



Cite this: *Phys. Chem. Chem. Phys.*,
2017, **19**, 28876

Exploring the effect of fluorinated anions on the CO₂/N₂ separation of supported ionic liquid membranes†

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The CO₂ and N₂ permeation properties of ionic liquids (ILs) based on the 1-ethyl-3-methylimidazolium cation ([C₂mim]⁺) and different fluorinated anions, namely 2,2,2-trifluoromethylsulfonyl-*N*-cyanoamide ([TFSAM]⁻), bis(fluorosulfonyl) imide ([FSI]⁻), nonafluorobutanesulfonate ([C₄F₉SO₃]⁻), tris(pentafluoroethyl)trifluorophosphate ([FAP]⁻), and bis(pentafluoroethylsulfonyl)imide ([BETI]⁻) anions, were measured using supported ionic liquid membranes (SILMs). The results show that pure ILs containing [TFSAM]⁻ and [FSI]⁻ anions present the highest CO₂ permeabilities, 753 and 843 Barrer, as well as the greatest CO₂/N₂ permselectivities of 43.9 and 46.1, respectively, with CO₂/N₂ separation performances on top of or above the Robeson 2008 upper bound. The re-design of the [TFSAM]⁻ anion by structural unfolding was investigated through the use of IL mixtures. The gas transport and CO₂/N₂ separation properties through a pure [C₂mim][TFSAM] SILM are compared to those of two different binary IL mixtures containing fluorinated and cyano-functionalized groups in the anions. Although the use of IL mixtures is a promising strategy to tailor gas permeation through SILMs, the pure [C₂mim][TFSAM] SILM displays higher CO₂ permeability, diffusivity and solubility than the selected IL mixtures. Nevertheless, both the prepared mixtures present CO₂ separation performances that are on top of or above the Robeson plot.

Received 14th September 2017,
Accepted 9th October 2017

DOI: 10.1039/c7cp06297d

rs.c.li/pccp

Introduction

The development of supported ionic liquid membranes (SILMs) for CO₂ separation has been widely investigated in recent years mainly due to their easy preparation and versatility.^{1–3} In contrast to traditional liquid membranes, which are produced by impregnating a porous membrane support with common organic solvents,⁴ SILMs use ionic liquids (ILs) and thus benefit from negligible displacement of the liquid phase from the membrane pores through evaporation,^{5,6} due to the low volatility of ILs.⁷ It should also be emphasized that within the CO₂ separation context, the most important features of ILs are their high CO₂ affinity over light gases^{8–10} and their inherent designer nature that enables the tailoring of IL properties by proper selection

of cations and/or anions or *via* the addition of specific functional groups.

Numerous works have investigated the effect of IL chemical structure on the gas permeation properties of SILMs. A broad diversity of cations, such as imidazolium,¹¹ triazolium,¹² thiazolium,¹³ pyridinium,¹⁴ cholinium,¹⁵ ammonium,¹⁶ and phosphonium,¹⁷ combined with halogens, sulfonates, carboxylates, fluorinated or cyano-functionalized anions, have been studied. Other works, mostly focusing on imidazolium-based ILs, have also explored the effect of alkyl,¹⁸ fluoroalkyl,¹⁹ etoxyalkyl,²⁰ and aminoalkyl²¹-functionalized cations. Since IL anions have a stronger influence on the CO₂ separation performance of SILMs than IL cations,¹ they deserved from the start a closer look. The first studies on SILMs made use of fluorinated anions such as bis(trifluoromethylsulfonyl)imide [NTf₂]⁻, tetrafluoroborate [BF₄]⁻, and hexafluorophosphate [PF₆]⁻ and enabled drawing conclusions about the CO₂-phylic behaviour and high CO₂ permeabilities of these anions.²² More recently, low viscous ILs with cyano-functionalized anions, such as tricyanomethanide [C(CN)₃]⁻ and tetracyanoborate [B(CN)₄]⁻,^{23–25} have been recognized as better candidates for the development of improved SILMs, because of their superior CO₂ permeabilities and permselectivities when compared to the most used [NTf₂]⁻ anion. Task-specific ILs bearing amine groups, such as those containing amino acid

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† Electronic supplementary information (ESI) available. See DOI: 10.1039/c7cp06297d



anions,^{26–28} have also been proposed to prepare SILMs, since amine groups can chemically bond CO₂ and act as carriers for CO₂ facilitated transport through SILMs at low pressures. However, the high viscosity of these task-specific ILs is undoubtedly a key limitation, as CO₂ diffusion is strongly compromised.

In an effort to improve the CO₂ permeability and permselectivity properties of SILMs, our recent studies explored the use of IL mixtures by fixing the [C₂mim]⁺ cation and researching on different anion chemical structures. Initially, SILMs based on IL mixtures combining anions with different CO₂ solubility behaviours were investigated: thiocyanate ([SCN][−]), dicyanamide ([N(CN)₂][−]) and bis(trifluoromethylsulfonyl)imide ([NTf₂][−]) that present physical solubility; acetate ([Ac][−]) and lactate ([Lac][−]), which additionally have chemical solubility.²⁹ Afterwards, we focused on IL mixtures based on sulfate ([CH₃SO₄][−]) and cyano-functionalized anions ([SCN][−], [N(CN)₂][−], [C(CN)₃][−] and [B(CN)₄][−]).³⁰ Moreover, we studied IL mixtures containing [C(CN)₃][−] and different amino acid anions, so that one IL component maintains the low viscosity, while the other provides the desired chemical characteristics for the active transport of CO₂.³¹ The overall results of these studies showed that mixing anions with specific chemical features allows variations in IL viscosity and molar volume that significantly impact the gas transport through SILMs, and thus tailored permeabilities and permselectivities can be achieved.^{29–31}

In the present work, the gas permeation properties and CO₂/N₂ separation performance of SILMs prepared with pure ILs bearing the [C₂mim]⁺ cation and different less conventional fluorinated anions, namely [TFSAM][−], [FSI][−], [C₄F₉SO₃][−], [FAP][−] and [BETI][−], were evaluated and the effect of the fluorinated moieties in the IL anion was discussed. Despite the fact that several SILMs with common fluorinated anions have already been reported,^{1–3} the gas permeation properties of SILMs containing fluorinated anions, such as those selected herein, have still not been properly studied and discussed. Only three studies have reported SILMs that made use of [C₄F₉SO₃][−], [FAP][−] and [BETI][−] anions. Pereira *et al.*³² conducted single gas permeation experiments through a [C₂C₁py][C₄F₉SO₃] SILM, at 294 K and 75 kPa, using CO₂, N₂, O₂, hydrocarbon gases (CH₄, C₂H₆, C₃H₈, C₃H₆) and perfluorocarbon gases (CF₄, C₂F₆, C₃F₈). Scovazzo *et al.*²² determined ideal/mixed CO₂/CH₄ and CO₂/N₂ permselectivities in a [C₄mim][BETI] SILM at 303 K and 200 kPa, while Althuluth *et al.*³³ reported ideal/mixed CO₂/CH₄ permselectivities in a [C₂mim][FAP] SILM at 313 K and 700 kPa. Nevertheless, the obtained results cannot be directly compared due to the different measurement conditions, as well as the use of diverse IL cation structures.

Additionally, this work investigates the impact on gas transport through SILMs of using a pure IL *versus* a structurally similar IL mixture as the liquid phase. Inspired by the fact that the [TFSAM][−] anion has an unusual asymmetric chemical structure, which combines both fluorinated and cyano functionalities, the re-design of the chemical structure of a pure [C₂mim][TFSAM] IL through the use of IL mixtures is explored. For that purpose, different pairs of ILs, based on the [C₂mim]⁺ cation and anions containing fluorinated or cyano functionalities, were selected and

their gas permeation properties were compared to those of the pure [C₂mim][TFSAM] SILM. One of the IL mixtures contains [NTf₂][−] and [N(CN)₂][−] anions, whose gas permeation properties were previously determined,²⁹ whereas the other IL mixture is based on [OTf][−] and [SCN][−] anions and its gas transport properties are reported here for the first time.

Results and discussion

Gas permeation through SILMs having fluorinated anions

The structures of the pure ILs bearing fluorinated anions are depicted in Fig. 1.

The water content (wt%), molar mass (*M*), viscosity (*η*), density (*ρ*) and molar volume (*V_m*) values of the pure ILs used as liquid phases in the studied SILMs are summarized in Table 1. The thermophysical properties of the conventional [C₂mim][NTf₂] IL are also included for comparison.²⁹ From Table 1, it can be observed that the IL containing the [C₄F₉SO₃][−] anion shows the highest viscosity, while [C₂mim][FSI] presents the lowest viscosity. The IL viscosity values can be organized following the IL anion order: [C₄F₉SO₃][−] > [BETI][−] > [FAP][−] > [NTf₂][−] > [TFSAM][−] > [FSI][−]. A slightly different trend was observed for molar volumes, with the IL comprising the [FAP][−] anion showing the highest molar volume, while [C₂mim][FSI] exhibiting the lowest molar volume. These data will be used subsequently in the understanding of the gas permeation results.

The experimental gas permeability (*P*) values obtained through the prepared SILMs having ILs with fluorinated anions, measured at 293 K with a *trans*-membrane pressure differential of 100 kPa, are shown in Table 2. To the best of our knowledge, the CO₂/N₂ separation properties of [C₂mim][TFSAM], [C₂mim][FSI], [C₂mim][FAP], [C₂mim][BETI] and [C₂mim][C₄F₉SO₃] SILMs are reported here for the first time, while those of the [C₂mim][NTf₂] SILM were previously determined using the same experimental conditions.²⁹ It is important to mention that in order to attain stable SILMs, both hydrophilic and hydrophobic supports were used according to the hydrophobicity of ILs, and the results are compared in this section, irrespective of the support used.

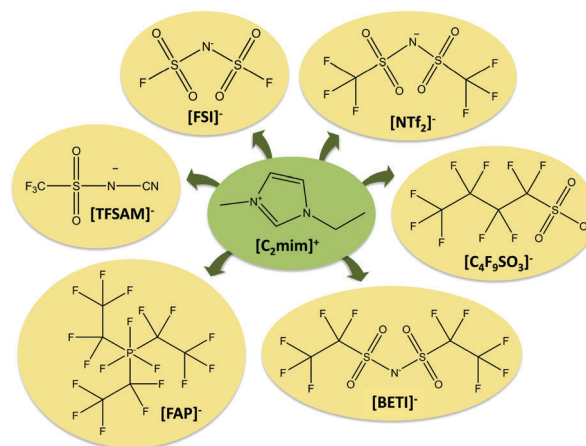


Fig. 1 Chemical structures of the pure ionic liquids (ILs) used in this work to prepare SILMs.



Table 1 Water contents and thermophysical properties of the pure ILs

IL sample	H ₂ O (wt%)	<i>M</i> (g mol ⁻¹)	η^a (mPa s)	ρ^b (g cm ⁻³)	<i>V_m</i> (cm ³ mol ⁻¹)
[C ₂ mim][TFSAM]	0.02	284.26	23.7	1.3518	210.3
[C ₂ mim][FSI]	0.09	291.30	22.4	1.4480	201.2
[C ₂ mim][NTf ₂] ^c	0.02	391.31	39.1	1.5240	256.8
[C ₂ mim][FAP] ^d	—	556.17	76.4	1.7081	325.6
[C ₂ mim][BETI]	0.02	491.33	85.5	1.5989	307.3
[C ₂ mim][C ₄ F ₉ SO ₃]	0.08	410.26	109.7	1.5420	266.1

^a Viscosity (η) and density (ρ) measured at 293 K, except for [C₂mim][C₄F₉SO₃] IL (303 K). ^b Molar volume (*V_m*) obtained for 293 K, except for [C₂mim][C₄F₉SO₃] IL (303 K). ^c The values of [C₂mim][NTf₂] were taken from Tomé *et al.*²⁹ ^d The density and viscosity values of [C₂mim][FAP] were taken from Neves *et al.*⁵²

Table 2 Gas permeabilities (*P*)^a through the prepared SILMs of pure ILs^b

SILM sample	<i>P</i> (CO ₂)	<i>P</i> (N ₂)	α (CO ₂ /N ₂)
[C ₂ mim][TFSAM]	753 ± 0.2	17 ± 0.1	43.9 ± 0.1
[C ₂ mim][FSI]	843 ± 0.5	18 ± 0.2	46.1 ± 0.5
[C ₂ mim][NTf ₂] ^c	589 ± 1.0	17 ± 0.1	35.5 ± 0.3
[C ₂ mim][FAP]	624 ± 0.4	24 ± 0.1	26.0 ± 0.1
[C ₂ mim][BETI]	437 ± 1.9	18 ± 0.1	24.8 ± 0.2
[C ₂ mim][C ₄ F ₉ SO ₃]	32 ± 0.2	6 ± 0.1	5.5 ± 0.1

^a Barrer (1 Barrer = 10⁻¹⁰ cm_(STP)³ cm cm⁻² s⁻¹ cmHg⁻¹). ^b The listed uncertainties represent the standard deviations, based on three experiments. ^c Values taken from Tomé *et al.*²⁹

From Table 2, the same trend in gas permeability is valid for all the studied SILMs: *PCO*₂ ≫ *PN*₂, as expected. Regarding the influence of the fluorinated-based anions, SILMs having the [FSI]⁻, [TFSAM]⁻ and [FAP]⁻ anions present higher CO₂ permeabilities of 843, 753 and 624 Barrer, respectively, than the SILM containing the [NTf₂]⁻ anion, which is well-known for its high CO₂ permeability (589 Barrer).²⁹ It should be noted that in spite of the similar structures of [NTf₂]⁻ and [FSI]⁻ anions, in which the difference consists in two extra ⁻CF₃ groups in the [NTf₂]⁻ anion structure (Fig. 1), the CO₂ permeability through the [C₂mim][FSI] SILM is ~1.5 times higher than that through the [C₂mim][NTf₂] SILM. Generally, CO₂ permeabilities through the studied SILMs are found to decrease in the following IL anion order: [FSI]⁻ > [TFSAM]⁻ > [FAP]⁻ > [NTf₂]⁻ > [BETI]⁻ > [C₄F₉SO₃]⁻ (Table 2). Considering the IL anion viscosity trend, obtained at 293 K (Table 1), [FSI]⁻ < [TFSAM]⁻ < [NTf₂]⁻ < [FAP]⁻ < [BETI]⁻ < [C₄F₉SO₃]⁻, it can be concluded that these experimental data are in agreement with the general trend usually observed in the literature, where ILs with high viscosities yield SILMs with low gas permeabilities.^{3,14,15,23,40} However, [C₂mim][FAP] is the only exception since it presents a different behaviour (Table 2): despite its high viscosity (76.4 mPa s), it also exhibits high CO₂ permeability (624 Barrer), higher than those of the [C₂mim][NTf₂] IL (39.1 mPa s and 589 Barrer). Notice that the [FAP]⁻ anion has the most different chemical structure among all the IL anions studied in this work, consisting of a phosphorus atom surrounded by fluorine atoms, without sulfonyl functional groups (Fig. 1). Moreover, taking a closer look at the gas permeabilities obtained through SILMs immobilized with the remaining ILs, it can be seen that

higher CO₂ permeabilities are achieved for ILs with anions bearing a smaller number of fluorine elements, such as [TFSAM]⁻ and [FSI]⁻ anions (Table 2).

Gas diffusivity (*D*) is a mass transfer property that directly accounts for gas permeability (eqn (1)). Typically, the higher the gas diffusivity, the faster is the gas flux through the SILM. The experimental CO₂ and N₂ diffusivity values obtained through the prepared SILMs are presented in Table 3. The CO₂ diffusivities of the SILMs with fluorinated anions can be ordered as follows: [FSI]⁻ > [TFSAM]⁻ > [FAP]⁻ > [NTf₂]⁻ > [BETI]⁻ > [C₄F₉SO₃]⁻, which fully corresponds to the IL anion order observed for CO₂ permeabilities (Table 2). As for N₂ diffusivities the subsequent order is attained: ([FAP]⁻ > [FSI]⁻ > [TFSAM]⁻ > [BETI]⁻ > [NTf₂]⁻ > [C₄F₉SO₃]⁻), which is nearly the same IL anion order observed for N₂ permeabilities, but different from that obtained for CO₂ diffusivities and permeabilities.

The relationship between IL viscosity and gas diffusion is in fact the basis of the dependence of permeability on viscosity that we have shown above. A number of works have reported the inversely proportional relationship between gas diffusivity and IL viscosity.^{3,16,17,41} Along this line of thought, the relationship between experimental CO₂ diffusivities and IL viscosity for the studied SILMs is depicted in Fig. 2.

In agreement to what was previously observed in the literature for other SILMs,^{4,21–23} the CO₂ diffusivity through SILMs having fluorinated anions decreases as the IL viscosity increases. The SILMs with the lowest CO₂ diffusivities are [C₂mim][C₄F₉SO₃] (55 × 10⁻¹² m² s⁻¹) and [C₂mim][BETI] (167 × 10⁻¹² m² s⁻¹), which also have the lowest CO₂ permeabilities (437 and 32 Barrer), depicting the highest viscosities (109.7 and 85.5 mPa). Similarly to what is mentioned above for CO₂ permeabilities, again a deviant behaviour can be observed for the [C₂mim][FAP] SILM, since its CO₂ diffusivity is in between those of [C₂mim][NTf₂] and [C₂mim][TFSAM] (Table 3), but its viscosity values (76.4 mPa s) are higher than those of [C₂mim][NTf₂] (39.1 mPa s) and [C₂mim][TFSAM] (23.7 mPa s) (Fig. 2).

The gas solubility (*S*) values calculated using Eqn 1 are listed in Table 4. It can be seen that the [C₂mim][C₄F₉SO₃] SILM presents the lowest CO₂ solubility (4 × 10⁻⁶ m_(STP)³ m⁻³ Pa⁻¹), while the previously reported [C₂mim][NTf₂] SILM has the highest CO₂ solubility (26 × 10⁻⁶ m_(STP)³ m⁻³ Pa⁻¹). The CO₂ solubility values of the remaining studied SILMs having different fluorinated anions are very similar (ranging from 20 × 10⁻⁶

Table 3 Gas diffusivity (*D*) values through the prepared SILMs of pure ILs

SILM sample	Gas diffusivity (×10 ¹²) (m ² s ⁻¹)	
	<i>D</i> (CO ₂)	<i>D</i> (N ₂)
[C ₂ mim][TFSAM]	258 ± 1.1	291 ± 4.5
[C ₂ mim][FSI]	315 ± 4.4	353 ± 7.5
[C ₂ mim][NTf ₂] ^a	172 ± 1.7	203 ± 2.5
[C ₂ mim][FAP]	214 ± 1.9	326 ± 0.2
[C ₂ mim][BETI]	167 ± 2.1	216 ± 2.4
[C ₂ mim][C ₄ F ₉ SO ₃]	55 ± 1.2	88 ± 2.0

^a The gas diffusivity values through the [C₂mim][NTf₂] SILM were taken from Tomé *et al.*²⁹



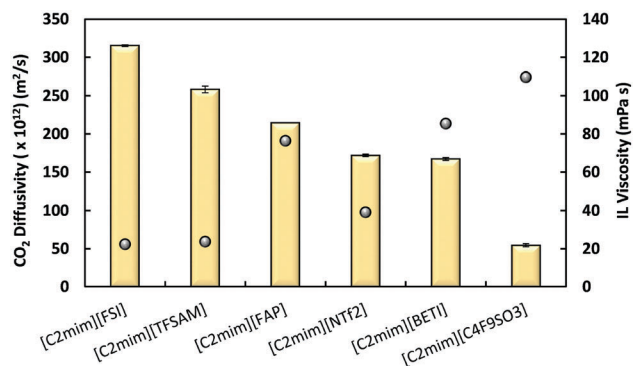


Fig. 2 Relationship between the CO₂ diffusivities determined through the SILMs prepared with pure ILs and the respective IL viscosities measured at 293 K. Error bars represent standard deviations based on three experimental replicas.

to $22 \times 10^{-6} \text{ m}_{(\text{STP})}^3 \text{ m}^{-3} \text{ Pa}^{-1}$) and can be ordered as follows: $[\text{TFSAM}]^- \approx [\text{FAP}]^- > [\text{FSI}]^- \approx [\text{BETI}]^-$. Regarding the N₂ solubility, its values are always significantly lower (between 0.39×10^{-6} and $0.61 \times 10^{-6} \text{ m}_{(\text{STP})}^3 \text{ m}^{-3} \text{ Pa}^{-1}$) than those of CO₂ for all the SILMs studied.

Over the past few years, a number of correlations have been proposed with the intention of understanding the relationships between CO₂ solubility and the intrinsic properties of ILs.^{42–44} The proposed models showed that CO₂ solubility increases with increasing IL molecular weight, molar volume and free volume.¹⁰ Taking into consideration the gas solubility results obtained in this work (Table 4), as well as the range of molar volumes (from 201.2 up to 325.6 cm³ mol⁻¹) and molecular weights (from 284.3 up to 556.2 g mol⁻¹) of the ILs used, divergences from the abovementioned trends can be found for the studied SILMs having fluorinated anions. For example, the [C₂mim][NTf₂] SILM shows the highest CO₂ solubility ($26 \times 10^{-6} \text{ m}_{(\text{STP})}^3 \text{ m}^{-3} \text{ Pa}^{-1}$), but it does not have the highest IL molar volume and molecular weight (Table 1). Likewise, the lowest CO₂ solubility ($4 \times 10^{-6} \text{ m}_{(\text{STP})}^3 \text{ m}^{-3} \text{ Pa}^{-1}$) belongs to the [C₂mim][C₄F₉SO₃] SILM, although it does not present the lowest IL molar volume and molecular weight (Table 1). The effect of fluorination, either in the IL cation or anion, on CO₂ solubility has been studied by different researchers.^{19,45–47} Tagiuri *et al.*⁴⁸ explored the effect of cation on the CO₂ solubility of three different ILs combining the [FSI]⁻ anion. Moreover, Kroon *et al.*⁴⁹ determined the CO₂ solubility in the [C₂mim][FAP] IL by measuring the bubble point

Table 4 Gas solubility (*S*) values through the prepared SILMs of pure ILs

SILM sample	Gas solubility ($\times 10^6$) ($\text{m}_{(\text{STP})}^3 \text{ m}^{-3} \text{ Pa}^{-1}$)	
	<i>S</i> (CO ₂)	<i>S</i> (N ₂)
[C ₂ mim][TFSAM]	22 ± 0.09	0.44 ± 0.01
[C ₂ mim][FSI]	20 ± 0.29	0.39 ± 0.01
[C ₂ mim][NTf ₂] ^a	26 ± 0.28	0.61 ± 0.01
[C ₂ mim][FAP]	22 ± 0.21	0.55 ± 0.001
[C ₂ mim][BETI]	20 ± 0.17	0.61 ± 0.01
[C ₂ mim][C ₄ F ₉ SO ₃]	4 ± 0.07	0.49 ± 0.002

^a The gas solubility values through the [C₂mim][NTf₂] SILM were taken from Tomé *et al.*²⁹

pressures of the binary mixture of [C₂mim][FAP] + CO₂. The results showed that the CO₂ solubility in [C₂mim][FAP] is higher when compared to that of ILs having the same cation combined with other fluorinated anions such as [NTf₂]⁻, [BF₄]⁻ and [PF₆]⁻, due to the fact that [FAP]⁻ has a large size and it is highly fluorinated. Although it has been recognized that introducing fluorination into the cation and/or anion can effectively improve CO₂ solubility,⁴⁶ it was recently reported after critical analysis that no special effect of the fluorination upon the CO₂ solubility has been observed for both perfluorocarbon and heavily fluorinated ILs.⁵⁰ In fact, the introduction of fluorination into the anions of the ILs studied in this work does not significantly affect the obtained gas solubility values (Table 4), except for the case of the [C₂mim][C₄F₉SO₃] IL that displays a very low CO₂ solubility.

Re-designing the [TFSAM]⁻ anion by structural unfolding: effect on gas permeation

Taking into account that the [TFSAM]⁻ anion has an unconventional and asymmetric chemical structure, combining both fluorinated and cyano functionalities, which have both been recognized to be responsible for high CO₂ separation performance, we explore here the effect of structural unfolding of the pure [C₂mim][TFSAM] IL on gas permeation properties of SILMs using IL mixtures. Thus, two equimolar IL mixtures were used for this purpose: [C₂mim][SCN][OTf] (Fig. 3), which is studied here for the first time, and [C₂mim][N(CN)₂][NTf₂], whose gas permeation and thermophysical properties were previously determined by us.²⁹ Both these mixtures have IL anions that show structural similarities to the [TFSAM]⁻ anion (Fig. 3). The composition description, water content (wt%), molar mass (*M*), viscosity (*η*), density (*ρ*) and molar volume (*V_m*) values of the pure ILs, [C₂mim][TFSAM], [C₂mim][SCN], [C₂mim][OTf], [C₂mim][N(CN)₂], [C₂mim][NTf₂], and the selected IL mixtures are listed in Table 5, while their gas permeability, diffusivity and solubility values are depicted in Fig. 4(a)–(c), respectively.

Fig. 4(a) and (b) show that the pure [C₂mim][TFSAM] SILM exhibits higher gas permeabilities and diffusivities than SILMs

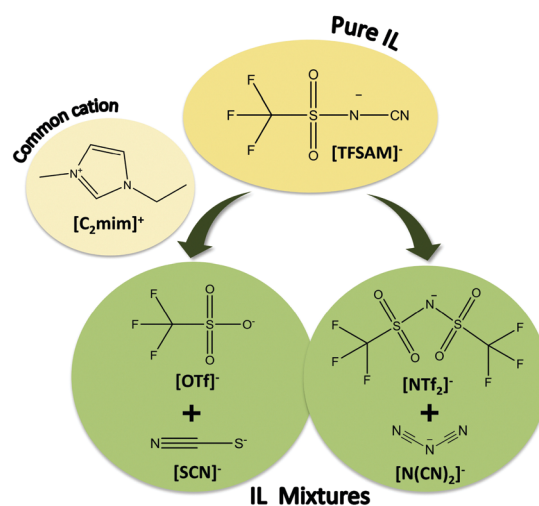


Fig. 3 Chemical structures of the pure IL and IL mixtures studied in this work.



Table 5 Composition descriptions, water contents and thermophysical properties of the pure ILs and IL mixtures

IL sample	Composition description (mole fraction)	H ₂ O (wt%)	<i>M</i> (g mol ⁻¹)	η^a (mPa s)	ρ^b (g cm ⁻³)	<i>V_m</i> (cm ³ mol ⁻¹)
[C ₂ mim][TFSAM]	Pure	0.02	284.26	23.7	1.3518	210.3
[C ₂ mim][SCN] ^c	Pure	0.09	169.25	27.9	1.1190	151.2
[C ₂ mim][SCN][OTf]	0.5 [C ₂ mim][SCN] + 0.5 [C ₂ mim][OTf]	0.16	214.74	36.4	1.2638	169.9
[C ₂ mim][OTf]	Pure	0.10	260.23	49.5	1.3850	187.9
[C ₂ mim][N(CN) ₂] ^c	Pure	0.09	177.21	18.0	1.1060	160.2
[C ₂ mim][N(CN) ₂][NTf ₂] ^c	0.5[C ₂ mim][N(CN) ₂] + 0.5 [C ₂ mim][NTf ₂]	0.12	284.26	29.2	1.3620	208.7
[C ₂ mim][NTf ₂] ^c	Pure	0.02	391.31	39.1	1.5240	256.8

^a Viscosity (η) and density (ρ) measured at 293 K. ^b Molar volume (*V_m*) obtained at 293 K. ^c The values of [C₂mim][N(CN)₂][NTf₂] were taken from Tomé *et al.*²⁹

composed of both [C₂mim][SCN][OTf] and [C₂mim][N(CN)₂][NTf₂] IL mixtures. Moreover, the addition of 0.5 mole fraction of [C₂mim][SCN] did not significantly affect CO₂ permeability but decreased to almost half the N₂ permeability compared to those of the pure [C₂mim][OTf] SILM. From Fig. 4(b), where the relationship between CO₂ diffusivity and IL viscosity is illustrated, it can also be seen that both [C₂mim][N(CN)₂][NTf₂] and [C₂mim][SCN][OTf] IL mixtures present slightly higher viscosities (Table 5) and lower CO₂ diffusivities than those of the pure [C₂mim][TFSAM] SILM. Also, the presence of 0.5 mole fraction of [C₂mim][SCN] leads to a decrease in IL viscosity and an increase in CO₂ diffusivity. The same behaviour was found for [C₂mim][N(CN)₂][NTf₂] with the addition of 0.5 mole fraction of [C₂mim][N(CN)₂]. These results are in accordance with the general trend observed in the literature: the CO₂ diffusivity decreases with the increase in IL viscosity.^{4,21–23}

Concerning gas solubility, from Fig. 4(c), it can be observed that CO₂ solubilities of both IL mixtures are in between those of the individual IL components. Moreover, the presence of 0.5 mole fraction of [C₂mim][OTf] or [C₂mim][NTf₂] in the corresponding mixtures leads to an increase in the CO₂ solubilities, probably due to the fact that they present higher molar volume compared to the pure [C₂mim][SCN] or [C₂mim][N(CN)₂], respectively. Furthermore, the [C₂mim][N(CN)₂][NTf₂] mixture displays a CO₂ solubility ($20 \times 10^{-6} \text{ m}_{(\text{STP})}^3 \text{ m}^{-3} \text{ Pa}^{-1}$) closer to that of the pure [C₂mim][TFSAM] SILM ($22 \times 10^{-6} \text{ m}_{(\text{STP})}^3 \text{ m}^{-3} \text{ Pa}^{-1}$), probably due to the fact that these IL phases have very close molar volumes (208.7 and 210.3 cm³ mol⁻¹, respectively) and the same molecular weight (284.3 g mol⁻¹). Conversely, the [C₂mim][SCN][OTf] mixture has lower CO₂ solubility ($15 \times 10^{-6} \text{ m}_{(\text{STP})}^3 \text{ m}^{-3} \text{ Pa}^{-1}$) than that observed for the pure [C₂mim][TFSAM] SILM, which is in agreement with the lower molar volume (169.9 cm³ mol⁻¹) and molecular weight (214.7 cm³ mol⁻¹) presented by this IL mixture containing the [SCN]⁻ and [OTf]⁻ anions. Despite the fact that it is possible to obtain similar viscosities and molar volumes of the pure [C₂mim][TFSAM] IL by mixing the [C₂mim][N(CN)₂] and [C₂mim][NTf₂] ILs, improved gas permeabilities, diffusivities and solubilities were obtained through the pure [C₂mim][TFSAM] SILM.

Comparison of CO₂/N₂ separation performance

The gas permeabilities and CO₂/N₂ permselectivities of all the studied SILMs are listed in Table 6. Amongst the pure SILMs with fluorinated anions, the [C₂mim][FSI] and [C₂mim][TFSAM] SILMs not only show the highest gas permeabilities (843 and

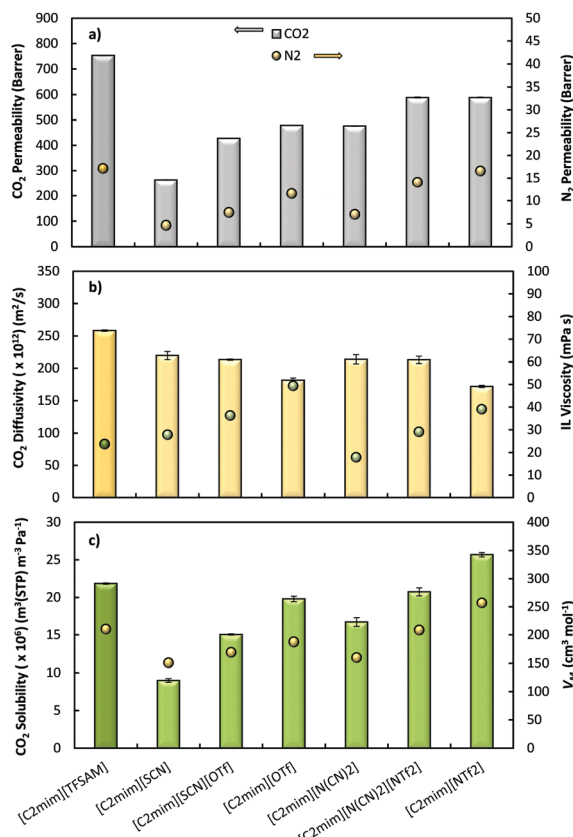


Fig. 4 (a) Gas permeability, (b) CO₂ diffusivity as a function of IL viscosity and (c) CO₂ solubility as a function of IL molar volume for the studied SILMs. Error bars represent standard deviations based on three experimental replicas. The gas permeability, diffusivity and solubility values of [C₂mim][N(CN)₂][NTf₂] and [C₂mim][NTf₂] were taken from Tomé *et al.*²⁹

753 Barrer, respectively), but also have the largest CO₂/N₂ permselectivities (43.9 and 46.1, respectively). In contrast, the lowest gas permeabilities and CO₂/N₂ permselectivity belong to the [C₂mim][C₄F₉SO₃] SILM. The CO₂/N₂ permselectivities of the pure SILMs decrease as the fluorinated chain increases in the IL anion: [FSI]⁻ > [TFSAM]⁻ > [NTf₂]⁻ > [BETI]⁻, the [C₂mim][FAP] and [C₂mim][C₄F₉SO₃] SILMs being the only exceptions.

Concerning the effect of the structural unfolding of the [TFSAM]⁻ anion, and as previously discussed, the pure [C₂mim][TFSAM] SILM presents higher gas permeabilities compared to its structurally similar IL mixtures (Table 6). Nevertheless, the greatest CO₂/N₂



Table 6 Single gas permeability (P)^a and ideal permselectivities (α) of all the studied SILMs^b

SILM sample	P (CO ₂)	P (N ₂)	α (CO ₂ /N ₂)
[C ₂ mim][TFSAM]	753 ± 0.2	17 ± 0.1	43.9 ± 0.1
[C ₂ mim][FSI]	843 ± 0.5	18 ± 0.2	46.1 ± 0.5
[C ₂ mim][FAP]	624 ± 0.4	24 ± 0.1	26.0 ± 0.1
[C ₂ mim][BETI]	437 ± 1.9	18 ± 0.1	24.8 ± 0.2
[C ₂ mim][C ₄ F ₉ SO ₃]	32 ± 0.2	6 ± 0.1	5.5 ± 0.1
[C ₂ mim][SCN] ^c	263 ± 0.6	5 ± 0.2	56.6 ± 1.9
[C ₂ mim][SCN][OTf]	428 ± 0.5	7 ± 0.1	57.6 ± 0.4
[C ₂ mim][OTf]	479 ± 0.6	12 ± 0.1	41.4 ± 0.3
[C ₂ mim][N(CN) ₂] ^c	476 ± 0.8	7 ± 0.1	67.8 ± 0.6
[C ₂ mim][N(CN) ₂][NTf ₂] ^c	589 ± 1.9	14 ± 0.2	41.8 ± 0.7
[C ₂ mim][NTf ₂] ^c	589 ± 1.0	17 ± 0.1	35.5 ± 0.3

^a Barrer (1 Barrer = 10⁻¹⁰ cm_(STP)³ cm cm⁻² s⁻¹ cmHg⁻¹). ^b The listed uncertainties represent the standard deviations, based on three experiments. ^c Values taken from Tomé *et al.*²⁹

permselectivity (57.6) was achieved for the SILM containing the [C₂mim][SCN][OTf] IL mixture.

With the purpose of comparing the performance results obtained in this work to those reported in the literature for other SILMs, Fig. 5 displays the Robeson plot for CO₂/N₂ separation, where the CO₂/N₂ permselectivity is plotted against CO₂ permeability and the solid black line represents the empirical 2008 upper bound for this gas pair.⁵¹ It can be seen that among the SILMs immobilized with the pure IL having fluorinated anions, both the [C₂mim][TFSAM] and [C₂mim][FSI] SILMs fall or exceed the Robeson 2008 upper bound, meaning that these two ILs are the most promising candidates for CO₂/N₂ separation processes. Comparing the results of the pure [C₂mim][TFSAM] SILM with those of selected IL mixtures, Fig. 5 clearly shows that the CO₂/N₂ separation performance of the SILM immobilized with the [C₂mim][SCN][OTf] IL mixture surpasses the upper bound, primarily due to its higher CO₂/N₂ permselectivity (57.6). On the other hand, the CO₂/N₂ separation efficiency of the [C₂mim][N(CN)₂][NTf₂] SILM is on top of the

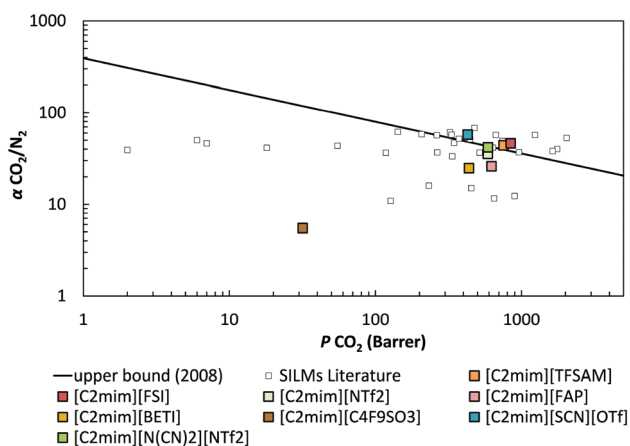


Fig. 5 CO₂/N₂ separation performance of all the studied SILMs. The experimental error is within the data points. Data are plotted on a log–log scale and the upper bound is adapted from Robeson.⁵¹ The literature data previously reported for other SILMs (□)^{15,17,22,24,29,30,32,53} are also illustrated for comparison. The values of [C₂mim][N(CN)₂][NTf₂] and [C₂mim][NTf₂] SILMs were taken from Tomé *et al.*²⁹

upper bound, since it presents lower permselectivity (41.8), despite its high CO₂ permeability (589 Barrer) in comparison to that of the [C₂mim][SCN][OTf] SILM (428 Barrer). Actually, it is the [C₂mim][N(CN)₂][NTf₂] SILM that discloses the most similar CO₂/N₂ separation performance results to the pure synthesized [C₂mim][TFSAM] IL (Fig. 5).

Experimental

Materials

Lithium bis(pentafluoroethylsulfanyl)imide (Li(CF₃CF₂SO₂)₂N, LiBETI, 98%, Chameleon Reagent) and lithium nonafluoro-1-butananesulfonate (LiC₄F₉SO₃, > 95%, TCI Chemicals) were used without purification. Reagent-grade dichloromethane, acetonitrile, hexane and ethyl acetate were obtained from Aldrich or Merck and were dried by vacuum distillation over P₂O₅. *N*-Methylimidazole (98%, Aldrich) and bromoethane (98%, Acros) were distilled under an inert atmosphere over CaH₂.

1-Ethyl-3-methylimidazolium bis(fluorosulfanyl)imide ([C₂mim][FSI], 99.5 wt%, Solvionic), 1-ethyl-3-methylimidazolium tris(pentafluoro-ethyl)trifluorophosphate ([C₂mim][FAP], 98 wt%, Merck), 1-ethyl-3-methylimidazolium thiocyanate ([C₂mim][SCN], > 98 wt%, IoLiTec) and 1-ethyl-3-methylimidazolium trifluoromethanesulfonate ([C₂mim][OTf], ≥ 98 wt%, Aldrich) were obtained from the specified suppliers. To reduce the content of water and other volatile substances, the pure ILs were dried at approximately 1 Pa and 318 K for at least 4 days.

IL synthesis and characterization

1-Ethyl-3-methylimidazolium bromide ([C₂mim][Br]). [C₂mim][Br] was synthesized by the reaction between *N*-methylimidazole and an excess of bromoethane following the method used in a previous work.³⁴ Spectroscopic data of the target compound were in accordance with those reported in the literature.

1-Ethyl-3-methylimidazolium 2,2,2-trifluoromethylsulfonyl-*N*-cyanoamide ([C₂mim][TFSAM]). [C₂mim][TFSAM] was prepared by ion exchange between [C₂mim][Br] and KTFSAM, in an aqueous medium in accordance with a described procedure.³⁵ Yield: 85%; anal. calcd for C₈H₁₁N₄F₃SO₂ (284.26), %: N, 19.71%; C, 33.80%; H, 3.90%; Found, %: N, 19.68%; C, 33.78%; H, 3.99%; ¹H NMR (300 MHz, DMSO-*d*₆): 9.08 (s, 1H, 2H (Im)), 7.72 (s, 1H, H4 (Im)), 7.64 (s, 1H, H5 (Im)), 4.22–4.15 (m, 2H, CH₂CH₃), 3.84 (s, 3H, CH₃), 1.42 (t, 3H, CH₂CH₃, *J*_{HH} = 7.5 Hz); ¹³C NMR (100.6 MHz, DMSO-*d*₆): 136.1, 125.0–115.4 (q, ¹*J*_{CF} = 325 Hz), 123.4, 121.8, 44.1, 35.5, 14.8; ¹⁹F NMR (282.4 MHz, DMSO-*d*₆): –77.8; IR (KBr pellet): 3158 (m, ν_{C–H}), 3117 (m, ν_{C–H}), 2192 (vs, ν_{C–N}), 1573 (m), 1468 (w), 1333 (vs, ν_{asSO₂}), 1236 (s), 1217 (vs, ν_{CF}), 1170 (vs, ν_{ssSO₂}), 1119 (s, ν_{CF}), 832 (s), 752 (w), 639 (m), 595 (s), 479 (m) cm⁻¹.

1-Ethyl-3-methylimidazolium nonafluorobutanesulfonate ([C₂mim][C₄F₉SO₃]). Lithium nonafluoro-1-butananesulfonate (8.00 g, 0.026 mol) was dissolved in 20 mL of distilled water and added dropwise to a solution of [C₂mim][Br] (3.84 g, 0.020 mol) in 15 mL of H₂O at ambient temperature. The solution was stirred for 2 h at room temperature and then [C₂mim][C₄F₉SO₃] was extracted



with dichloromethane (4 × 40 mL). The combined CH₂Cl₂ solution was washed with a small amount of water and dried over anhydrous MgSO₄. The magnesium sulfate was filtered off and dichloromethane was stripped off under reduced pressure. The product was obtained as slightly yellow transparent fluid oil, which was finally dried at 323 K and 100 Pa for 12 h using a special flask filled with P₂O₅ and introduced into the vacuum line. Yield: 5.61 g (68%); anal. calcd for C₁₀H₁₁N₂F₉SO₃ (410.26), %: C, 29.28%; H, 2.70%; F, 41.68%; Found, %: C, 28.99%; H, 2.83%; F, 41.39%; ¹H NMR (300 MHz, DMSO-d₆): 9.09 (s, 1H, H2 (Im)), 7.77 (s, 1H, H4 (Im)), 7.68 (s, 1H, H5 (Im)), 4.22–4.17 (m, 2H, CH₂CH₃), 3.85 (s, 3H, CH₃), 1.43–1.39 (m, 3H, CH₂CH₃); ¹³C NMR (100.6 MHz, DMSO-d₆): 136.2, 123.5, 121.9, 117.1 (qt, -CF₂-CF₃, ¹J = 288 Hz, ²J = 34 Hz), 113.4 (tt, (-)O₃S-CF₂-, ¹J = 288 Hz, ²J = 34 Hz), 110.4 (tp, (-)O₃S-CF₂-CF₂-, ¹J = 266, ²J = 33 Hz), 109.8 (qq, -CF₂-CF₃, ¹J = 268 Hz, ²J = 38 Hz), 44.1, 35.6, 14.9; ¹⁹F NMR (282.4 MHz, DMSO-d₆): -80.8, -114.9, -121.6, -125.9; IR (KBr pellet): 3156 (s, ν_{C-H}), 3117 (s, ν_{C-H}), 2993 (m, ν_{C-H}), 1574 (s), 1462 (m), 1432 (w), 1393 (w), 1353 (s), 1261 (vs, ν_{asSO₂}), 1236 (vs), 1214 (vs, ν_{CF}), 1170 (vs, ν_{ssO₂}), 1134 (s, ν_{CF}), 1057 (vs), 1119 (m), 1006 (m), 988 (w), 870 (m), 845 (m), 802 (m), 736 (m), 699 (m), 679 (w), 656 (s), 620 (s), 596 (m), 564 (m), 532 (s) cm⁻¹.

1-Ethyl-3-methylimidazolium bis(pentafluoroethylsulfonyl)imide ([C₂mim][BETI]). The procedure previously described for [C₂mim][C₄F₉SO₃] was also used for the synthesis of [C₂mim]-[BETI]. After purification and drying, [C₂mim][BETI] was obtained as colorless transparent fluid liquid. Yield: 82%; anal. calcd for C₁₀H₁₁N₃F₁₀S₂O₄ (491.32), %: C, 24.45%; H, 2.26%; N, 8.55%; Found, %: C, 24.50%; H, 2.03%; N, 8.49%; ¹H NMR (300 MHz, DMSO-d₆): 9.11 (s, 1H, H2 (Im)), 7.76 (s, 1H, H4 (Im)), 7.68 (s, 1H, H5 (Im)), 4.22–4.16 (m, 2H, CH₂CH₃), 3.84 (s, 3H, CH₃), 1.43–1.40 (m, 3H, CH₂CH₃); ¹³C NMR (100.6 MHz, DMSO-d₆): 136.7, 123.9, 122.4, 117.7 (qt, -CF₂-CF₃, ¹J = 287 Hz, ²J = 34 Hz), 110.1 (tq, -CF₂-CF₃, ¹J = 293 Hz, ²J = 37 Hz), 44.6, 36.0, 15.4; ¹⁹F NMR (282.4 MHz, DMSO-d₆): -78.8, -117.6; IR (KBr pellet): 3160 (m, ν_{C-H}), 3124 (m, ν_{C-H}), 2993 (w, ν_{C-H}), 1574 (m), 1472 (w), 1432 (w), 1355 (vs, ν_{asSO₂}), 1331 (vs, ν_{CF}), 1223 (vs, ν_{CF}), 1172 (vs, ν_{ssO₂}), 1087 (s, ν_{CF}), 978 (s), 824 (w), 775 (w), 755 (m), 742 (m), 701 (w), 644 (m), 616 (s), 536 (m), 525 (m) cm⁻¹.

NMR spectra were recorded on an AMX-400 spectrometer (Bruker) at 298 K in the indicated deuterated solvent and are listed in ppm. The signal corresponding to the residual protons of the deuterated solvent was used as an internal standard for ¹H and ¹³C NMR, while for ¹⁹F NMR CHCl₂F was used as an external standard. IR spectra were acquired on a Nicolet Magna-750 Fourier IR-spectrometer using KBr pellets (128 scans, resolution was 2 cm⁻¹).

IL mixture preparation. The IL mixture, [C₂mim][SCN][OTf], containing 0.5 mole fraction of both [C₂mim][OTf] and [C₂mim][SCN], was prepared using an analytical high precision balance with an uncertainty of ±10⁻⁵ g by syringing known masses of the IL components into a glass vial. Good mixing was ensured by magnetic stirring for 30 min at 298 K. Then, the IL mixture was dried at roughly 1 Pa and 318 K for at least 4 days immediately prior to use. The water contents of all IL samples were determined by Karl Fischer titration using a 831 KF Coulometer (Metrohm).

Density and viscosity determination. The density and viscosity measurements of the pure ILs and the [C₂mim][SCN][OTf] IL mixture were performed at 293 K and atmospheric pressure using an SVM 3000 Anton Paar rotational Stabinger viscometer-densimeter, where the standard uncertainty for the temperature was 0.02 K. The repeatability of density and dynamic viscosity of this equipment was 0.0005 g cm⁻³ and 0.35%, respectively. Measurements of each sample were performed in triplicate to ensure accuracy and the reported results are average values. The highest relative standard uncertainty registered for the density and dynamic viscosity measurements was 1 × 10⁻⁴ and 0.03, respectively.

Gas permeation measurements. Porous hydrophobic poly(vinylidene fluoride) (PVDF) membranes supplied by Millipore Corporation (USA), with a pore size of 0.22 μm and an average thickness of 125 μm, were used to support [C₂mim][FSI], [C₂mim][FAP] and [C₂mim][BETI]. Since the impregnation of the remaining IL samples into hydrophobic PVDF resulted in unstable SILMs, the other IL samples were supported on porous hydrophilic poly(tetrafluoroethylene) (PTFE) membranes acquired from Merck Millipore, with a pore size of 0.2 μm and an average thickness of 65 μm. All the SILM configurations were prepared by the vacuum method.²⁹

Ideal gas permeabilities and diffusivities through the prepared SILMs were measured using a time-lag apparatus.³⁶ First, each SILM was degassed under vacuum inside the permeation cell for 12 h. Then, CO₂ and N₂ permeation experiments were carried out at 293 K with a *trans*-membrane pressure differential of 100 kPa. All the permeation data were measured at least in triplicate on a single SILM sample. The highest relative standard uncertainty registered for gas permeability measurements was 0.03. The permeation cell and lines were evacuated until the pressure was below 0.1 kPa before each run. No residual IL was found inside the permeation cell at the end of the experiments. The thickness of the SILMs was assumed to be equivalent to the membrane filter thickness.

Gas transport through the prepared SILMs was assumed to follow a solution-diffusion mass transfer mechanism,³⁷ where the permeability (*P*) is related to diffusivity (*D*) and solubility (*S*) as follows:

$$P = D \times S \quad (1)$$

The permeate flux of each gas (*J_i*) was determined experimentally using eqn (2),³⁸ where *V^p* is the permeate volume, Δ*p_d* is the variation of downstream pressure, *A* is the effective membrane surface area, *t* is the experimental time, *R* is the gas constant and *T* is the temperature.

$$J_i = \frac{V^p \Delta p_d}{A t R T} \quad (2)$$

Ideal gas permeability (*P_i*) was then determined from the steady-state gas flux (*J_i*), the membrane thickness (*ℓ*) and the trans-membrane pressure difference (Δ*p_i*), as shown in eqn (3).³⁸

$$P_i = \frac{J_i}{\Delta p_i / \ell} \quad (3)$$



Gas diffusivity (D_i) was determined according to eqn (4). The time-lag parameter (θ) was calculated by extrapolating the slope of the linear portion of the p_d vs. t curve back to the time axis, where the intercept was equal to θ .³⁹

$$D_i = \frac{\ell^2}{6\theta} \quad (4)$$

After P_i and D_i were known, the gas solubility (S_i) was calculated using the relationship shown in eqn (1). The ideal permeability selectivity (or permselectivity), α_{ij} , was obtained by dividing the permeability of the more permeable species i to the permeability of the less permeable species j . The permselectivity can also be expressed as the product of the diffusivity selectivity and the solubility selectivity:

$$\alpha_{ij} = \frac{P_i}{P_j} = \left(\frac{D_i}{D_j}\right) \times \left(\frac{S_i}{S_j}\right) \quad (5)$$

Conclusions

In this work, ILs containing a common cation ($[\text{C}_2\text{mim}]^+$) and different fluorinated anions ($[\text{TFSAM}]^-$, $[\text{FSI}]^-$, $[\text{C}_4\text{F}_9\text{SO}_3]^-$, $[\text{BETI}]^-$, $[\text{FAP}]^-$) were synthesized and used as liquid phases to prepare SILMs for flue gas separation (CO_2/N_2). The single CO_2 and N_2 permeation properties through the prepared SILMs were determined. The viscosity and density of the IL phases were also evaluated. The results showed that CO_2 permeabilities and diffusivities through the studied SILMs follow the same fluorinated anion order: $[\text{FSI}]^- > [\text{TFSAM}]^- > [\text{FAP}]^- > [\text{NTf}_2]^- > [\text{BETI}]^- > [\text{C}_4\text{F}_9\text{SO}_3]^-$, which is inversely related to IL viscosity, with the only outlier being $[\text{C}_2\text{mim}][\text{FAP}]$. Conversely, the introduction of fluorination in the IL anions did not significantly affect gas solubility, except for the case of the $[\text{C}_2\text{mim}][\text{C}_4\text{F}_9\text{SO}_3]$ SILM displaying a very low CO_2 solubility. Among the pure SILMs, it is worth noting that the best separation performances were achieved for $[\text{C}_2\text{mim}][\text{TFSAM}]$ and $[\text{C}_2\text{mim}][\text{FSI}]$ SILMs that fall on top of or surpassed the Robeson 2008 upper bound, with CO_2 permeabilities of 753 and 843 Barrer and CO_2/N_2 permselectivities of 43.9 and 46.1, respectively.

Furthermore, the effect of structural unfolding of the $[\text{TFSAM}]^-$ anion on gas permeation properties of SILMs was investigated using IL mixtures comprising both fluorinated and cyano functionalities in the anions. The pure $[\text{C}_2\text{mim}][\text{TFSAM}]$ IL provided a membrane with improved CO_2 permeabilities, diffusivities and solubilities compared to those of the SILMs based on the selected $[\text{C}_2\text{mim}][\text{SCN}][\text{OTf}]$ and $[\text{C}_2\text{mim}][\text{N}(\text{CN})_2][\text{NTf}_2]$ IL mixtures. Overall, and despite the fact that the $[\text{C}_2\text{mim}][\text{SCN}][\text{OTf}]$ SILM revealed better CO_2/N_2 separation performance essentially due to its higher CO_2/N_2 permselectivity (57.6), the $[\text{C}_2\text{mim}][\text{N}(\text{CN})_2][\text{NTf}_2]$ IL mixture disclosed the most similar results to the pure synthesized $[\text{C}_2\text{mim}][\text{TFSAM}]$ IL, not only in terms of thermophysical properties, but also regarding gas transport and CO_2/N_2 separation performance.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

Andreia S. L. Gouveia and Liliana C. Tomé are grateful to FCT (Fundação para a Ciência e a Tecnologia) for their Doctoral (SFRH/BD/116600/2016) and Post-doctoral research grants (SFRH/BPD/101793/2014), respectively, Isabel M. Marrucho acknowledges FCT/MCTES (Portugal) for a contract under Investigador FCT 2012. This work was partially supported by FCT through the project PTDC/CTM-POL/2676/2014, R&D unit UID/Multi/04551/2013 (GreenIT), R&D unit UID/QUI/00100/2013 (CQE) and through the grant of President of the Russian Federation "For Young Outstanding Professors" (project no. MD-2371.2017.3).

Notes and references

- L. C. Tomé and I. M. Marrucho, *Chem. Soc. Rev.*, 2016, **45**, 2785–2824.
- Z. Dai, R. D. Noble, D. L. Gin, X. Zhang and L. Deng, *J. Membr. Sci.*, 2016, **497**, 1–20.
- P. Scovazzo, *J. Membr. Sci.*, 2009, **343**, 199–211.
- P. K. Parhi, *J. Chem.*, 2013, 11.
- L. J. Lozano, C. Godínez, A. P. de los Ríos, F. J. Hernández-Fernández, S. Sánchez-Segado and F. J. Alguacil, *J. Membr. Sci.*, 2011, **376**, 1–14.
- P. Luis, T. Van Gerven and B. Van der Bruggen, *Prog. Energy Combust. Sci.*, 2012, **38**, 419–448.
- M. J. Earle, J. M. S. S. Esperanca, M. A. Gilea, J. N. Canongia Lopes, L. P. N. Rebelo, J. W. Magee, K. R. Seddon and J. A. Widegren, *Nature*, 2006, **439**, 831–834.
- C. Cadena, J. L. Anthony, J. K. Shah, T. I. Morrow, J. F. Brennecke and E. J. Maginn, *J. Am. Chem. Soc.*, 2004, **126**, 5300–5308.
- M. Hasib-ur-Rahman, M. Sijaj and F. Larachi, *Chem. Eng. Process.*, 2010, **49**, 313–322.
- M. Ramdin, T. W. de Loos and T. J. H. Vlugt, *Ind. Eng. Chem. Res.*, 2012, **51**, 8149–8177.
- L. A. Neves, J. G. Crespo and I. M. Coelho, *J. Membr. Sci.*, 2010, **357**, 160–170.
- P. C. Hillesheim, J. A. Singh, S. M. Mahurin, P. F. Fulvio, Y. Oyola, X. Zhu, D.-e. Jiang and S. Dai, *RSC Adv.*, 2013, **3**, 3981–3989.
- P. C. Hillesheim, S. M. Mahurin, P. F. Fulvio, J. S. Yeary, Y. Oyola, D.-e. Jiang and S. Dai, *Ind. Eng. Chem. Res.*, 2012, **51**, 11530–11537.
- S. M. Mahurin, T. Dai, J. S. Yeary, H. Luo and S. Dai, *Ind. Eng. Chem. Res.*, 2011, **50**, 14061–14069.
- L. C. Tomé, D. J. S. Patinha, R. Ferreira, H. Garcia, C. Silva Pereira, C. S. R. Freire, L. P. N. Rebelo and I. M. Marrucho, *ChemSusChem*, 2014, **7**, 110–113.
- R. Condemarin and P. Scovazzo, *Chem. Eng. J.*, 2009, **147**, 51–57.
- L. Ferguson and P. Scovazzo, *Ind. Eng. Chem. Res.*, 2007, **46**, 1369–1374.
- J. J. Close, K. Farmer, S. S. Moganty and R. E. Baltus, *J. Membr. Sci.*, 2012, **390–391**, 201–210.
- J. E. Bara, C. J. Gabriel, T. K. Carlisle, D. E. Camper, A. Finotello, D. L. Gin and R. D. Noble, *Chem. Eng. J.*, 2009, **147**, 43–50.



- 20 P. Cserjési, N. Nemestóthy, A. Vass, Z. Csanádi and K. Bélafi-Bakó, *Desalination*, 2009, **245**, 743–747.
- 21 C. Myers, H. Pennline, D. Luebke, J. Ilconich, J. K. Dixon, E. J. Maginn and J. F. Brennecke, *J. Membr. Sci.*, 2008, **322**, 28–31.
- 22 P. Scovazzo, D. Havard, M. McShea, S. Mixon and D. Morgan, *J. Membr. Sci.*, 2009, **327**, 41–48.
- 23 S. M. Mahurin, J. S. Lee, G. A. Baker, H. Luo and S. Dai, *J. Membr. Sci.*, 2010, **353**, 177–183.
- 24 S. M. Mahurin, P. C. Hillesheim, J. S. Yeary, D.-e. Jiang and S. Dai, *RSC Adv.*, 2012, **2**, 11813–11819.
- 25 S. M. Mahurin, J. S. Yeary, S. N. Baker, D.-e. Jiang, S. Dai and G. A. Baker, *J. Membr. Sci.*, 2012, **401–402**, 61–67.
- 26 E. Kamio, T. Matsuki, S. Kasahara and H. Matsuyama, *Sep. Sci. Technol.*, 2017, **52**, 209–220.
- 27 S. Kasahara, E. Kamio, T. Ishigami and H. Matsuyama, *J. Membr. Sci.*, 2012, **415–416**, 168–175.
- 28 S. Kasahara, E. Kamio, T. Ishigami and H. Matsuyama, *Chem. Commun.*, 2012, **48**, 6903–6905.
- 29 L. C. Tomé, D. J. S. Patinha, C. S. R. Freire, L. P. N. Rebelo and I. M. Marrucho, *RSC Adv.*, 2013, **3**, 12220–12229.
- 30 L. C. Tomé, C. Florindo, C. S. R. Freire, L. P. N. Rebelo and I. M. Marrucho, *Phys. Chem. Chem. Phys.*, 2014, **16**, 17172–17182.
- 31 A. S. L. Gouveia, L. C. Tomé and I. M. Marrucho, *J. Membr. Sci.*, 2016, **510**, 174–181.
- 32 A. B. Pereira, L. C. Tomé, S. Martinho, L. P. N. Rebelo and I. M. Marrucho, *Ind. Eng. Chem. Res.*, 2013, **52**, 4994–5001.
- 33 M. Althuluth, J. P. Overbeek, H. J. van Wees, L. F. Zubeir, W. G. Haije, A. Berrouk, C. J. Peters and M. C. Kroon, *J. Membr. Sci.*, 2015, **484**, 80–86.
- 34 Y. U. Paulechka, G. J. Kabo, A. V. Blokhin, A. S. Shaplov, E. I. Lozinskaya and Y. S. Vygodskii, *J. Chem. Thermodyn.*, 2007, **39**, 158–166.
- 35 A. S. Shaplov, E. I. Lozinskaya, P. S. Vlasov, S. M. Morozova, D. Y. Antonov, P. H. Aubert, M. Armand and Y. S. Vygodskii, *Electrochim. Acta*, 2015, **175**, 254–260.
- 36 L. C. Tomé, D. Mecerreyes, C. S. R. Freire, L. P. N. Rebelo and I. M. Marrucho, *J. Membr. Sci.*, 2013, **428**, 260–266.
- 37 J. G. Wijmans and R. W. Baker, *J. Membr. Sci.*, 1995, **107**, 1–21.
- 38 S. Matteucci, Y. Yampolskii, B. D. Freeman and I. Pinnau, *Materials Science of Membranes for Gas and Vapor Separation*, John Wiley & Sons, Ltd, 2006, pp. 1–47.
- 39 S. W. Rutherford and D. D. Do, *Adsorption*, 1997, **3**, 283–312.
- 40 E. Santos, J. Albo, C. I. Daniel, C. A. M. Portugal, J. G. Crespo and A. Irabien, *J. Membr. Sci.*, 2013, **430**, 56–61.
- 41 D. Morgan, L. Ferguson and P. Scovazzo, *Ind. Eng. Chem. Res.*, 2005, **44**, 4815–4823.
- 42 D. Camper, J. Bara, C. Koval and R. Noble, *Ind. Eng. Chem. Res.*, 2006, **45**, 6279–6283.
- 43 P. J. Carvalho and J. A. P. Coutinho, *J. Phys. Chem. Lett.*, 2010, **1**, 774–780.
- 44 M. S. Shannon, J. M. Tedstone, S. P. O. Danielsen, M. S. Hindman, A. C. Irvin and J. E. Bara, *Ind. Eng. Chem. Res.*, 2012, **51**, 5565–5576.
- 45 J. L. Anderson, J. K. Dixon and J. F. Brennecke, *Acc. Chem. Res.*, 2007, **40**, 1208–1216.
- 46 M. J. Muldoon, S. N. V. K. Aki, J. L. Anderson, J. K. Dixon and J. F. Brennecke, *J. Phys. Chem. B*, 2007, **111**, 9001–9009.
- 47 Y.-F. Hu, Z.-C. Liu, C.-M. Xu and X.-M. Zhang, *Chem. Soc. Rev.*, 2011, **40**, 3802–3823.
- 48 A. Tagiuri, K. Z. Sumon, A. Henni, K. Zanganeh and A. Shafeen, *Fluid Phase Equilib.*, 2014, **375**, 324–331.
- 49 M. Althuluth, M. T. Mota-Martinez, M. C. Kroon and C. J. Peters, *J. Chem. Eng. Data*, 2012, **57**, 3422–3425.
- 50 P. J. Carvalho, K. A. Kurnia and J. A. P. Coutinho, *Phys. Chem. Chem. Phys.*, 2016, **18**, 14757–14771.
- 51 L. M. Robeson, *J. Membr. Sci.*, 2008, **320**, 390–400.
- 52 C. M. S. S. Neves, K. A. Kurnia, K. Shimizu, I. M. Marrucho, L. P. N. Rebelo, J. A. P. Coutinho, M. G. Freire and J. N. Canongia Lopes, *Phys. Chem. Chem. Phys.*, 2014, **16**, 21340–21348.
- 53 P. Cserjési, N. Nemestóthy and K. Bélafi-Bakó, *J. Membr. Sci.*, 2010, **349**, 6–11.

