Tuning Polybenzimidazole Membrane by Immobilizing a Novel Ionic Liquid with Superior Oxygen Reduction Reaction Kinetics

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ABSTRACT: Protic ionic liquid (PIL) is a promising nonaqueous electrolyte alternative to replacing phosphoric acid for fuel cells operating at temperatures above 100 °C. In this study, the physical and electrochemical properties of stoichiometric and nonstoichiometric PILs are investigated focusing on their acid/base ratio. The study involves a series of PILs, generically indicated as \( \text{N}_2\text{N}-\text{diethyl-3-sulfopropane-1-ammonium trifluoromethanesulfonate} \) ([DESPA\(^+\)]\([\text{TfO}^-])\), varying from an excess of the proton acceptor (\( \text{N}_2\text{N}-\text{diethyl-3-aminopropane-1-sulfonic acid} \)) to an excess of the proton donor (trifluoromethanesulfonic acid, \( \text{TfOH} \)). Compared to a state-of-the-art electrolyte, i.e., concentrated phosphoric acid, the nonstoichiometric [DESPA\(^+\)]\([\text{TfO}^-]) shows superior oxygen reduction reaction kinetics on the investigated Pt catalysts and oxygen permeation ability (\( D_{\text{O}_2} \)). [DESPA\(^+\)]\([\text{TfO}^-]) with a base-to-acid molar ratio of 1:2 achieves a current density \( \sim 10 \) times larger than that of concentrated phosphoric acid at 110 °C and 0.8 V. Membranes including polybenzimidazole as a host polymer and stoichiometric and nonstoichiometric [DESPA\(^+\)]\([\text{TfO}^-]) as the conductive electrolyte exhibit promising properties in terms of thermal stability and conductivity. At 120 °C and 40% relative humidity, conductivities of 2 and 16 mS cm\(^{-1}\) are achieved by the membranes employing stoichiometric and excess acid [DESPA\(^+\)]\([\text{TfO}^-])

1. INTRODUCTION

In the last few decades, polymer electrolyte fuel cells (PEFCs) have attracted increased research attention as they allow the use of hydrogen as an alternative (chemical) energy storage medium for mobility as well as other applications like spacecraft or marine vessel.\(^1\)\(^-\)\(^3\) PEFCs are also of interest due to the rapid depletion of fossil fuel resources that is tied to the current global environmental concerns mainly anthropogenic CO\(_2\) emissions. PEFCs based on proton-conducting membranes consisting of perfluorinated polymers with sulfonic acid moieties (PFSA), e.g., AQUIVION or NAFION, have reached a mature stage of development regarding mobile applications.\(^4\)\(^-\)\(^6\) However, due to the limited operational range (up to 80 °C) of PEFCs, they suffer from a number of drawbacks such as large cooling system required, insufficient (if not any) waste heat recovery, and need for a water recirculation/feed gas humidification system.\(^5\)\(^7\)\(^-\)\(^10\) In fact, the proton conduction of the sulfonated fluorocarbon polymers depends on sufficient swelling with water. The solution to this latter limitation requires an electrolyte (membrane) with a conduction mechanism independent of the presence of high vapor pressure/liquid H\(_2\)O.

Several advantages are accompanied when increasing temperature above 100 °C: (i) A more efficient heat management/usable waste heat, (ii) an easier water manage-
mechanical stability, low gas permeability, and high proton conductivity at very low water vapor pressures. Experimental studies have shown that a conductivity of $4 \times 10^{-2} \; \text{S cm}^{-1}$ could be achieved at 190 °C for the H$_3$PO$_4$/PBI mole ratio of 1:5. Nevertheless, a major drawback of using H$_3$PO$_4$ as the electrolyte is the sluggish ORR kinetics due to the blocking effect of the catalytic sites and the low oxygen diffusion and solubility. This results in significantly lower power densities compared to PFSA-based electrolyte membranes.

Promising alternatives for nonaqueous electrolytes to H$_3$PO$_4$ are ionic liquids (ILs). In general, ILs are compounds that consist of a bulky (organic) cation and large anion with a low charge density, resulting in high lattice energies in the solid state and thus very low melting points. ILs exhibit low volatility and flammability, (often) moderate ionic conductivity, and a wide electrochemical stability window. A subgroup of the ILs are protic ionic liquids (PILs), which consist of a cation or an anion that possesses an exchangeable proton, which can serve as a mobile protonic charge carrier. When PILs are utilized as a nonaqueous electrolyte in PEFCs, high proton conductivity can be achieved. The properties of ILs can be tailored depending on the combination of cationic and anionic moieties which are selected. Among the several advantages of PILs, it is worth stressing that anions of superacids, e.g., TiOH, bis(trifluoromethanesulfonil)imide (TFSI), and fluoroboric acid (HBF$_4$) possess the advantage of weak adsorption on the catalysts, resulting in improved fuel cell performance due to the high numbers of free catalytic sites per unit area.

Many experimental studies have focused on the development of highly proton-conducting membranes employing different doping electrolytes other than H$_3$PO$_4$. Van de Ven et al. achieved a conductivity of 1.35 mS cm$^{-1}$ at 129 °C and 1.86 mS cm$^{-1}$ at 190 °C by impregnating a porous PBI membrane into IL 1-H-3-methylimidazolium bis(trifluoromethanesulfonil)imide. The proton conductivity of the pristine porous PBI membrane ($3 \times 10^{-3}$ mS cm$^{-1}$ at 134 °C and $7 \times 10^{-3}$ mS cm$^{-1}$ at 190 °C) was increased by a factor of 250–450 due to the uptake of the IL. Skorikova et al. proved that introducing an additional PIL into a H$_3$PO$_4$-doped PBI membrane is also beneficial to extend the lifetime of the gas diffusion electrode. Nair et al. prepared a membrane by blending poly(vinylidene fluoride-co-hexafluoropropylene) (PVDF-HFP) with the PIL diethylmethylammonium trifluoromethanesulfonate ([DEMA][TfO]) and nanosilica modified with HClO$_4$. With a loading of 80 wt % [DEMA][TfO], the blend membrane achieved a maximum conductivity of 0.6 mS cm$^{-1}$ at 100 °C. Wang et al. described the immobilization of 1-hexyl-3-methylimidazolium trifluoromethanesulfonate into a dense PBI membrane. The blend membrane exhibited good performance in terms of thermal stability, mechanical stability, and low methanol crossover. Neves et al. modified a NAFION membrane via immobilization of a certain fraction of IL cations. This led to the discovery of an effective approach to obtain a membrane with tailored properties. Two PILs, i.e., bis-(2-ethyl-hexyl)-ammonium hydrogen phosphate ([EHNH$_2$][H$_3$PO$_4$]) and imidazolium hexanolate ([Im][Hex]), were immobilized in a poly(vinylidene fluoride) (PVDF) membrane. The blended membranes showed enhanced proton conductivity, 33 mS cm$^{-1}$ for [EHNH$_2$][H$_3$PO$_4$]/PVDF and 69 mS cm$^{-1}$ for [Im][Hex]/PVDF at 60 °C, and good mechanical stability.

There are two widely applied methods for preparing blend membranes composed of an IL and a host polymer, namely, swelling and solution casting. Sood et al. investigated the influence of the blending method of NAFION and PILs based on a triethylammonium cation (TEA), leading to the finding that the preparation method has a great impact on the membrane’s morphology. The cast membranes have better long-range crystalline order and slightly higher correlation distance in the ionic domains. The morphology also plays an important role with respect to the thermomechanical properties and gas permeability. Cast NAFION membranes with 5–20 wt % TEA-based IL maintain a storage modulus of at least 0.5 MPa, even at 150–190 °C, whereas the swollen membranes with the same composition collapse before reaching 100 °C. Finally, the H$_2$/O$_2$ permeability coefficients of cast membranes are smaller by a factor of 1.1–1.4 than those of doped membranes.

In this study, a PIL with a highly acidic cation is investigated based on a potential application as an electrolyte in IT-PEFCs. This type of PIL can be prepared by means of a proton exchange reaction between an organic Bronsted base B and a Bronsted superacid HA, as per eq 1

$$B + HA \rightarrow HB^- + A^+ \quad (1)$$

In general, the anion $A^-$ (conjugated base) of a superacid (e.g., TiOH) only has a very small basicity and a negligible nucleophilicity. The acidity of the cation HB$^-$ depends on the strength of base B. A PIL with high acidity (or proton mobility) offers several advantages for future fuel cell applications. In the case of the presence of (residual) water or of an excess of the base B, a highly acidic PIL will significantly enhance conductivity. The proton can be transferred from the acidic cation to the free base or to the amphoteric water or back to another base molecule, respectively, mimicking the Grotthuss mechanism. Only slight conformational rearrangements of the proton-carrying ions are necessary to enable long-range motion. Besides, in a nonaqueous protic electrolyte, the protons necessary for the cathodic oxygen reduction reaction (ORR) cannot be delivered by the ubiquitous solvent water. In recent kinetic studies on PILs with various cation acidities, it emerged that the proton donor properties of the cation become rate-limiting; thus, it is an important parameter. An acidic PIL can also protonate the alkaline moieties in an ionogene polymer, e.g., a PBI. This will increase the interactions between the IL and the host material as well as help to immobilize the PIL at high levels.

An in-house synthesized PIL, N,N-diethyl-3-sulphopropyl-1-ammonium trifluoromethanesulfonate ([DESPA$^-$][TfO$^-$.]), was selected in this study due to its good thermal stability, conductivity, ORR kinetics, and fast oxygen transport, as reported in our previous work. Considering the water production and possible leaching of the PIL electrolyte, respectively, of the free acid and base during PEFC operation, an investigation of the PIL/H$_2$O blends with...
2. EXPERIMENTAL SECTION

2.1. Preparation of the Protic Ionic Liquids. All of the chemicals were used as received. The stoichiometric and nonstoichiometric PIL samples were prepared by mixing the calculated amounts of the \( N,N\)-diethyl-3-amino propane-1-sulfonic acid, with trifluoromethanesulfonic acid (reagent grade, 98%, Sigma-Aldrich), labeled as stoichiometric molar ratios \( xy \). A more detailed description of the preparation of the \( N,N\)-diethyl-3-amino propane-1-sulfonic acid can be found elsewhere.\(^\text{39,40}\) The base-to-acid molar ratio \( xy \) was varied from 1.5:1 to 1:2.5. As shown in a preceding study, the amount of \([\text{DESPA}^+][\text{TfO}^-]\) immobilized in the host PBI polymer has a major impact on the cell performance. Thus, the maximum degree of loading in the case of the nonstoichiometric PILs and the influence of the nonstoichiometry on the properties of the blend membranes are evaluated.

2.2. Characterization of the Stoichiometric and Nonstoichiometric \([\text{DESPA}^+][\text{TfO}^-]\) Electrolytes. The thermal behavior of the electrolytes was investigated by means of thermogravimetry analysis (TGA, PerkinElmer STA 6000). The samples were heat treated at 90 °C for 1 h to avoid a release of water during the measurement, followed by a heating program from 50 to 400 °C with a rate of 1 °C min \(^{-1}\) in an open quartz glass crucible under \( N_2 \) atmosphere with a flow rate of 20 mL min \(^{-1}\). The phase transition behavior was investigated by performing differential scanning calorimetry (DSC, TA Instruments Q2000 equipped with a liquid \( N_2 \) cooling system). The samples were sealed hermetically in Al pans and cycled in the range of -140 to 80 °C at a rate of 5 °C min \(^{-1}\). The electric conductivity was determined using a conductometer combined with a frequency analyzer and a thermostatic bath (MMates Italy). The water content was determined by adding ultrapure water (Millipore) and verified using Karl–Fischer titration (825 KF Titrand, Metrohm).

2.3. Preparation of the PBI–PIL Blend Membrane. The PBI–PIL blend membranes were prepared by means of solution casting. The PBI solution (O-PBI, Fumion AMLD, PBI polymer solution in DMAc, Fumatech) was dried under air at 80 °C for 3 days, which was followed by drying at 80 °C under vacuum overnight to eliminate the residual DMAc solvent. The \( ^{13} \text{C} \) NMR and \( ^1 \text{H} \) NMR spectra of the pristine PBI membrane are depicted in Figure S1. The average molar mass of the PBI sample was determined via gel permeation chromatography, and the results are shown in the Supporting Information. The dry PBI was dissolved in dimethyl sulfoxide (DMSO, ≥99.9%, VWR Chemicals) with a ratio of 1:50 (w/w) at 80 °C. An appropriate amount of \([\text{DESPA}^+][\text{TfO}^-]\) was mixed with the PBI solution to maintain a desired molar ratio (\( z \) mol %) in the blend membrane. The blend membranes were named in the form of: \( z \) mol \% - \( xy \). The PBI–PIL solutions were poured into a Petri dish, and the DMSO solvent was evaporated slowly at 50 °C under air for 2 days, followed by a drying process at 70 °C for 2 days. The residuals of the solvent were removed by a heat treatment in a vacuum oven overnight at 80 °C. An identical process was used for fabricating the blend membranes with the nonstoichiometric PILs (base-to-acid molar ratio of 1:1.5). A schematic sketch of the blend membrane preparation is shown in Figure 2. A visual inspection of the blend membranes allows a rough estimation of the homogeneity. The membrane becomes more inhomogeneous with an increasing amount of the PIL loading. The maximum molar ratio of stoichiometric \([\text{DESPA}^+][\text{TfO}^-]\) is up to 74 mol % and up to 65 mol % for the nonstoichiometric \([\text{DESPA}^+][\text{TfO}^-]\) (acid-to-base molar ratio of 1:1.5). A schematic drawing of the blend membrane preparation and a photograph of the blend membrane 74 mol %—1:1.

\[
f(r) = \frac{\tau}{4} + \frac{\tau^{1/2}}{2\tau^{1/2}} + 0.2146e^{-0.7823r^{-0.5}}
\]

\[
f(r) = \frac{4D_{O_2}t}{r^2}
\]

where \( n \) denotes the number of transferred electrons, \( F \) is the Faraday constant, \( r \) is the radius of the working electrode (WE), and \( t \) is the elapsed time.
1.5); see also Figure S2. Thus, only the homogeneous membrane samples are characterized in the following.

2.4. PBI−PIL Blend Membrane Characterization. The IR spectra of the PBI−PIL membrane were recorded at ambient temperature by means of an FT-IR spectrometer (Nicolet iS 20 FT-IR spectrometer, Thermo Fisher Scientific GmbH) equipped with an ATR unit using a diamond window (GladiATR single reflection, PIKE Technologies). The through-plane ionic conductivity was measured with an impedance analyzer (Solartron 1260) in a temperature- and humidity-controlled climate chamber (WK3600/40, Weiss Umwelttechnik GmbH). The cell was in-house-designed with a PVDF interior. Strips of the membrane samples with a size of 2 × 1.5 cm² were sandwiched between two Pt electrodes with a diameter of 1 mm (99.9%, MaTeck Material Technologie & Kristalle GmbH). The impedance was measured in a frequency range of 1−10⁶ Hz. The conductivity yields as

\[
\sigma = \frac{1}{R A}
\]

where \(\sigma\) is the total ionic conductivity, \(R\) is the measured membrane resistance, \(l\) is the distance between the electrode, and \(A\) is the cross-sectional area of the membrane.⁴⁵

The mechanical properties were measured under ambient conditions using a stress−strain material testing machine (ZwickRoell GmbH & Co. KG). The membrane samples were cut into strips 10 cm in length and 1 cm in width. The elongation rate was set to 5 mm min⁻¹. Each sample was tested three times, and the average value is provided here.

The atomic force microscopy (AFM) images were recorded with a Cypher S (Asylum Research, Santa Barbara) in tapping mode under ambient conditions. The images were processed using the Gwyddion 2.57 software.

The small-angle X-ray scattering (SAXS) patterns of the blend membranes were collected using a Xeuss 3.0c (Xenocs GmbH) equipped with an Eiger2 1M detector. The sample-to-SAXS detector distance was set to 800 mm. A Cu-Kα source was exploited with a beam size of 0.9 × 0.9 mm², by which a flux of ~10⁹ photons s⁻¹ could be obtained. The pattern collected for the direct beam was subtracted in the SAXS patterns as a background. [DESPA+[TfO⁻] blends with different stoichiometries are investigated as a function of the molar fraction of water (see Figure 4). The DSC patterns are shown in Figure S3. All of the IR spectra of the PBI−PIL blend membranes were collected using a Xeuss 3.0c (Xenocs − Grenoble, France) equipped with an Eiger2 1M detector. The wide-angle X-ray scattering (WAXS) patterns were collected using the same setup but equipped with an Eiger2 500K detector. The sample-to-WAXS detector distance was set to 800 mm. A Cu-Kα source was exploited with a beam size of 0.9 × 0.9 mm², by which a flux of ~10⁹ photons s⁻¹ could be obtained. The pattern collected for the direct beam was subtracted in the WAXS patterns as a background. The IR spectra of the PIL blend membranes were characterized in the following.

3. RESULTS AND DISCUSSION

3.1. Thermal Analysis of the Stoichiometric and Nonstoichiometric [DESPA]⁺[TfO⁻] Electrolyte. When performing TGA measurements, the samples with stoichiometry varying from 1.5:1 to 1:1 exhibited a one-step decomposition above 150 °C, indicating a sufficiently high thermal stability. With an increasing amount of excess acid present in the sample, a two-step decomposition process becomes apparent (see Figure 3). The first decomposition steps may be attributed to a loss of TfOH. Taking the sample with a stoichiometry of 1:2 as an example, the first weight loss of 31 wt % in the temperature range of 86−214 °C can be assigned to the loss of the (free) excess TfOH and residue water in the sample. This agrees with the initial composition of 30 wt % excess TfOH. The second decomposition step starts at 214 °C. The weight loss in the temperature range of 214−392 °C may be caused by PIL decomposition. Up to a temperature of 392 °C, a weight loss of 62 wt % can be observed. This may be explained by the decomposition of the PIL first through re-protonation of the trifluoromethanesulfonate anions and subsequent release of TfOH, then follows the decomposition of the remaining base. Thus, the total weight loss (62 wt %) in the temperature range of 214−392 °C agrees well with the adjusted sample composition of 70 wt % of the bound base and acid. The weight losses of the samples with excess acids are listed in Table S1.

Because of water production during PEFC operation, an investigation of the influence of residual water on the PIL electrolyte is necessary. The glass-transition temperatures \(T_g\) of the [DESPA⁺][TfO⁻] blends with different stoichiometries are investigated as a function of the molar fraction of water (see Figure 4). The DSC patterns are shown in Figure S3. All of the blends showed no crystallization or melting behavior. Only a glass transition could be observed in the investigated temperature range.
The glass-transition temperature \( T_g \) is influenced by two factors, namely, the amount of free excess acid and the water fraction. In general, \( T_g \) is decreasing with an increasing fraction of the excess acid. A similar trend can be observed in blends of 1.5:1 and 1:1 when considering the water influence, i.e., an increasing \( x_{H_2O} \) could lead to a decreasing \( T_g \). The influence of water on \( T_g \) is different in blends with excess acid. \( T_g \) first increases with an increasing water fraction and then decreases again. Taking a blend with a stoichiometry of 1:1.5 as an example, by adding water, the free acid may bond to it. When most of the free acid is bound at a composition of \( x_{H_2O} = 0.33 \), it corresponds to a (free) excess acid and \( H_2O \) molar ratio of 1:1, which accords well with the observed decreasing of \( T_g \) when the water fraction exceeds 0.33. A similar trend is observed in other blends with excess acid.

### 3.2. Conductivity of the Stoichiometric and Non-stoichiometric [DESPA]^+\[TfO^-] Electrolytes.

The conductivity measurements specify the total conductivity \( \sigma \) of the blends, i.e., the sum of the proton (cation) and anion partial conductivities. The temperature dependence of the conductivity \( \sigma \) of a viscous electrolyte (close to the glass-transition temperature) can be described by the Vogel–Fulcher–Tamman (VFT) equation

\[
\sigma = \sigma_0 \exp \left( -\frac{B_\sigma}{R(T - T_0)} \right)
\]

where \( B_\sigma \) denotes the pseudo-activation energy, \( T_0 \) is the temperature of zero configurational entropy, and \( \sigma_0 \) is the preexponential factor. The conductivity data of all investigated stoichiometries clearly obey a VFT behavior; see Figures S5a and S4a–d. The conductivity is influenced by stoichiometry, temperature, and water content. In the case of the base-excess blends (Figure S5a), the conductivity is enhanced with increasing water content. There is no experimental study regarding the pK\(_a\) value of the [DESPA]\(^+\), but for an analogue cation 2-sulfoethylmethylammonium, a pK\(_a\) value of 0.94 has been estimated in the literature.\(^{38,46}\) In addition, since the two cations have similar electron density distributions, as shown in Figure S6, assuming a pK\(_a\) value of the [DESPA]\(^+\) of around zero is reasonable. It is in the same order of magnitude as the hydroxonium cation with a pK\(_a\) = 0, which indicates significant protolysis.\(^{47,48}\)

\[
(C_2H_5)_2NH^+(CH_2)\_2SO_3H + H_2O
\]

\[
\rightarrow (C_2H_5)_2NH^+(CH_2)\_2SO_3^- + H_2O^+
\]

As was illustrated in our previous work, in the case of the stoichiometric [DESPA]\(^+[TfO^-]\) being neat or slightly moist \((x_{H_2O} < 0.25)\), the proton transfer may be mainly dominated by a vehicular mechanism.\(^{39,40}\) At a higher water concentration, more proton acceptor species (water, free base) are provided. The contribution to the conductivity via the cooperative mechanism may increase. As is depicted in Figure 5b, the conductivity increases with increasing water molar fraction.

In the case of the excess acid blends, water will also serve as a proton acceptor. Depending on the water fraction, it can be either protonated by the highly acidic excess acid (TfOH) or by the [DESPA]\(^+\) cation; see eqs 7 and 8

\[
\text{CR}_3\text{SO}_3H + H_2O \rightarrow \text{CR}_3\text{SO}_3^- + H_3O^+
\]
influence may be counteracted by the increased viscosity and the electrophoretic and relaxation effects.49

3.3. Electrochemical Properties. According to the thermal analysis results shown in Figure 3, the nonstoichiometric [DESPA⁺][TfO⁻] with a base-to-acid molar ratio of 1:2.5 shows a clear weight loss starting from 90 °C. Therefore, the electrochemical measurements were only performed with PILs with a base-to-acid molar ratio varying from 1:1 to 1:2.

The open-cell voltage (OCV) was determined by directly measuring the cell voltage at j = 0 using the potentiostat/impedance analyzer as a voltmeter with a high internal resistance. An OCV of 1.05 V can be measure for a Pt cathode at 90 °C under O₂-saturated conditions vs Pd-H using a stoichiometric [DESPA⁺][TfO⁻] electrolyte. Under the same conditions, values of 1.07 and 1.10 V can be measured for the blends with base-to-acid stoichiometries of 1:1.5 and 1:2, respectively. A similar trend can be observed at other experimental temperatures (100–120 °C). The OCV can be regarded as a rough indicator for the electrocatalytic activity, i.e., the higher the electrocatalytic activity, the higher OCV can be achieved and the closer to its Nernst potential.38

The polarization curves for the blends with a stoichiometry varied from 1:1 to 1:2 were recorded in the temperature range of 90–120 °C by scanning the potential in the negative direction from OCV to a potential where the limiting current density was reached. The results are depicted in Figure 7a. The ORR onset potential shifts significantly in the positive direction with an increasing temperature or when adding an excess of TfOH; see Figures 6 and 7a. An excess of TfOH accelerates the proton transfer to O₂ thus boosting the overall ORR kinetics. The significant shift of the ORR onset potentials in the blends with an excess TfOH could be explained by a change of the proton donor species, from H₃O⁺ and [DESPA⁺]⁺ (pKₐ = 0) to superacid TfOH (pKₐ = −14), as per eq 9. A similar effect in [DEMA][TfO] blends was observed by Goodwin et al.50 The ORR onset potential is shifted in the positive direction by 0.8 V when adding the acid in neat PIL.

\[ \frac{1}{2} O_2 + 2 e^- + TfOH \rightarrow 2[TfO]^- + H_2O \]  

Figure 6. Cyclic voltammograms in blends with a base-to-acid stoichiometry of 1:1 to 1:2 under a N₂ or O₂ atmosphere at 90 °C, with a scan rate of 100 mV s⁻¹.

Compared with the state-of-the-art electrolyte, concentrated H₃PO₄, a current density is achieved of 0.037 mA cm⁻² at a potential of 0.8 V (vs Pd-H) and 110 °C. Under the same conditions, stoichiometric and nonstoichiometric [DESPA⁺][TfO⁻] yield values of 0.15 mA cm⁻² (base-to-acid molar ratio of 1:1), 0.2 mA cm⁻² (1:1.5), and 0.37 mA cm⁻² (1:2). This is obviously a pronounced improvement in the presence of stoichiometric and nonstoichiometric [DESPA⁺][TfO⁻].

The value of the oxygen diffusion coefficient D_O₂ and equilibrium solubility c_O₂ in the blends with a stoichiometry varying from 1:1 to 1:2 are compiled in Table S2. The product of D_O₂ and c_O₂ and thus the O₂ permeability increases with increasing temperature and with increasing excesses of TfOH; see Figure 8. This correlates well with the measured limiting current densities |j|; see Figure 7b. The O₂ permeability D_O₂ c_O₂ in the 1:2 blend at 90 °C achieves an ∼5 times higher value compared to the 1:1 blend. All blends showed a significantly enhanced D_O₂ c_O₂ compared to concentrated H₃PO₄, e.g., 98% H₃PO₄ has a value of 1.05 × 10⁻¹² mol cm⁻¹ s⁻¹ at 100 °C.51

3.4. Thermal Analysis of the PIL–PBI Blend Membrane. A sufficient thermal stability of the PBI–IL blend membrane is crucial for application as an electrolyte in PEFCs above 100 °C. The thermal stabilities of the PIL, pristine PBI, and blend membranes are depicted in Figure 9.
The PBI blend membranes showed improved thermal stability compared to \([\text{DESPA}^+][\text{TfO}^-]\). The weight loss near 100 °C could be ascribed to a loss of water. Given that the PBI samples were blended with stoichiometric \([\text{DESPA}^+][\text{TfO}^-]\), the second decomposition step occurred between 285 and 550 °C. With respect to the thermal behavior of the stoichiometric PIL, the decomposition was presumably caused by the loss TfOH (back-transfer of the proton from \([\text{DESPA}]^+\) and from the imidazolium moieties of the PBI) and by the decomposition of the remaining free base. Most of the immobilized \([\text{DESPA}^+][\text{TfO}^-]\) decomposes below 550 °C. This assumption is in line with the initial sample composition (see Table S3). The PBI backbone degrades above 600 °C.

The PBI samples blended with the nonstoichiometric PIL exhibited a clear weight loss (about 3 wt %) in the temperature range of 157–244 °C, which could be attributed to the evaporation of a part of the excess acid (see Table S4). All investigated blend membranes showed sufficient thermal stability in the operating temperature range (100–120 °C).

3.5. ATR-IR. The ATR spectra of the free base, \([\text{DESPA}^+][\text{TfO}^-]\), the pristine PBI membrane, and the PBI–PIL blend membranes are depicted in Figure 10. In the spectrum of the pristine PBI, the bands present in the region between 4000 and 2000 cm\(^{-1}\) can be assigned to the stretching vibration modes of N–H, O–H, and C–H. The characteristic band at 3415 cm\(^{-1}\) is due to the isolated N–H stretching mode of the PBI imidazole moiety, i.e., a free N–H group. The bands at 3250–2500 cm\(^{-1}\) appear in the range that can be assigned to the stretching mode of the self-associated N–H bonds. The broad bands become stronger in the blend membrane samples, which may indicate a protonation of the imidazole ring by \([\text{DESPA}^+][\text{TfO}^-]\). Similar spectra were also observed in H\(_3\)PO\(_4\)-doped PBI membranes. The band at 1606 cm\(^{-1}\) corresponds to the aromatic C=\(\equiv\)N stretching mode of PBI and at 1534 cm\(^{-1}\) to the aromatic C–C stretching mode of the imidazole ring. These bands are stronger with the presence of the PIL in the PBI matrix and shift toward higher wavenumbers of 1632 and 1601 cm\(^{-1}\), respectively. This could be due to the protonation of the imidazole ring. No studies are available in the literature focusing on investigations by vibrational spectroscopy on \([\text{DESPA}^+][\text{TfO}^-]\). Thus, the literature data on other triflate-based ILs are used for comparison. The bands located at 1270 and 1032 cm\(^{-1}\) are assigned to the asymmetric and symmetric SO\(_3\) stretching modes. The asymmetric and symmetric stretching modes of CF\(_3\) are present at 1149 and 1222 cm\(^{-1}\).

3.6. Conductivity of the PIL–PBI Blend Membranes. In addition to the ORR kinetics, conductivity is one key parameter for assessing the fuel cell’s performance. To conduct a general evaluation of the PIL loading, water, and temperature influences on the conductivity, the blend membranes were measured from low to high relative humidity (10–40% RH) in the temperature range of 80–120 °C.

The temperature dependence of the total conductivity can be described by the Arrhenius equation, as is conveyed in eq 10.

\[
\sigma = \frac{\sigma_0}{T} \exp\left(-\frac{E_1}{RT}\right)
\]

Figure 8. Temperature dependences of the product of \(D_{O_2} \times c_{O_2}\) in blend electrolytes with a base-to-acid stoichiometry varied from 1:1 to 1:2.

Figure 9. Dynamic TG and DTG curves of the \([\text{DESPA}^+][\text{TfO}^-]\), pristine PBI, and blend membranes. The heating rate was 10 °C min\(^{-1}\). The membranes were pretreated at 100 °C for 30 min to reduce the influence of water.

Figure 10. ATR-IR spectra of the free \(N,N\)-diethyl-3-aminopropane-1-sulfonic acid, \([\text{DESPA}^+][\text{TfO}^-]\), pristine PBI, and the blend membranes.
where $E_a$ is the activation energy for the proton conduction, $\sigma_0$ is the preexponential factor, and $R$ is the ideal gas constant. The activation energy is an indicator of the energy required to perform the (rate-limiting) elementary step of the conduction mechanism.\footnote{The activation energies for proton conduction in the blend membranes are depicted in Figure 13. In general, $E_a$ decreases with increasing PIL loading. This indicates that a decreasing density of the polymer network due to a higher PIL fraction causes higher mobility among the proton-carrying ions. Increasing humidity (RH%) also has a positive effect on proton conduction. This can be explained by the increasing amount of $\text{H}_3\text{O}^+$ due to a protolysis with water, as discussed for the neat PILs. This is beneficial to proton conduction because a cooperative transport mechanism is possible, and the proton may move along a network of the H-bond chains formed by $\text{H}_2\text{O}^+$ and $\text{H}_2\text{O}$. Moreover, considering the geometry factor, $\text{H}_2\text{O}^+$ may diffuse faster than $[\text{DESPA}]^+$, which could also lead to faster vehicular transport. The membrane samples with an excess of TfOH exhibit the lowest values of $E_a$ compared to other blend membranes under the same conditions. In the case of the membrane sample 65 mol \%--1:1.5, a value of only 26.1 kJ mol\(^{-1}\) is measured at 120 °C and 40% RH. The incorporation of TfOH in the blend membrane may have a positive influence on the formation of a domain structure of the polymer and the incorporated PIL by forming H bonds to $[\text{DESPA}]^+$ and $[\text{TfO}]^-$\footnote{The lifetime of a polymer electrolyte fuel cell is not only influenced by the chemical degradation of the catalyst and membrane but also by the latter’s mechanical failure. Performing stress–strain tests, all blend membranes exhibit a higher elongation compared to the pristine PBI membranes. The results of this are listed in Table 1. From the literature, it is known that PBI membranes with a high degree of $\text{H}_3\text{PO}_4$ doping also suffer losses of mechanical stability.\footnote{When introducing the acidic PIL into the PBI membrane, an analogous deterioration of the mechanical properties can be observed.} Both may significantly accelerate proton transport.}.

The conductivity is improved by increasing the temperature or relative humidity. It is decreased with an increasing PBI molar fraction, as is depicted in Figure 11a,b. Moreover, Figure 11b clearly suggests that the excess acid in the blend membrane brings a significant improvement in conductivity. At 120 °C and 10% RH, the conductivity of the blend membrane (74 mol \%--1:1) exhibits a value of $9.1 \times 10^{-4}$ S cm\(^{-1}\), in the blend membrane (65 mol \%--1:1.5) of $9.5 \times 10^{-4}$ S cm\(^{-1}\) under the same experimental conditions. The conductivity increases by 1 order of magnitude.

The $[\text{TfO}]^-$ anions can interact with the PBI’s imidazole rings (N–H). Depending on the PIL/PBI ratio and the stoichiometry of the PIL (acid excess), hydronium ions ($\text{H}_2\text{O}^+$) are formed due to the protolysis equilibrium in the presence of the water. The proton can move through the H-bond network, as depicted in Figure 12. Furthermore, a vehicular mechanism is possible via the movement of the proton-carrying ions, i.e., $[\text{DESPA}]^+$ and $\text{H}_3\text{O}^+$\footnote{This is beneficial to proton conduction because a cooperative transport mechanism is possible, and the proton may move along a network of the H-bond chains formed by $\text{H}_2\text{O}^+$ and $\text{H}_2\text{O}$. Moreover, considering the geometry factor, $\text{H}_2\text{O}^+$ may diffuse faster than $[\text{DESPA}]^+$, which could also lead to faster vehicular transport. The membrane samples with an excess of TfOH exhibit the lowest values of $E_a$ compared to other blend membranes under the same conditions. In the case of the membrane sample 65 mol \%--1:1.5, a value of only 26.1 kJ mol\(^{-1}\) is measured at 120 °C and 40% RH. The incorporation of TfOH in the blend membrane may have a positive influence on the formation of a domain structure of the polymer and the incorporated PIL by forming H bonds to $[\text{DESPA}]^+$ and $[\text{TfO}]^-$. Both may significantly accelerate proton transport.}. The proton conduction in the blend membrane under humidified conditions may follow a cooperative mechanism and support by means of a vehicular mechanism. However, considering only the results from this study, it is difficult to specify the contribution of these two mechanisms and further investigation is necessary. Moreover, the PIL may also act as a plasticizer that is beneficial to the flexibility of the PBI backbone.\footnote{The lifetime of a polymer electrolyte fuel cell is not only influenced by the chemical degradation of the catalyst and membrane but also by the latter’s mechanical failure. Performing stress–strain tests, all blend membranes exhibit a higher elongation compared to the pristine PBI membranes. The results of this are listed in Table 1. From the literature, it is known that PBI membranes with a high degree of $\text{H}_3\text{PO}_4$ doping also suffer losses of mechanical stability.\footnote{When introducing the acidic PIL into the PBI membrane, an analogous deterioration of the mechanical properties can be observed.} Both may significantly accelerate proton transport.} The activation energy is an indicator of the energy required to perform the (rate-limiting) elementary step of the conduction mechanism.\footnote{The activation energies for proton conduction in the blend membranes are depicted in Figure 13. In general, $E_a$ decreases with increasing PIL loading. This indicates that a decreasing density of the polymer network due to a higher PIL fraction causes higher mobility among the proton-carrying ions. Increasing humidity (RH%) also has a positive effect on proton conduction. This can be explained by the increasing amount of $\text{H}_3\text{O}^+$ due to a protolysis with water, as discussed for the neat PILs. This is beneficial to proton conduction because a cooperative transport mechanism is possible, and the proton may move along a network of the H-bond chains formed by $\text{H}_2\text{O}^+$ and $\text{H}_2\text{O}$. Moreover, considering the geometry factor, $\text{H}_2\text{O}^+$ may diffuse faster than $[\text{DESPA}]^+$, which could also lead to faster vehicular transport. The membrane samples with an excess of TfOH exhibit the lowest values of $E_a$ compared to other blend membranes under the same conditions. In the case of the membrane sample 65 mol \%--1:1.5, a value of only 26.1 kJ mol\(^{-1}\) is measured at 120 °C and 40% RH. The incorporation of TfOH in the blend membrane may have a positive influence on the formation of a domain structure of the polymer and the incorporated PIL by forming H bonds to $[\text{DESPA}]^+$ and $[\text{TfO}]^-$. Both may significantly accelerate proton transport.}

The conductivity is improved by increasing the temperature or relative humidity. It is decreased with an increasing PBI molar fraction, as is depicted in Figure 11a,b. Moreover, Figure 11b clearly suggests that the excess acid in the blend membrane brings a significant improvement in conductivity. At 120 °C and 10% RH, the conductivity of the blend membrane (74 mol \%--1:1) exhibits a value of $9.1 \times 10^{-4}$ S cm\(^{-1}\), in the blend membrane (65 mol \%--1:1.5) of $9.5 \times 10^{-4}$ S cm\(^{-1}\) under the same experimental conditions. The conductivity increases by 1 order of magnitude.

The conductivity is improved by increasing the temperature or relative humidity. It is decreased with an increasing PBI molar fraction, as is depicted in Figure 11a,b. Moreover, Figure
topography correspond to a higher phase; see Figure 14b and the line profiles in Figure 14c. Also, on other positions on the surface, an increase in the phase beside the grooves can be identified; see Figure 14e,f. It can be concluded that the surface has a certain degree of inhomogeneity, which could indicate the formation of clusters or channels consisting of soft PIL-rich material embedded in the hydrophobic PBI matrix.

As it is visible to the naked eye, all of the blend membranes appear similar at ambient pressure. When applying vacuum, the appearance of the blend membranes with high PIL loading (74 mol %−1:1 and 70 mol %−1:1) is changed; see Figure 15. The blend membrane (70 mol %−1:1) shrinks, and the surface takes on a dry and brittle appearance. Cracks are visible. When the PIL loading is increased to 74 mol %, the membrane seems to leach out the PIL and a wormlike droplet structure can be observed on the surface. However, the samples, 70 mol %−1:1 and 74 mol %−1:1 revert to their original appearance when the vacuum is broken, which indicates the reversibility of the process.

The SAXS and WAXS patterns of the PIL, pristine PBI, and blend membranes are depicted in Figure 16a,b. The characteristic $I(q)$ in low-$q$ regions (at 0.03 Å$^{-1}$) change with the PIL contents; see Figure 16. It disappears in the pristine PBI and the 54 mol %−1:1 sample and reaches a maximum in the 63 mol %−1:1 sample. When increasing the PIL loading beyond 63 mol %, it decreases again. This is generally associated with a crystallization of the polymer backbone (intercrystalline domains). At high $q$ values of 0.25 Å$^{-1}$, a characteristic peak appears when adding the PIL to the host polymer. This is attributed to side-chain aggregation (cluster domain) and corresponds to the size of the hydrophilic channels. The ionomer peak shifts from 0.258 to 0.206 Å$^{-1}$ when increasing the PIL content (from 54 to 74 mol %) and the characteristic size of the cluster domain changes from 2.434 to 3.048 nm as expected; see Table 2. The peak becomes less recognizable for the samples, 70 mol %−1:1 and 74 mol %−1:1. This may be explained by the crystallization of the polymer and leaching out of the PIL; see Figure 15 (C4 and CS).

Table 1. Mechanical Properties of the Pristine PBI and the Blend Membranes with a Stoichiometric PIL Loading Varied from 54 to 74 mol % and a Blend Membrane with a Nonstoichiometric PIL (Base-to-Acid Molar Ratio of 1:1.5) Loading of 65 mol %

<table>
<thead>
<tr>
<th>membrane</th>
<th>Young's modulus (MPa)</th>
<th>ultimate tensile strength (MPa)</th>
<th>elongation at break (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>pristine PBI</td>
<td>177.5 ± 25.0</td>
<td>151.8 ± 14.7</td>
<td>65.5 ± 7.6</td>
</tr>
<tr>
<td>54 mol %−1:1</td>
<td>21.5 ± 3.5</td>
<td>71.3 ± 8.1</td>
<td>184.0 ± 5.4</td>
</tr>
<tr>
<td>63 mol %−1:1</td>
<td>8.7 ± 2.3</td>
<td>24.8 ± 6.0</td>
<td>188.7 ± 11.0</td>
</tr>
<tr>
<td>74 mol %−1:1</td>
<td>6.6 ± 0.6</td>
<td>23.7 ± 4.8</td>
<td>260.3 ± 9.8</td>
</tr>
<tr>
<td>65 mol %−1:1.5</td>
<td>8.7 ± 2.1</td>
<td>13.8 ± 2.9</td>
<td>125.9 ± 11.6</td>
</tr>
<tr>
<td>65 mol %−1:1.5</td>
<td>8.7 ± 2.1</td>
<td>13.8 ± 2.9</td>
<td>125.9 ± 11.6</td>
</tr>
</tbody>
</table>
Figure 14. Tapping-mode AFM analysis of the blend membrane: 74 mol %−1:1. Maps of topography and phase with a representative line profile for a 7 × 7 μm² scan (a−c) and for a 1 × 1 μm² scan (d−f).

Figure 15. Photographs of the blend membranes in the SAXS instrument sample chamber (p = 15 μbar). C1: pristine PBI; C2: sample 54 mol %−1:1; C3: sample 63 mol %−1:1; C4: sample 70 mol %−1:1; C5: sample 74 mol %−1:1.
electrolytes were prepared via solution casting. In general, the conductivity and mechanical stability relate to the immobilized amounts of PIL. A large amount of PIL is advantageous in terms of conductivity but results in a loss of mechanical stability. Moreover, the morphology of the blend membrane is significantly influenced by the amount of PIL that is utilized. This study opens the possibility of applying highly acidic PILs that have been immobilized in a host polymer to future fuel cell applications.

**Associated Content**

**Supporting Information**
The Supporting Information is available free of charge at https://pubs.acs.org/doi/10.1021/acschemlett.1c03819.

$^{13}$C NMR and $^1H$ NMR spectra of the PBI in DMSO-$d_6$, photographs of PBI blend membranes, molar mass of PBI sample, calculated weight loss from TG curves of samples with a stoichiometry (x/y) from 1:1.5 to 1:2.5, DSC heating traces of stoichiometric [DESFA$^+$$][$TFO$^-]$ with varied water molar fractions, conductivity dependence on water molar fractions in blends with a base-to-acid stoichiometry from 1:1 to 1:2.5 in the temperature range of 40–120 °C, electron density distributions of the [DESFA$^+$] and [2-Sema$^-$] cations, value of the oxygen diffusion coefficient and solubility in the blends, calculated weight loss of the pristine PBI and PIL–PBI blend membranes from the TG curves, and leakage tests of the PIL–PBI blend membranes (PDF).

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**Figure 16.** (a) SAXS patterns and (b) WAXS patterns of the pristine PBI, blend PBI membranes, and [DESFA$^+$$][$TFO$^-]$.

**Table 2. Characteristic Peak Positions of the Blend Membranes and Typical Size of the Ionic Cluster**

<table>
<thead>
<tr>
<th>Membrane</th>
<th>Ionomeric peak position ($\text{Å}^{-1}$)</th>
<th>Characteristic size (nm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>54 mol %:1:1</td>
<td>0.258</td>
<td>2.434</td>
</tr>
<tr>
<td>63 mol %:1:1</td>
<td>0.246</td>
<td>2.553</td>
</tr>
<tr>
<td>70 mol %:1:1</td>
<td>0.233</td>
<td>2.695</td>
</tr>
<tr>
<td>74 mol %:1:1</td>
<td>0.206</td>
<td>3.048</td>
</tr>
</tbody>
</table>

**4. CONCLUSIONS**

In this study, stoichiometric and nonstoichiometric samples of the PIL, N,N-diethyl-3-sulfopropyl-ammonium trifluoromethanesulfonate [DESFA$^+$][TFO$^-$], were prepared and their physical and electrochemical properties were evaluated. An excess of the base, N,N-diethyl-3-aminopropane-1-sulfonic acid, enhances the thermal stability and an excess of trifluoromethanesulfonic acid is beneficial to the proton conduction, ORR kinetics, and oxygen permeation. The physical and electrochemical analysis offers a promising prospect by adding an excess of acid to a PIL for use in future PEFCs at elevated operating temperatures.

PBI blend membranes may play a key role in developing PEFC applications operating at 100–120 °C. Blend membranes based on stoichiometric and nonstoichiometric PIL
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Notes
The authors declare no competing financial interest.

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REFERENCES


