®-111 sulfonyl fluoride membrane (Dupont, *ca.* 30 μm thickness) was reacted with 14DMP at 60 °C for up to 24 h with a subsequent OH^- anion-exchange process; they compared the Nafion-based AEMs [Nafion-DMP(OH^-)] to polysulfone AEMs. A more recent 2012 report was published by Elabd *et al.*⁶ who reacted Nafion- SO_2F membranes (111P, Ion Power, 25 μm thickness) with 14DMP at room temperature for 3 h followed by a subsequent OH^- anion-exchange process. Both studies used infrared spectroscopic, water uptake and conductivity measurements to characterise the membranes; measurements such as ion-exchange capacity determinations (IEC), thermogravimetry, small angle X-ray scattering and fuel cell testing were also conducted. Note: Elabd *et al.* have also

published a follow up paper on Nafion-based AEMs with different head-group chemistries.⁷

The Nafion-DMP(OH^-) produced by Ramani *et al.*⁵ yielded ion conductivities of over 45 mS cm^{-1} (40 mS cm^{-1} for Nafion-DMP(CO_3^{2-})) at 70 °C, but they only referred to the theoretical cation-exchange capacity (CEC) of Nafion- SO_3H (0.91 meq. g^{-1} = 1100 equivalent weight) and they did not determine the IEC directly. However, some of the data presented in these prior studies did not correlate well with the proposed synthesis of the AEM. For example, the Nafion-DMP(OH^-) produced by Elabd *et al.*⁶ yielded conductivities up to 5 mS cm^{-1} at 50 °C. Unexpectedly, the conductivities at 50 °C for Nafion-DMP(HCO_3^-),



Scheme 1 The proposed reaction of Nafion® sulfonyl precursor [Nafion- SO_2F] with 1,4-dimethylpiperazine [14DMP] to form an AEM in X^- anion form [Nafion-DMP(X)]. Also shown is the process to generate the standard PEM form of Nafion® [Nafion- SO_3H]. X^- = anion being studied.

Department of Chemistry, Faculty of Engineering and Physical Sciences, University of Surrey, Guildford GU2 7XH, UK. E-mail: j.varcoe@surrey.ac.uk; Tel: +44 (0)1483 686838

† Electronic supplementary information (ESI) available: Additional details on the syntheses and titrations used and additional spectroscopic results (Raman, FTIR and solid state NMR). See DOI: 10.1039/c2ta00955b



Nafion-DMP(CO₃²⁻), and **Nafion-DMP(Cl⁻)** were higher at 7, 7, and 12 mS cm⁻¹ respectively. Could this be evidence of AEM degradation at high pHs in these prior studies? Additionally, the prior determined IECs [measured using the Warder back-titration technique, which yields the anion-exchange capacities (AEC)] were ≤ 0.1 meq. g⁻¹; these values are <13% of the theoretical AEC value of 0.81 meq. g⁻¹ (calculated from the mono-reaction of **14DMP** with **Nafion-SO₂F** as in Scheme 1). These very low AEC values then translated to the unexpectedly high reported λ values (mol H₂O molecules per mol anion X⁻).

The results reported in this communication detail experiments that investigate this intriguing system further. To complement the prior studies, we focused on characterising the membranes using vibrational spectroscopy, solid state NMR, and IEC methods.

All **Nafion-DMP(F⁻)** syntheses followed the exemplar procedure detailed in the ESI† but with variations in **Nafion-SO₂F** + **14DMP** reaction times (3–503 h) and temperatures (room temperature, 60 °C and 80 °C). Work up involved removal of excess **14DMP**, washing with water, and then storage in water until required. Despite washing with water, a continuous smell of **14DMP** could still be detected from the vessels containing the reacted transparent brown membranes in their storage water over long periods of time (even with multiple replacement of the storage water). The membranes' colour also faded on prolonged exposure to water.

Ion-exchange of the as-synthesised **Nafion-DMP(F⁻)** membranes, into to the various other anion forms, were conducted by immersion of the membranes in excess aqueous solutions (at least 1 mol dm⁻³ concentrations) for at least 1 day followed by soaking over at least 1 day in 18.2 MΩ cm water. Despite this, the continuous presence of traces of F⁻ anions (even after multiple changes of water) was observed using ion-chromatography of the post-exchange water washings; this tentative observation may suggest either the slow decomposition of the **Nafion-DMP(X)** AEM or the slow reaction of residual -SO₂F groups in the aqueous environment (potentially slightly alkaline considering the presence of amine groups).

Before the Raman spectra of select **Nafion-DMP(X)** AEMs can be investigated, the Raman spectrum of the starting material (and the potential cation-exchange byproduct) needs to be studied. Fig. S1 in the ESI† gives the FT-Raman spectra of **Nafion-SO₂F** and the commercially available Nafion®-117 proton-exchange membrane (**Nafion-SO₃H** in Scheme 1, EW = 1100 g eq.⁻¹, 0.007 inch in thickness, Alfa Aesar UK). The FT-Raman spectrum of **Nafion-SO₂F** has bands at 1470, 1250, and 648 cm⁻¹, that are well demarcated and not present in the spectrum of **Nafion-SO₃H**; these are therefore diagnostic of the -SO₂F functional group. A band at 1470 cm⁻¹ in the FTIR spectrum of **Nafion-SO₂F** was tentatively assigned to the S-F bond motion in a recent prior study⁸ and to the asymmetric -SO₂- stretch (of the -SO₂F group) in an earlier prior study.⁹ The Raman spectrum of **Nafion-SO₃H** is well assigned (Table 2 in ref. 10). For example, the medium intensity band at 1060 cm⁻¹ is due to the symmetrical -SO₃⁻ stretch, while the medium intensity band at 973 cm⁻¹ is due to the symmetrical C-O-C stretch.

Fig. 1 below shows how the Raman spectra develop starting with **Nafion-SO₂F** and then **Nafion-DMP(F⁻)** membranes synthesised using increasing room temperature reaction times. First consider the disappearance of the -SO₂F related band at 1250 cm⁻¹ and note

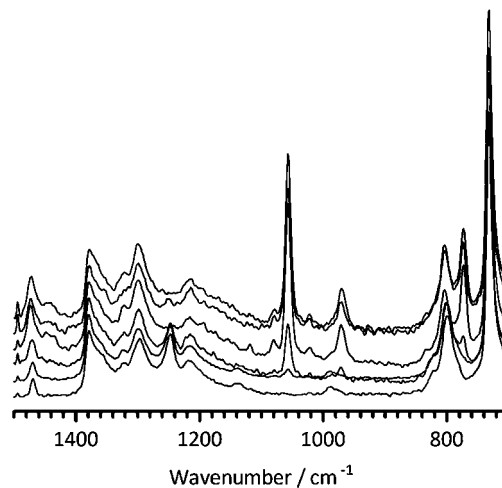


Fig. 1 The FT-Raman spectra (diagnostic region 1500–700 cm⁻¹) of the **Nafion-DMP(F⁻)** membranes formed on the reaction of **Nafion-SO₂F** and **14DMP** at room temperature for (from bottom to top): 0 h (*i.e.* pristine **Nafion-SO₂F**), 3 h, 24 h, 100 h, 144 h, and 196 h. Spectra were normalised to the band at 1380 cm⁻¹ and then stacked for clarity of presentation.

that Elabd *et al.* used 3 h reaction time with 25 μm thick **Nafion-SO₂F** precursor membranes to synthesise their membranes. With the [thicker] 50 μm **Nafion-SO₂F** precursor membranes used in this study, 3 h is not adequate for reaction at room temperature as this band remains visible until 100+ h of reaction time. This is backed up by the gradual appearance of bands at 1057, 972 and 774 cm⁻¹, all of which plateau in intensity after *ca.* 100 h of room temperature reaction time; the SO₂F-related band at 1470 cm⁻¹ in **Nafion-SO₂F** also appears to increase in intensity on increasing reaction time. A C-H stretch band at 2980 cm⁻¹ (not shown) also develops and plateaus in intensity around 100 h reaction time (the most intense C-H stretch for in the spectrum of **14DMP** is located at 2950 cm⁻¹ – see ESI†). Solid state NMR Data (see ESI†) yields further evidence of reaction of the -SO₂F functional group of the **Nafion-SO₂F** precursor (this time with a **Nafion-DMP(F⁻)** synthesised at 80 °C and 240 h).

The Raman spectra of the different **Nafion-DMP(F⁻)** membranes are complex. The bands at 1057 and 972 cm⁻¹ that develop on reaction are also present in the spectrum of **Nafion-SO₃H**, but the new band at 774 cm⁻¹ is absent from both the spectra of **Nafion-SO₃H** and **Nafion-SO₂F**; however, there is a strong band at 779 cm⁻¹ in the spectrum of pristine **14DMP** [there is also a medium intensity band at 1470 cm⁻¹ in the Raman spectrum of **14DMP**]. The bands located in the range 1051–1059 cm⁻¹ in the FTIR spectrum of the **Nafion-DMP(F⁻)** and **Nafion-DMP(OH⁻)** produced in the prior study by Elabd *et al.*⁶ were assigned to the -SO₂-N⁺R₃ group. However, considering the spectroscopic data discussed above, this assignment is risky.

An alternative hypothesis is that there is a reaction [or partial reaction] between the **14DMP** and **Nafion-SO₂F** but the product membranes [or residual -SO₂F groups in the membranes] react with the water [in the aqueous work up and in the presence of the alkaline piperazine-related groups] and form the proton-exchange form **Nafion-SO₃H** membranes. The infrared spectroscopic data



presented in the ESI† provides evidence of degradation of a **Nafion-DMP(F⁻)** membrane when exposed to alkaline conditions.

Fig. 2 shows an AEC curve (Cl⁻ method)¹¹ of an example **Nafion-DMP(Cl⁻)** AEM. It is immediately obvious that the Cl⁻ AEC cannot be determined as the release of Cl⁻ is below the detection limit of the technique (and well below expected from the theoretical AEC of 0.81 meq. g⁻¹). This was consistently the case for **Nafion-DMP(Cl⁻)** membranes synthesised with different reaction times and temperatures [**Nafion-DMP(Cl⁻)** membranes made from reactions at room temperature for 144–503 h, at 80 °C for 240 h, and 60 °C for 25 days were tested for AEC]. Zero ion-exchange capacities were also measured when the pre-titration KCl exchange solution were replaced with aqueous HCl (1 mol dm⁻³) [theoretically this modified titration method would enable the Cl⁻ titrations to probe for the presence of non-ionic tertiary amine groups in addition to the ionic quaternary ammonium groups].

AEMs containing the -SO₂-N(R')-R-N⁺R₃'' functional group chemistry (where the N connected to the -SO₂- group is not charged) are known,¹² but there is little precedence for the functional group -SO₂-N⁺R₃ (1-methyl-1-[(4-methylphenyl)sulfonyl]pyrrolidinium is a rare example on ChemSpider¹³). The question really must be – How stable is the S–N bond in the proposed -SO₂-N⁺ functional group in Scheme 1? To test this, the **Nafion-DMP(F⁻)** AEM [synthesised at room temperature and 144 h] was tested for CEC (see ESI† for the standard method used): CEC = 0.79 ± 0.08 meq. g⁻¹ (sd, n = 3). As a control experiment, **Nafion-SO₂F** was hydrolyzed in a solution of 15% mass KOH, 35% mass DMSO, and 50% mass de-ionized water at 80 °C for several hours, followed by thorough washing in water and final conversion to the H⁺ form **Nafion-SO₃H** membrane by immersion in nitric acid; the CEC of this control membrane = 0.92 ± 0.05 meq. g⁻¹ (sd, n = 3), which compares well to the theoretical 0.91 meq. g⁻¹ for **Nafion-SO₃H**

membranes produced from Nafion® PFSA R-1100 sulfonyl fluoride precursor. This is very strong evidence that the end result of the synthesis procedure reported is membranes that are predominantly in the *cation-exchange form*. These membranes will therefore not be suitable for application in electrochemical devices requiring *anion-exchange* polymer electrolytes (*i.e.* APEFCs and alkaline electrolysers).

The authors thank the UK's Engineering & Physical Sciences Research Council (EPSRC grant EP/H025340/1 and Dr John Varcoe's EPSRC Leadership Fellowship funded with grant EP/I004882/1), the University of Surrey (for funds for Susan Stephen's final year undergraduate project) and the EPSRC Solid State NMR service at the University of Durham.

References

- 1 E. H. Yu, X. Wang, U. Krewer, L. Li and K. Scott, *Energy Environ. Sci.*, 2012, **5**, 5668; J. Pan, C. Chen, L. Zhuang and J. Lu, *Acc. Chem. Res.*, 2012, **45**, 473; G. Merle, M. Wessling and K. Nijmeijer, *J. Membr. Sci.*, 2011, **377**, 1; R. Zeng and J. R. Varcoe, *Recent Pat. Chem. Eng.*, 2011, **4**, 93; G. Couture, A. Alaeddine, F. Boschet and B. Ameduri, *Prog. Polym. Sci.*, 2011, **36**, 1521; B. Pivovar, *Alkaline Membrane Fuel Cell Workshop Final Report NREL/BK-5600-54297*, National Renewable Energy Laboratory, Golden CO, USA, 2011, <http://www.osti.gov/bridge>; E. Antolini and E. R. Gonzalez, *J. Power Sources*, 2010, **195**, 3431; S. Lu, J. Pan, A. Huang, L. Zhuang and J. Lu, *Proc. Natl. Acad. Sci. U. S. A.*, 2008, **105**, 20611; L. A. Adams, S. D. Poynton, C. Tamain, R. C. T. Slade and J. R. Varcoe, *ChemSusChem*, 2008, **1**, 79.
- 2 N. J. Robertson, H. A. Kostalik IV, T. J. Clark, P. F. Mutolo, H. D. Abruña and G. W. Coates, *J. Am. Chem. Soc.*, 2010, **132**, 3400; M. R. Hibbs, C. H. Fujimoto and C. J. Cornelius, *Macromolecules*, 2009, **42**, 8316; J. R. Varcoe, R. C. T. Slade, E. Lam How Yee, S. D. Poynton, D. J. Driscoll and D. C. Apperley, *Chem. Mater.*, 2007, **19**, 2686.
- 3 O. D. Thomas, K. J. W. Soo, T. J. Peckham, M. P. Kulkarni and S. Holdcroft, *J. Am. Chem. Soc.*, 2012, **134**, 10753; Y. Zha, M. L. Disabb-Miller, Z. D. Johnson, M. A. Hickner and G. N. Tew, *J. Am. Chem. Soc.*, 2012, **134**, 4493; S. Gu, J. Skovgard and Y. S. Yan, *ChemSusChem*, 2012, **5**, 843; X. Lin, L. Wu, Y. Liu, A. L. Ong, S. D. Poynton, J. R. Varcoe and T. Xu, *J. Power Sources*, 2012, **217**, 373; X. Wang, M. Li, B. T. Golding, M. Sadeghi, Y. Cao, E. H. Yu and K. Scott, *Int. J. Hydrogen Energy*, 2011, **36**, 1002; Y. Ye and Y. A. Elabd, *Macromolecules*, 2011, **44**, 8494; S. Gu, R. Cai and Y. Yan, *Chem. Commun.*, 2011, **47**, 2856; Q. Zhang, S. Li and S. Zhang, *Chem. Commun.*, 2010, **46**, 7495; X. Kong, K. Wadhwa, J. G. Verkade and K. Schmidt-Rohr, *Macromolecules*, 2009, **42**, 1659; R. Schwesinger, R. Link, P. Wenzl, S. Kossek and M. Keller, *Chem.–Eur. J.*, 2006, **12**, 429; M. Tomoi, K. Yamaguchi, R. Ando, Y. Kantake, Y. Aosaki and H. Kubota, *J. Appl. Polym. Sci.*, 1997, **64**, 1161; K. J. T. Noonan, K. M. Hugar, H. A. Kostalik IV, E. B. Lobkovsky, H. D. Abruña and G. W. Coates, *J. Am. Chem. Soc.*, 2012, **134**, 18161.

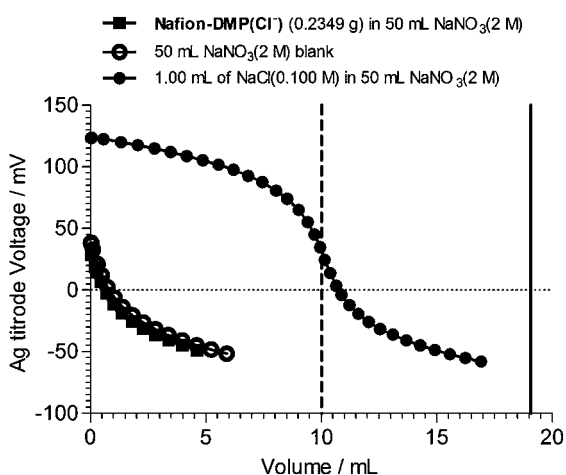


Fig. 2 The anion-exchange capacity (AEC) titration curves of a **Nafion-DMP(Cl⁻)** AEM synthesised from **Nafion-SO₂F** and **14DMP** at room temperature and 503 h. The method used determines the amount of Cl⁻ anions released from the AEM on immersion in excess NaNO₃ (2 mol dm⁻³). The solid vertical line indicates the expected end point for 0.2349 g of AEM with the theoretical AEC of 0.81 meq. g⁻¹. The dashed vertical line gives the theoretical endpoint for an aqueous solution consisting of 1.00 cm³ of NaCl (0.1000 mol dm⁻³) added to 50 cm³ NaNO₃ (2.0 mol dm⁻³) which matches the actual control experiment conducted.



- 4 R. Zeng, J. Handsel, S. D. Poynton, A. J. Roberts, R. C. T. Slade, H. Herman, D. C. Apperley and J. R. Varcoe, *Energy Environ. Sci.*, 2011, **4**, 4925; M. Tanaka, M. Koike, K. Miyatake and M. Watanabe, *Polym. Chem.*, 2011, **2**, 99; W. Li, J. Fang, M. Lv, C. Chen, X. Chi, Y. Yang and Y. Zhang, *J. Mater. Chem.*, 2011, **21**, 11340.
- 5 M.-S. J. Jung, C. G. Arges and V. Ramani, *J. Mater. Chem.*, 2011, **21**, 6158.
- 6 H. L. S. Salerno, F. L. Beyer and Y. A. Elabd, *J. Polym. Sci., Part B: Polym. Phys.*, 2012, **50**, 552.
- 7 H. L. S. Salerno and Y. A. Elabd, *J. Appl. Polym. Sci.*, 2013, **127**, 298.
- 8 S. A. Perusich, *J. Appl. Polym. Sci.*, 2011, **120**, 165.
- 9 A. J. Greso, R. B. Moore, K. M. Cable, W. L. Jarrett and K. A. Mauritz, *Polymer*, 1997, **38**, 1345.
- 10 A. Gruger, A. Régis, T. Schmatko and P. Colomban, *Vib. Spectrosc.*, 2001, **26**, 215.
- 11 O. I. Deavin, S. Murphy, A. Ong, S. D. Poynton, R. Zeng, H. Herman and J. R. Varcoe, *Energy Environ. Sci.*, 2012, **5**, 8584.
- 12 R. Ei Moussaoui and R. Martin, *Eur. Pat.* EP1612874, 2004; R. Ei Moussaoui and R. Martin, *PCT Pat.* WO2006003182, 2006.
- 13 <http://www.chemspider.com/Chemical-Structure.3526107.html>, (accessed 15:39, 14 September 2012) [CSID: 3526107].

