Model Long Side-Chain PPO-Based Anion Exchange Ionomers: Properties and Alkaline Stability

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Abstract

The stability of anion exchange membranes is paramount for the use in alkaline fuel cells. Long chain ionomers are supposed to be more alkaline-resistant with respect to short chain isomers. In this paper the synthesis, properties and stability of ionomers with a long side chain are investigated. Poly(2,6-dimethyl-1,4-phenylene)oxide (PPO) is chosen as backbone, due to its reported stability in alkaline conditions. The functional group is pentyl-ammonium with trimethylamine (TMA) or 1,4-diazabicyclo[2.2.2]octane (DABCO) as model amines. The synthesis is carried out via metalation reaction and is optimized as a function of temperature and time. The water uptake is relatively low, in accordance with the large hydrophobicity of the PPO backbone. The through-plane ionic conductivity is consistent with literature data; it amounts to 15.3 mS/cm at 80 °C for the TMA derivative. The mechanical properties are typical of ionomers below the glass transition temperature (for the TMA derivative at ambient humidity: Young Modulus = 1310 ± 30 MPa). The stability in alkaline conditions, studied by thermogravimetric analysis and measurements of ionic conductivity and ion exchange capacity, is higher than that of short-side chain ionomers with the same basic group. The decrease of ionic conductivity (57 vs 22% residual conductivity after 72 h in 2 M NaOH at 80 °C) and IEC is monitored showing that the degradation is fast in the first hours and may be described by second order kinetics. These results help in selecting high performance anion exchange membranes for electrochemical energy technologies.

Introduction

Anion exchange ionomers are used as innovative separators in alkaline membrane fuel cells¹⁻³. The major advantage of these devices is that the oxygen reduction reaction (ORR) can be catalysed in alkaline conditions by non-noble metals, such as Ag or Ni³⁻⁵. The use of anion exchange membranes (AEM) avoids the severe carbonation issues observed in alkaline fuel cells with liquid electrolytes. However, the practical application of AEM in fuel cells is hampered by their instability in alkaline environment due to the attack of the ammonium groups by the hydroxide ions⁶⁻⁹. Furthermore, the polymer backbone can also be attacked and chain scission processes occur¹⁰⁻¹¹. The instability of the widely studied poly(arylene ether sulfones) (PSU) is promoted by the sulfone groups present in the backbone through their inductive effect increasing the reactivity in specific positions¹³. AEM based on poly(2,6-dimethyl-1,4-phenylene oxide) (PPO)¹¹⁻¹² are expected to have a higher alkaline stability in comparison with PSU, due to the less activated backbone for a nucleophilic attack¹².

The alkaline attack may occur by several mechanisms^{8-9, 12}. The most difficult to avoid is the nucleophilic substitution reaction S_N2: the rate of this bimolecular reaction is proportional to the IEC of the ionomer and the OH⁻ concentration in the medium¹³⁻¹⁴. The bimolecular Hofmann elimination E2 leads to the formation of an alkene. In sterically very unfavourable conditions, the monomolecular elimination reaction E1 can also occur. Various approaches are discussed in the literature to mitigate the alkaline degradation^{1-2, 15-25}, including a change of the polymer backbone^{24, 26-28}, the introduction of side chains^{18, 22, 29-32}, the presence of a secondary phase³³⁻³⁵, a delocalization of the positive charge^{7, 21}, or a protection by steric hindrance^{7, 36-37}.

In this study, we investigate the alkaline stability of model long side-chain anion-conducting ionomers based on PPO with N-pentyl-N,N,N-trimethylammonium side groups, given that the theoretical work of Pivovar and coworkers shows that the most stable side-chain length corresponds to 4-5 CH₂ groups³⁰. Two model amines, small and flexible trimethylamine (TMA) and bulky and rigid 1,4-diaza-bicyclo[2.2.2]octane (DABCO), are used for functionalization. The water uptake, thermal, mechanical and electrical properties are reported before and after a harsh alkaline treatment in 2 M NaOH at 80 °C during various times.

Experimental section

Materials

Poly(2,6-dimethyl-1,4-phenylene)oxide (PPO, MW = 50000 g/mol) trimethylamine (TMA, 4.2 M

in ethanol), 1,4-diazabicyclo[2.2.2] octane (DABCO), n-butyllithium (BuLi, 2.5 M in hexane), tetrahydrofuran (THF, anhydrous), 1,4-dibromobutane, and N-methyl-2-pyrrolidone (NMP) were reagent-grade, purchased from Sigma-Aldrich.

Ionomer synthesis

Synthesis of PPO-(CH₂)₅Br.

PPO (5 g, 41.7 meq) was dissolved in dry THF (500 mL) under nitrogen at 55 °C. The solution was cooled to RT and 25 mL of n-BuLi (2.5 M, 62.6 meq) was added dropwise until the appearance of a persistent light yellow coloration. The reaction was carried out during 3 h at two temperatures: 25 or 55 °C in order to optimize the final products (see Results and Discussion). The solution was then cooled to -70 °C and 1,4-dibromobutane (15 mL, 125 meq) was added. A light yellow solution was immediately formed and the reaction was kept under stirring overnight at RT. The product was precipitated in ethanol and the yellow polymer was filtered, washed repeatedly with fresh ethanol and dried at 55 °C during 2 days.

The products were characterized by ¹H NMR. The degree of benzyl bromination was 0.24 at 25 °C and 0.35 at 55 °C.

Synthesis of PPO-(CH₂)₅TMA and PPO-(CH₂)₅DABCO.

The brominated polymer obtained at 55 °C (degree of benzyl bromination 0.35) was dissolved in NMP and the amine was added in excess of 5.5 (TMA) or 1.5 (DABCO) with respect to the degree of functionalization, due to the well-known volatility of TMA. The solution was kept at 80 °C for 72 h and then the polymer was precipitated in deionized water, filtered, washed with water and dried under vacuum at 60 °C.

Membranes were obtained by casting at 95 °C from NMP solution (0.3 g polymer in 25 mL solvent).

NMR spectroscopy

¹H NMR spectroscopy was performed using a Bruker Avance 400 apparatus (400.13 MHz) using deuterated solvents (CDCl₃, DMSO-d₆). Chemical shifts were referenced to tetramethylsilane.

Ion exchange capacity

The IEC was determined by NMR spectroscopy (see Results) and by Mohr titration. For the Mohr titration, the chloride form was obtained by treating the AEM in 1 M NaCl solution during

24 h. The membranes were then washed carefully with deionized water to remove any excess NaCl. The Cl⁻ anions contained in the membranes were exchanged with SO₄²⁻ by immersion in 1 M Na₂SO₄ solution for 24 h. This solution was titrated using 0.02 M AgNO₃. The end-point indicator was prepared dissolving 1 g of K₂CrO₄ in 20 mL of water.

By Mohr titration, PPO-(CH₂)₅-TMA had an IEC = 1.8 meq/g and PPO-(CH₂)₅-DABCO an IEC = 1.3 meq/g.

Water uptake

The membrane mass was measured after 24 h immersion in deionized water at 25 °C, removing excess surface water by rapid blotting with absorbent paper (m_{wet}). The dry mass was obtained after drying in a desiccator over P₂O₅ during 72 h (m_{dry}). The water uptake WU was calculated according to the equation:

$$WU = 100 \cdot \frac{m_{wet} - m_{dry}}{m_{dry}} \quad (1)$$

The hydration numbers λ , which are the number of water molecules per ammonium group, are calculated using the IEC of the ionomer according to the equation:

$$\lambda = \frac{10 \cdot WU}{M(H_2O) \cdot IEC} \quad (2)$$

M(H₂O) is the molar mass of water.

Ionic conductivity

Prior to the measurements, the membrane samples were immersed during 24 h in 2 M NaCl (Cl form) or 2 M NaOH (OH form) solution and then thoroughly washed in deionized water. In the latter case, the solution was kept under nitrogen to avoid contamination by CO₂. The ionic conductivity was measured between 25 and 80 °C by impedance spectroscopy using a Biologic VSP300 equipment. The frequency range was between 1 Hz and 6 MHz and the a.c. voltage amplitude was 20 mV. The measurements were made on fully humidified samples in two-point through-plane configuration using a swagelok cell with electrode area $A = 0.264 \text{ cm}^2$. The membrane resistance R was determined at the intersection of the electrode arc with the real axis. The membrane thickness d was determined before and after the measurements using a Mitutuyo micrometer.

The ionic conductivity was calculated according to the equation:

$$\sigma = \frac{d}{R \cdot A} \quad (3)$$

Mechanical measurements

The tensile stress-strain measurements were performed on membranes in Cl form using an ADAMEL Lhomargy DY30 apparatus at room temperature and ambient humidity ((50 +/- 10) % RH) or at full humidification.

Thermogravimetry

High-resolution thermogravimetric analysis of membranes (Cl form) was made using a TA Q500 apparatus. The samples were placed in Pt sample holders. The experiments were performed in air flux between 50 and 600 °C. The maximum heating rate was 3 K/min.

Results and Discussion

Synthesis

Figure 1a shows the molecular formulas of long side-chain ionomers functionalized by TMA and DABCO. Butyllithium may attack PPO protons on two different positions in the side chain or in the main chain leading to the formation of a benzyl (A) or an aryl (B) carbanion (Figure 1b). The following reaction with 1,4-dibromobutane gives the bromine derivatives in benzylic or aryl position. The distribution of the products depends on the temperature and time. In our case, the time was fixed at 3 h while the temperature was varied (25 or 55 °C). At low temperature the ratio A:B (Figure 1b) was 2.4:1, while at higher temperature the ratio was 8.8:1. These data correspond to a competitive reaction under thermodynamic or kinetic control³⁸. The substitution at benzylic position appears to be more favored than that at the aromatic ring position under both kinetic and thermodynamic control, although a larger selectivity is observed under thermodynamic control. The larger stability of the benzyl anion can be ascribed to resonance delocalization of the negative charge that is not allowed for the sp² centered aryl anion.

Figure 1. a) Molecular formulas of long side-chain anion exchange polymers (substitution in benzylic position); b) reaction pathways for the synthesis of brominated PPO.

The ^1H NMR spectra of the brominated derivatives obtained at the two temperatures are shown in Figure 2.

Figure 2. ^1H NMR spectra of PPO-(CH₂)₅Br in CDCl₃ obtained at a) 25 °C and b) 55 °C.

The presence of the substitution of alkylbromine in aryl position is evidenced by the signal of the hydrogen *b* (1H, 6.03 ppm), which appears with higher intensity in the reaction carried out at 25

°C (Figure 2a). The signals of the methylene group in α to bromine (2H, *b* and *b'*) appear at 3.34 ppm for the benzyl derivative and at 3.48 ppm for the aryl one ³⁹. The latter signal is not observable in the spectrum of the compound obtained at higher temperature (Figure 2b) due to the lower quantity of the aryl substitution. The integration of the signal *f* (2H, 1.40 ppm), present only in the derivative obtained from the attack in benzylic position (about 90% of the peaks *b* and *b'*, Figure 2b), confirms that a part of the product is due to the formation of the aryl (B) carbanion (Figure 1b). The total degree of bromination DBr is calculated from the integration of the protons *b* and *b'* with respect to the protons *a* of PPO; DBr amounts to 0.34 and 0.39 at 25 and 55 °C respectively.

The reaction with tertiary amines was conducted only with the derivative obtained at higher temperature.

Figure 3a shows the ¹H NMR spectrum of PPO-(CH₂)₅-TMA. The peak *a* is assigned to the hydrogens (2H) of the aromatic ring containing the pentyl chain originating from the attack of BuLi in benzylic position. The attack on the aryl moiety generates the signal *b* (approximately 4%). The peak *b* is not integrable in the case of PPO-(CH₂)₅-DABCO (Figure 3b). The degree of amination (DAM) of PPO-(CH₂)₅-TMA is obtained from the integration of the aromatic hydrogens of PPO (*a*, 2H), the hydrogens of methyl groups on the ammonium ions (*i*, 9H), and the internal hydrogens of the pentyl chain (*e, f, g* 6H) resulting in 0.28. The other signals overlap with the solvents (DMSO, H₂O). The spectrum of PPO-(CH₂)₅-DABCO is more complex. The peak *l* due to hydrogens in α to the nitrogen are shifted to highfield with respect to hydrogens near to the quaternary nitrogen (*i*) and overlap with the signals *b* of the CH₂ belonging to the pentyl chain. The integration of NMR peaks *e, f, g* of the DABCO derivative shows the absence of crosslinks in spite of the two amine groups in the molecule ²⁴.

Figure 3. ^1H NMR spectra of (a) PPO- $(\text{CH}_2)_5$ -TMA with DAM = 0.28 and (b) PPO- $(\text{CH}_2)_5$ -DABCO with DAM = 0.20 in DMSO d_6 .

The DAM is lower for DABCO showing a lower reaction yield probably due to the more bulky molecule that reduces the $\text{S}_{\text{N}}2$ reaction rate by steric hindrance. The IEC obtained from NMR using eq. 4 is 1.8 meq/g for TMA and 1.3 meq/g for DABCO in good agreement with the values obtained by Mohr titration (see experimental).

$$\text{DAM} = \frac{\text{IEC} \times M_{\text{PPO}}}{1 - \text{IEC} \times M_{(\text{CH}_2)_4\text{TMA or } (\text{CH}_2)_4\text{DABCO}}} \quad (4)$$

DAM, $M_{PPO} = 120$ g/mol and $M_{(CH_2)_4TMA \text{ or } (CH_2)_4DABCO} = 115$ or 168 g/mol are respectively the degree of amination, the molar mass of the polymer backbone and the molar mass of the functional unit, which contains 4 CH_2 groups given that one CH_2 belongs to the PPO structure.

Properties

The water uptake and ion conductivity values of PPO-(CH_2)₅-TMA and PPO-(CH_2)₅-DABCO are reported in Table 1. The water uptake values are relatively low in comparison with PSU-based ionomers^{14, 40}, certainly due to the very hydrophobic PPO backbone, known to exhibit the lowest water sorption among engineering plastics⁴¹, and increase relatively little with temperature. The room temperature data are consistent with similar PPO-based ionomers⁴²⁻⁴³. The hydration numbers are comparable with literature data for cross-linked copolymers¹¹, but clearly above ionomers made by a bromination route and with very long side chains¹⁶. The higher hydration numbers for the DABCO derivative might be attributed to the larger volume of the DABCO molecule increasing the inter-chain space.

The conductivity data are comparable with recent literature⁴³⁻⁴⁴. They are also consistent with the conductivity reported at room temperature for sulfonated PPO with an IEC = 2.63 meq/g (12 mS/cm⁴¹), considering the lower mobility of hydroxide (at infinite dilution in water: $20.5 \cdot 10^{-5} \text{ cm}^2 \text{ s}^{-1} \text{ V}^{-1}$)⁴⁵⁻⁴⁶ vs. hydronium ions ($36.3 \cdot 10^{-5} \text{ cm}^2 \text{ s}^{-1} \text{ V}^{-1}$)⁴⁵⁻⁴⁶. The ion conductivity of the Cl form is below the OH form due to the lower mobility of the chloride ion (at infinite dilution in water: $7.9 \cdot 10^{-5} \text{ cm}^2 \text{ s}^{-1} \text{ V}^{-1}$)⁴⁵⁻⁴⁶. The higher values for the TMA derivative are consistent with its higher IEC. The activation energy of conductivity calculated for the OH form of PPO-(CH_2)₅-TMA and PPO-(CH_2)₅-DABCO are 0.11 and 0.13 eV, respectively, consistent with a conduction mechanism through hydrated channels¹¹. The activation energy for the Cl forms is higher (0.2 eV) in agreement with the expectation and the literature for halide ion conductors^{11, 47}.

Table 1. Ion conductivity σ , water uptake WU and hydration number λ of PPO-(CH₂)₅-TMA (IEC = 1.8 meq/g) and PPO-(CH₂)₅-DABCO (IEC = 1.3 meq/g).

PPO-(CH ₂) ₅ -TMA							PPO-(CH ₂) ₅ -DABCO					
T	$\sigma(\text{OH})/$	WU(OH)	$\lambda(\text{OH})$	$\sigma(\text{Cl})/$	WU(Cl)	$\lambda(\text{Cl})$	$\sigma(\text{OH})/$	WU(OH)	$\lambda(\text{OH})$	$\sigma(\text{Cl})/$	WU(Cl)	$\lambda(\text{Cl})$
/ °C	mS cm ⁻¹	/%		mS cm ⁻¹	/%		mS cm ⁻¹	/%		mS cm ⁻¹	/%	
25	8.4	47.8	14.7	1.9	20.2	6.2	4.3	49.8	21.3	1.4	21.0	9.0
45	8.9	53.0	16.3	3.3	25.1	7.7	5.3	-	-	2.0	-	-
60	12.9	-	-	5.5	31.0	9.6	7.8	56.1	24.0	3.1	-	-
80	15.3	58.8	18.1	9.3	36.0	11.1	9.6	62.5	26.7	4.7	36.1	15.4

Typical mechanical tensile stress-strain test curves for membranes in Cl form are shown in Figure 4. One notices a very good reproducibility, especially for PPO-(CH₂)₅-TMA. The average mechanical properties are reported for the two polymers in Table 2. They are comparable to the properties of other aromatic polymer electrolytes^{7, 35, 48-50} and are typical of stiff ionomers below their glass transition temperature. The glass transition temperature of pristine PPO is known to be very high (212 °C)⁴¹. The Young modulus is above 1 GPa at ambient humidity, related to the high stiffness and strength of pristine PPO (2.55 GPa and 63.4 MPa, respectively⁴¹) weakened by the introduction of the ionic groups that attract water molecules, as observed in all ionomers⁵¹. The high dielectric constant of water reduces the Van der Waals forces between the macromolecular chains, mainly attributed to dipole-dipole interactions of the benzene rings⁴¹, which are probed in the elastic domain. The lower Young modulus and yield stress at 100% RH is consistent with the plasticizing effect of water. The low elongation at break of all samples, irrespective of the water content, is in the order of magnitude of similar ionomers⁴⁸. The yield stress and tensile strength are much lower for the DABCO derivative, which is attributable to the large and rigid DABCO molecule, which increases the distance between macromolecular chains, enabling larger hydration numbers and weakening the Van der Waals forces between chains, as discussed above.

Figure 4. Tensile stress-strain curves of membranes (CI form) at 25 °C at RH = (50 ± 10) % and fully humidified (RH = 100 %).

Table 2. Average mechanical properties of PPO-(CH₂)₅-TMA and PPO-(CH₂)₅-DABCO: Young modulus E, yield stress YS and elongation at break ϵ .

Sample	RH / %	E / MPa	YS / MPa	ϵ / %
PPO-(CH ₂) ₅ -TMA	50	1310 ± 30	28.4 ± 0.3	3.3 ± 0.1
	100	820 ± 60	16.2 ± 3.4	3.5 ± 0.4
PPO-(CH ₂) ₅ -DABCO	100	730 ± 140	4.2 ± 0.8	2.5 ± 0.5

Alkaline stability

The thermal behavior of AEM is investigated by high-resolution thermogravimetric analysis before and after a degradation treatment during 168 h in 2 M NaOH at 80 °C (Figure 5a and b). Figure 5a shows the mass loss and Figure 5b the derivative thermograms. The peaks correspond to the principal mass loss phenomena. In pristine PPO-(CH₂)₅-TMA, the large peak around 200 °C can be attributed to the loss of TMA; the mass loss of about 12%, integrated between 160 and 230 °C, is in good agreement with the degree of functionalization of 0.28 (corresponding to 11%). This asymmetric peak is significantly reduced after the alkaline treatment. The mass loss, integrated over the same temperature range, is 8%, corresponding to a decrease of the

ammonium groups by about 40 %, In PPO-(CH₂)₅-DABCO, the loss of DABCO (red curves) is observed at a slightly higher temperature around 230 °C.

The second peak at 365 °C (starting around 300 °C) in the pristine ionomers is the trace of the decomposition of methyl and pentyl groups. Considering the degree of functionalization of the ionomers, the mass loss should be 27% for TMA and 25% for DABCO in good agreement with the experiment. The very similar broad mass loss centered around 415 °C for PPO-(CH₂)₅-TMA and PPO-(CH₂)₅-DABCO can be attributed to the decomposition of partially oxidized and cross-linked PPO, as shown previously⁵²⁻⁵³. The main-chain of PPO rearranges on heating in the degradation step, accompanied by the formation of C-C bonds, followed by the thermal degradation of the methylene bridges⁴¹. After the alkaline treatment, the PPO degradation shows instead a sharp double peak around 410 °C. Similar observations were reported previously¹³.

a)

b)

Figure 5. High-resolution thermogravimetric curves of PPO-(CH₂)₅-TMA (Cl form) before and after alkaline degradation in 2 M NaOH at 80°C during 168 h and PPO-(CH₂)₅-DABCO (Cl form). a) Mass loss curves, b) derivative curves.

Figure 6 presents the decrease of the ionic conductivity of PPO-(CH₂)₅-TMA and PPO-(CH₂)₅-DABCO with the time of the alkaline treatment. Furthermore, the decrease of the IEC of PPO-(CH₂)₅-TMA is represented. Inside the experimental uncertainty, the IEC and conductivity data are in qualitative agreement, showing a rapid degradation during the first hours and a slower kinetics at longer times. Both observations can be reconciled with the kinetics of second order reactions, such as S_N2 or E2, which are both possible given the presence of a β -hydrogen in the long side-chain AEM.

Figure 6. Ionic conductivity of PPO-(CH₂)₅-TMA and PPO-(CH₂)₅-DABCO and ion exchange capacity (IEC) of PPO-(CH₂)₅-TMA as function of the degradation time in 2 M NaOH at 80 °C.

For PPO-(CH₂)₅-DABCO, the alkaline degradation is continuous, showing that the bulky and rigid side group does not improve the alkaline stability, different from the results in short side chain ionomers¹³.

In the literature¹⁸, a low degradation was reported for a similar long side chain PPO-(CH₂)₅-TMA in 1 M NaOH; although we do not confirm this result completely, the degradation is anyway much lower than for the corresponding short side chain PPO-TMA ionomer¹³, as shown in the normalized conductivity plot represented in Figure 7. The higher degradation rate observed in this work is certainly also related to the higher NaOH concentration, which is a parameter of the rate of second order reactions.

Figure 7. Normalized conductivity dependence on the time of alkaline treatment in 2 M NaOH at 80 °C for long and short side-chain AEM ¹³.

The improvement of stability for the long-chain sample is evident in Figure 7; the spacer concept ⁵⁴ is verified. Several hypothesis explaining the better stability can be formulated: i) the S_N2 reaction is less effective, because the primary alkyl halide presents a more than two order of magnitude lower reactivity than the benzyl halide ⁵⁵; ii) the ether linkage in the backbone is less destabilized due to the long alkyl chain that screens the positive charge of the ammonium group; iii) the antiperiplanar conformation needed for the E2 reaction is less stable when a long chain is attached to ammonium ³⁰. A simple conformational analysis by Newman projections confirms this result.

Conclusions

Anion exchange membranes based on PPO with a pendant side chain functionalized with TMA or DABCO are synthesized using a metalation route. The optimization of the reaction conditions allows obtaining ionomers with the pendant groups in benzylic position, which is the thermodynamically favoured product.

The properties and alkaline stability are investigated by several techniques. Stress-strain tests are typical of stiff polymers below their glass transition temperature. The water uptake is lower than that of PSU-based ionomers, but comparable with short side chain PPO. The DABCO derivative presents a higher water uptake due to the bulky molecule. The ionic conductivity and its activation energy are consistent with the mobility of the conducting ions.

The stability in alkaline condition, studied by thermogravimetry, ion conductivity and IEC measurements, is clearly improved vs. short side chain ionomers. The alkaline degradation is rapid during the first hours of the reaction and shows a slower kinetics at longer times. Both observations can be reconciled with the kinetics of second order reactions, such as S_N2 or E2.

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References

1. Merle, G.; Wessling, M.; Nijmeijer, K., Anion exchange membranes for alkaline fuel cells: A review. *J. Membr. Sci.* **2011**, *377*, 1-35.
2. Couture, G.; Alaaeddine, A.; Boschet, F.; Ameduri, B., Polymeric materials as anion-exchange membranes for alkaline fuel cells. *Progress in Polymer Science* **2011**, *36*, 1521-1557.
3. Hickner, M. A.; Herring, A. M.; Coughlin, E. B., Anion Exchange Membranes: Current Status and Moving Forward. *Journal of Polymer Science Part B-Polymer Physics* **2013**, *51*, 1727-1735.
4. Pan, Z. F.; An, L.; Zhao, T. S.; Tang, Z. K., Advances and challenges in alkaline anion exchange membrane fuel cells. *Prog. Energy Combust. Sci.* **2018**, *66*, 141-175.
5. Arges, C. G.; Zhang, L., Anion Exchange Membranes' Evolution toward High Hydroxide Ion Conductivity and Alkaline Resiliency. *Acs Applied Energy Materials* **2018**, *1*, 2991-3012.
6. Arges, C. G.; Wang, L. H.; Parrondo, J.; Ramani, V., Best Practices for Investigating Anion Exchange Membrane Suitability for Alkaline Electrochemical Devices: Case Study Using Quaternary Ammonium Poly(2,6-dimethyl 1,4-phenylene)oxide Anion Exchange Membranes. *J. Electrochem. Soc.* **2013**, *160*, F1258-F1274.
7. Di Vona, M. L.; Narducci, R.; Pasquini, L.; Pelzer, K.; Knauth, P., Anion-conducting ionomers: Study of type of functionalizing amine and macromolecular cross-linking. *Int. J. Hydrogen Energy* **2014**, *39*, 14039-14049.
8. Sun, Z.; Pan, J.; Guo, J. N.; Yan, F., The Alkaline Stability of Anion Exchange Membrane for Fuel Cell Applications: The Effects of Alkaline Media. *Advanced Science* **2018**, *5*, 11.
9. Nunez, S. A.; Hickner, M. A., Quantitative H-1 NMR Analysis of Chemical Stabilities in Anion-Exchange Membranes. *Acs Macro Letters* **2013**, *2*, 49-52.
10. Miyanishi, S.; Yamaguchi, T., Ether cleavage-triggered degradation of benzyl alkylammonium cations for polyethersulfone anion exchange membranes. *Physical Chemistry Chemical Physics* **2016**, *18*, 12009-12023.
11. Pandey, T. P.; Sarode, H. N.; Yang, Y. T.; Yang, Y.; Vezzu, K.; Di Noto, V.; Seifert, S.; Knauss, D. M.; Liberatore, M. W.; Herring, A. M., A Highly Hydroxide Conductive, Chemically Stable Anion Exchange Membrane, Poly(2,6 dimethyl 1,4 phenylene oxide)-b-Poly(vinyl benzyl trimethyl ammonium), for Electrochemical Applications. *J. Electrochem. Soc.* **2016**, *163*, H513-H520.
12. Cheng, J.; He, G. H.; Zhang, F. X., A mini-review on anion exchange membranes for fuel cell applications: Stability issue and addressing strategies. *International Journal of Hydrogen Energy* **2015**, *40*, 7348-7360.
13. Becerra-Arciniegas, R. A.; Narducci, R.; Ercolani, G.; Antonaroli, S.; Sgreccia, E.; Pasquini, L.; Knauth, P.; Di Vona, M. L., Alkaline stability of model anion exchange membranes based on poly(phenylene oxide) (PPO) with grafted quaternary ammonium groups: Influence of the functionalization route. *Polymer* **2019**, *185*, 121931.
14. Kreuer, K. D.; Jannasch, P., A practical method for measuring the ion exchange capacity decrease of hydroxide exchange membranes during intrinsic degradation. *Journal of Power Sources* **2018**, *375*, 361-366.
15. Jasti, A.; Prakash, S.; Shahi, V. K., Stable and hydroxide ion conductive membranes for fuel cell applications: Chloromethylation and amination of poly(ether ether ketone). *Journal of Membrane Science* **2013**, *428*, 470-479.
16. Li, N. W.; Leng, Y. J.; Hickner, M. A.; Wang, C. Y., Highly Stable, Anion Conductive, Comb-Shaped Copolymers for Alkaline Fuel Cells. *Journal of the American Chemical Society* **2013**, *135*, 10124-10133.
17. Li, N. W.; Wang, L. Z.; Hickner, M., Cross-linked comb-shaped anion exchange membranes with high base stability. *Chemical Communications* **2014**, *50*, 4092-4095.
18. Dang, H. S.; Jannasch, P., Exploring Different Cationic Alkyl Side Chain Designs for Enhanced Alkaline Stability and Hydroxide Ion Conductivity of Anion-Exchange Membranes. *Macromolecules* **2015**, *48*, 5742-5751.
19. Varcoe, J. R.; Slade, R. C. T., Prospects for alkaline anion-exchange membranes in low temperature fuel cells. *Fuel Cells* **2005**, *5*, 187-200.
20. Jagur-Grodzinski, J., Polymeric materials for fuel cells: concise review of recent studies. *Polymers for Advanced Technologies* **2007**, *18*, 785-799.
21. Marino, M. G.; Kreuer, K. D., Alkaline Stability of Quaternary Ammonium Cations for Alkaline Fuel Cell Membranes and Ionic Liquids. *ChemSuschem* **2015**, *8*, 513-523.

22. Pan, Y.; Zhang, Q. D.; Yan, X. M.; Liu, J. F.; Xu, X. W.; Wang, T. Y.; El Hamouti, I.; Ruan, X. H.; Hao, C.; He, G. H., Hydrophilic side chain assisting continuous ion-conducting channels for anion exchange membranes. *Journal of Membrane Science* **2018**, *552*, 286-294.
23. Narducci, R.; Ercolani, G.; Becerra-Arciniegas, R. A.; Pasquini, L.; Knauth, P.; Di Vona, M. L., "Intrinsic" Anion Exchange Polymers through the Dissociation of Strong Basic Groups: PPO with Grafted Bicyclic Guanidines. *Membranes* **2019**, *9*.
24. Bauer, B.; Strathmann, H.; Effenberger, F., Anion-exchange membranes with improved alkaline stability. *Desalination* **1990**, *79*, 125-144.
25. Zadok, I.; Dekel, D. R.; Srebnik, S., Effect of Ammonium Cations on the Diffusivity and Structure of Hydroxide Ions in Low Hydration Media. *Journal of Physical Chemistry C* **2019**, *123*, 27355-27362.
26. Zhang, M.; Liu, J. L.; Wang, Y. G.; An, L. A.; Guiver, M. D.; Li, N. W., Highly stable anion exchange membranes based on quaternized polypropylene. *Journal of Materials Chemistry A* **2015**, *3*, 12284-12296.
27. Mohanty, A. D.; Tignor, S. E.; Krause, J. A.; Choe, Y. K.; Bae, C., Systematic Alkaline Stability Study of Polymer Backbones for Anion Exchange Membrane Applications. *Macromolecules* **2016**, *49*, 3361-3372.
28. Lin, C. X.; Wang, X. Q.; Hu, E. N.; Yang, Q.; Zhang, Q. G.; Zhu, A. M.; Liu, Q. L., Quaternized triblock polymer anion exchange membranes with enhanced alkaline stability. *J. Membr. Sci.* **2017**, *541*, 358-366.
29. Pan, J.; Han, J. J.; Zhu, L.; Hickner, M. A., Cationic Side-Chain Attachment to Poly(Phenylene Oxide) Backbones for Chemically Stable and Conductive Anion Exchange Membranes. *Chemistry of Materials* **2017**, *29*, 5321-5330.
30. Long, H.; Kim, K.; Pivovar, B. S., Hydroxide Degradation Pathways for Substituted Trimethylammonium Cations: A DFT Study. *Journal of Physical Chemistry C* **2012**, *116*, 9419-9426.
31. Liu, L.; Chu, X. M.; Liao, J. Y.; Huang, Y. D.; Li, Y.; Ge, Z. Y.; Hickner, M. A.; Li, N. W., Tuning the properties of poly(2,6-dimethyl-1,4-phenylene oxide) anion exchange membranes and their performance in H₂/O₂ fuel cells. *Energy & Environmental Science* **2018**, *11*, 435-446.
32. Zhu, L.; Yu, X. D.; Hickner, M. A., Exploring backbone-cation alkyl spacers for multi-cation side chain anion exchange membranes. *Journal of Power Sources* **2018**, *375*, 433-441.
33. Di Vona, M. L.; Casciola, M.; Donnadio, A.; Nocchetti, M.; Pasquini, L.; Narducci, R.; Knauth, P., Anionic conducting composite membranes based on aromatic polymer and layered double hydroxides. *Int. J. Hydrogen Energy* **2017**, *42*, 3197-3205.
34. Pizzoferrato, R.; Ciotta, E.; Ferrari, I. V.; Narducci, R.; Pasquini, L.; Varone, A.; Richetta, M.; Antonaroli, S.; Braglia, M.; Knauth, P.; Di Vona, M. L., Layered Double Hydroxides Containing an Ionic Liquid: Ionic Conductivity and Use in Composite Anion Exchange Membranes. *Chemelectrochem* **2018**, *5*, 2781-2788.
35. Derbali, Z.; Fahs, A.; Chailan, J. F.; Ferrari, I. V.; Di Vona, M. L.; Knauth, P., Composite anion exchange membranes with functionalized hydrophilic or hydrophobic titanium dioxide. *Int. J. Hydrogen Energy* **2017**, *42*, 19178-19189.
36. Sun, Z.; Lin, B. C.; Yan, F., Anion-Exchange Membranes for Alkaline Fuel-Cell Applications: The Effects of Cations. *Chemsuschem* **2018**, *11*, 58-70.
37. Liu, L.; Li, Q.; Dai, J. W.; Wang, H.; Jin, B. K.; Bai, R. K., A facile strategy for the synthesis of guanidinium-functionalized polymer as alkaline anion exchange membrane with improved alkaline stability. *J. Membr. Sci.* **2014**, *453*, 52-60.
38. Chalk, A. J.; Hay, A. S., Direct metalation of poly(2,6-dimethyl-1,4-phenylene ether). *Journal of Polymer Science Part A-1: Polymer Chemistry* **1969**, *7*, 691-705.
39. Wang, Z. Y.; Parrondo, J.; Ramani, V., Alkaline Stability of Poly(Phenylene Oxide) Based Anion Exchange Membranes Containing Imidazolium Cations. *J. Electrochem. Soc.* **2016**, *163*, F824-F831.
40. Pasquini, L.; Di Vona, M. L.; Knauth, P., Effects of anion substitution on hydration, ionic conductivity and mechanical properties of anion-exchange membranes. *New J. Chem.* **2016**, *40*, 3671-3676.
41. Liang, M., *Poly(phenylene oxide)*. 2011; Vol. 3.
42. Arges, C. G.; Wang, L. H.; Jung, M. S.; Ramani, V., Mechanically Stable Poly(arylene ether) Anion Exchange Membranes Prepared from Commercially Available Polymers for Alkaline Electrochemical Devices. *Journal of the Electrochemical Society* **2015**, *162*, F686-F693.

43. Parrondo, J.; Jung, M. S. J.; Wang, Z. Y.; Arges, C. G.; Ramani, V., Synthesis and Alkaline Stability of Solubilized Anion Exchange Membrane Binders Based on Poly(phenylene oxide) Functionalized with Quaternary Ammonium Groups via a Hexyl Spacer. *Journal of the Electrochemical Society* **2015**, *162*, F1236-F1242.
44. Teresa Perez-Prior, M.; Urena, N.; Tannenber, M.; del Rio, C.; Levenfeld, B., DABCO-functionalized polysulfones as anion-exchange membranes for fuel cell applications: Effect of crosslinking. *Journal of Polymer Science Part B-Polymer Physics* **2017**, *55*, 1326-1336.
45. Kortüm, G., *Treatise on Electrochemistry, 2nd edition*. 2nd ed.; Elsevier: Amsterdam, New York, 1965.
46. Hamann, C. H.; Hamnett, A.; Vielstich, W., *Electrochemistry*. 2nd Edition ed.; Wiley-VCH: Weinheim, 2007.
47. Braglia, M.; Ferrari, I. V.; Pasquini, L.; Djenizian, T.; Sette, M.; Di Vona, M. L.; Knauth, P., Electrochemical synthesis of thin, dense, and conformal anion exchange membranes with quaternary ammonium groups. *Electrochim. Acta* **2018**, *265*, 78-88.
48. Pasquini, L.; Knauth, P.; Pelzer, K.; Di Vona, M. L., Anion-conducting sulfaminated aromatic polymers by acid functionalization. *Rsc Advances* **2015**, *5*, 56636-56644.
49. Narducci, R.; Pasquini, L.; Chailan, J. F.; Knauth, P.; Di Vona, M. L., Low-Permeability Poly(ether Ether Ketone)-Based Ampholytic Membranes. *Chempluschem* **2016**, *81*, 550-556.
50. Narducci, R.; Chailan, J. F.; Fahs, A.; Pasquini, L.; Di Vona, M. L.; Knauth, P., Mechanical Properties of Anion Exchange Membranes by Combination of Tensile Stress-Strain Tests and Dynamic Mechanical Analysis. *Journal of Polymer Science Part B-Polymer Physics* **2016**, *54*, 1180-1187.
51. Sgreccia, E.; Di Vona, M. L.; Licoccia, S.; Sganappa, M.; Casciola, M.; Chailan, J. F.; Knauth, P., Self-assembled nanocomposite organic-inorganic proton conducting sulfonated poly-ether-ether-ketone (SPEEK)-based membranes: Optimized mechanical, thermal and electrical properties. *J. Power Sources* **2009**, *192*, 353-359.
52. Li, X. G., High-resolution thermogravimetry of poly(2,6-dimethyl-1,4-phenylene oxide). *Journal of Applied Polymer Science* **1999**, *71*, 1887-1892.
53. Ehlers, G. F. L.; Fisch, K. R.; Powell, W. R., Thermal degradation of polymers with phenylene units in the chain. I. Polyphenylenes and poly(phenylene oxides). *Journal of Polymer Science Part A-1: Polymer Chemistry* **1969**, *7*, 2931-2953.
54. Tomoi, M.; Yamaguchi, K.; Ando, R.; Kantake, Y.; Aosaki, Y.; Kubota, H., Synthesis and thermal stability of novel anion exchange resins with spacer chains. *J. Appl. Polym. Sci.* **1997**, *64*, 1161-1167.
55. Streitwieser, A., *Solvolytic Displacement Reactions*. Literary Licensing, LLC: 2012.

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