Reactions of Ions with Ionic Liquid Vapors by Selected-Ion Flow Tube Mass Spectrometry (Preprint)

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Room temperature ionic liquids exert vanishingly small vapor pressures under ambient conditions. Under reduced pressure, certain ionic liquids have demonstrated volatility and they are thought to vaporize as intact cation-anion ion pairs. However, ion pair vapors are difficult to detect because their concentration is extremely low under these conditions. In this letter, we report the products of reacting ions such as NO+, NH4+, NO3-, and O2- with vaporized aprotic ionic liquids in their intact ion pair form. Ion pair fragmentation to the cation or anion, as well as ion exchange and ion addition processes are observed by selected ion flow tube mass spectrometry. Free energies of the reactions involving 1-ethyl-3-methylimidazolium bistrifluoromethylsulfonylimide determined by ab initio quantum mechanical calculations indicate that ion exchange or ion addition are energetically more favorable than charge transfer processes, whereas charge transfer processes can be important in reactions involving 1-butyl-3-methylimidazolium dicyanamide.
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Abstract

Room temperature ionic liquids exert vanishingly small vapor pressures under ambient conditions. Under reduced pressure, certain ionic liquids have demonstrated volatility and they are thought to vaporize as intact cation-anion ion pairs. However, ion pair vapors are difficult to detect because their concentration is extremely low under these conditions. In this letter, we report the products of reacting ions such as NO$^+$, NH$_4^+$, NO$_3^-$, and O$_2^-$ with vaporized aprotic ionic liquids in their intact ion pair form. Ion pair fragmentation to the cation or anion, as well as ion exchange and ion addition processes are observed by selected ion flow tube mass spectrometry. Free energies of the reactions involving 1-ethyl-3-methylimidazolium bis-trifluoromethylsulfonylimide determined by ab initio quantum mechanical calculations indicate that ion exchange or ion addition are energetically more favorable than charge transfer processes, whereas charge transfer processes can be important in reactions involving 1-butyl-3-methylimidazolium dicyanamide.
Table of Contents Figure: NH$_4^+$NTf$_2^-$EMIM$^+$ ion cluster geometry.

**Keywords:** RTIL, ion pair, SIFT, ion attachment, carbene.
Due to the high heats of vaporization involved with room temperature ionic liquids (RTIL), direct detection of RTIL vapors has proved difficult. Primary attempts to detect ionic liquid vapors under reduced pressure and to understand the nature of their volatilization have focused on mass spectrometric methods,\(^1\text{-}^4\) and, more recently, IR\(^5\text{-}^6\) and UV-VIS spectroscopy.\(^7\) Indirect methods involve transpiration,\(^8\) effusion\(^9\) or thermal gravimetric analyses.\(^10\text{-}^11\) To our knowledge, only one prior study has directly detected evaporated intact ion pairs, albeit under rather harsh thermal conditions.\(^12\) One drawback using positive mode ionization mass spectrometric techniques is that upon removing an electron from the ion pair, the coulombic attraction between the cation and anion is lost, and the remaining attractive forces between the cation-radical complex can be weak, and the complex can easily dissociate to the cation and radical. It is the detection of the intact cation which implies the presence of ion pairs in the vapor phase.\(^4\text{-}^13\) If, instead of removing an electron from the anion in the ion pair, a third ion can be gently attached to the ion pair, this would enable the detection of the larger charged product via mass spectrometry. Previous work using electrospray ionization mass spectrometry (ESI-MS),\(^14\) has detected multiple ion clusters: \((\text{C}^+)\text{nA}^-\text{(n-1)}\) in positive ion mode or \((\text{C}^+)\text{(n-1)A}^-\text{(n)}\) in negative ion mode. However, these clusters are formed under field evaporation conditions, not reduced pressure distillation conditions. Ion-ion pair reactions have been observed from reduced pressure evaporation of ionic liquids by Fourier transform ion cyclotron resonance mass spectrometry (FT-ICR-MS).\(^15\) In the FT-ICR-MS experiments, an evaporated ion pair dissociatively ionizes; the cation is trapped and can then add to another ion pair, forming stable clusters similar to those seen in ESI-MS experiments which are based on the original ionic liquid being evaporated.
The goal of this work was to expose vaporized ion pairs to different and more labile ions to look for intact RTIL ion pairs through proton transfer, charge exchange, clustering, or possible reactivity. Ions were chosen in order to observe: 1) a protonated ion pair via proton transfer from \( \text{NH}_4^+ \) (proton affinity (PA)\((\text{NH}_3)=8.85 \text{ eV}) \), 2) a negatively charged ion pair by electron transfer from \( \text{O}_2^- \) (electron affinity (EA)\((\text{O}_2)=0.45 \text{ eV}) \), and 3) soft chemical ionization of the ion pair by electron transfer from the ion pair to \( \text{NO}^+ \) (ionization potential (IP)\((\text{NO})=9.27 \text{ eV}) \). The relative PAs, EAs, and IPs of the ion pairs should indicate potential reactivity with the above ions. Apparently, in the systems considered here, the coulombic energy gained by ion addition or ion exchange can dominate the possible charge transfer processes.

Experiments were performed in the AFRL-Hanscom selected-ion flow tube (SIFT) mass spectrometric apparatus, which has been modified to introduce an effusive RTIL source. In order to produce conditions most likely to observe the intact ion pairs, the flow tube was cooled to 170 K. The reactions of \( \text{NO}^+ \), \( \text{NH}_4^+ \), \( \text{NO}_3^- \), and \( \text{O}_2^- \) with the vaporized ionic liquid 1-ethyl-3-methylimidazolium bis-trifluoromethylsulfonylimide (EMIM\(^+\)NTf\(_2^-\)) and \( \text{NO}^+ \) and \( \text{NH}_4^+ \) with the vaporized ionic liquid 1-butyl-3-methylimidazolium dicyanamide (BMIM\(^+\)dca\(^-\)) were monitored in positive ion mode for \( \text{NO}^+ \) and \( \text{NH}_4^+ \), and in negative ion mode for \( \text{NO}_3^- \), and \( \text{O}_2^- \) ions. When \( \text{EMIM}^+\text{NTf}_2^- \) (mass 391, IP=8.23 eV (MP2/6-311++G(d,p)) is reacted with \( \text{NO}^+ \) or \( \text{NH}_4^+ \), the major ion detected is mass 111, the EMIM\(^+\) cation, seen in Figure 1a and b. A small product peak also appears at mass 502 corresponding to the secondary reaction \( \text{EMIM}^+ + \text{EMIM}^+\text{NTf}_2^- \rightarrow \text{EMIM}^+\text{NTf}_2^-\text{EMIM}^+ \). For \( \text{NH}_4^+ \), a mass peak at 409 corresponding to \( \text{EMIM}^+\text{NTf}_2^-\text{NH}_4^+ \) is also detected (Figure 1b). \( \text{EMIM}^+\text{NTf}_2^- + \text{NO}_3^- \) yields a mass at 453, corresponding to \( \text{NO}_3^- \text{EMIM}^+\text{NTf}_2^- \) (Figure 2a). When \( \text{EMIM}^+\text{NTf}_2^- \) is reacted with \( \text{O}_2^- \), the major ion detected is mass 280, the \( \text{NTf}_2^- \) anion, seen in Figure 2b. When BMIM\(^+\)dca\(^-\) is reacted with \( \text{NH}_4^+ \) mass...
peaks include 139, 96, 125 and 82 (Figure 3a). When BMIM$^{+}$dca$^{-}$ is reacted with NO$^+$, mass peaks appear at 139, 125, 169, 179, 155, and 152 (Figure 3b). Possible reactions involved in the formation of the ions that correspond to the experimental mass peaks observed for EMIM$^{+}$NTf$_2$$^{-}$ are listed in Table 1, along with their enthalpies and free energies of reaction (at 298 K).

In the cases when EMIM$^{+}$NTf$_2$$^{-}$ is reacted with NO$^+$ or NH$_4^+$, the EMIM$^{+}$ (mass 111) ion is readily formed, presumably through a C$_i^+$A$^-$$C_j^+$ type of intermediate which dissociates to EMIM$^{+}$ cation plus a cofragment, either an ion pair, NO$^+$NTf$_2$$^{-}$, or a hydrogen bonded complex, NH$_3$HNTf$_2$, which are predicted to be exergonic ($\Delta G = -35.9$ and -25.6 kcal/mol, reactions 2 and 4, Table 1). Since masses 391 or 392 were not detected, simple charge transfer via electron removal or proton attachment to the ion pair apparently is not an important process. The relatively high concentration of EMIM$^{+}$ cation produced can then react with the RTIL ion pair to form an EMIM$^{+}$NTf$_2$$^{-}$EMIM$^{+}$ ion-trio ($\Delta G = -21.2$ kcal/mol, reaction 12, Table 1) at mass 502. Similarly, the formation of the NTf$_2$$^{-}$ anion is observed upon reaction of EMIM$^{+}$NTf$_2$$^{-}$ with O$_2^-$, but the formation of NTf$_2$$^{-}$EMIM$^{+}$NTf$_2$$^{-}$ (mass 671) or larger cluster ions cannot be detected as they are out of the mass range of the present detector (amu $\leq$600). The co-product of the NTf$_2$$^{-}$ anion formed upon reaction with O$_2^-$ is unclear. Some of the possible processes considered (reactions 8-11, Table 1) all appear to be endergonic, although the formation of the HO$_2$/EMIM carbene$^{16}$ complex product (reaction 8) has a $\Delta G$ of only +3.9±10 kcal/mol, and therefore would be the most likely route amongst reactions 8-11. The lack of signal at mass 391 suggests that electron transfer from O$_2^-$ or NO$_3^-$ is not an important process. Finally, small ion signals corresponding to ion addition to the ion pair is observed when EMIM$^{+}$NTf$_2$$^{-}$ is reacted with NH$_4^+$ or NO$_3^-$, resulting in EMIM$^{+}$NTf$_2$$^{-}$NH$_4^+$ (mass 409, $\Delta G = -40.8$ kcal/mol) or NO$_3^-$EMIM$^{+}$NTf$_2$$^{-}$ (mass 453, $\Delta G = -28.0$ kcal/mol) in reactions 6 and 7, respectively. Detection of the ion-trio
definitely shows the presence of neutral RTIL vapors in the flow tube. Some excess internal energy gained upon ion addition to the ion pair can be removed by collisions with the buffer gas at 170 K, which is consistent with the absence of the complex under room temperature conditions, indicating the bonds holding the complex together are weak. However, more exergonic processes may not be observed at 170 K if the dissociation rate of the internally excited complex is faster than thermalization by collisional de-excitation. The calculated geometry of EMIM\(^+\)NTf\(_2\)\(^-\)NH\(_4\)\(^+\) in Figure 4 shows a dramatic change in geometry of the EMIM\(^+\)NTf\(_2\)\(^-\) from the previously calculated geometries\(^5,17\) upon addition of NH\(_4\)\(^+\): The NTf\(_2\)\(^-\) migrates from being essentially coplanar with the EMIM\(^+\) ring in the ion pair to being perpendicular to the ring and is sandwiched between the EMIM\(^+\) and NH\(_4\)\(^+\) cations in the EMIM\(^+\)NTf\(_2\)\(^-\)NH\(_4\)\(^+\) ion-trio.

When BMIM\(^+\)dca\(^-\) (IP=7.4 eV, MP2/6-311++G(d,p)//B3LYP/6-311++G(d,p)) is reacted with NH\(_4\)\(^+\) or NO\(^+\), the observation of a peak at mass 139 indicates the formation of the BMIM\(^+\) cation. Similar to the EMIM\(^+\)NTf\(_2\)\(^-\) system, this could be from the dissociation of the C\(_i\)\(^+\)A\(^-\)C\(_j\)\(^+\) intermediate (BMIM\(^+\)dca\(^-\)NO\(^+\) or BMIM\(^+\)dca\(^-\)NH\(_4\)\(^+\)). However, unlike the EMIM\(^+\)NTf\(_2\)\(^-\) system, no peaks were detected corresponding to the secondary product BMIM\(^+\)dca\(^-\)BMIM\(^+\) at mass 344. This could be due to the lower vapor pressure, and thus lower concentration, of BMIM\(^+\)dca\(^-\) in the flow tube or weaker bonds. The presence of peaks with masses less than 139 could be due to fragmentation of the BMIM\(^+\) cation, as masses 82, 96 and 125 are seen in photoionization studies of BMIM\(^+\)dca\(^-\).\(^18\) The complexity of the BMIM\(^+\)dca\(^-\) mass spectra is likely due to the increased free energy release from charge transfer reactions compared to the similar charge transfer reactions in the EMIM\(^+\)NTf\(_2\)\(^-\) system (reactions 1 and 3, Table 1). The remaining mass peaks...
above mass 139 may result from more complex chemistry involving ion-neutral interactions, and this will be addressed in a future publication.

This work reports the experimental results of reacting vaporized ionic liquids 1-ethyl-3-methylimidazolium bis-trifluorosulfonylimide and 1-butyl-3-methylimidazolium dicyanamide with various ions including NO$^+$, NH$_4^+$, NO$_3^-$, and O$_2^-$. The presence of intact EMIM$^+$NTf$_2^-$ ion pairs in the gas phase is evidenced by the ion cluster (mass 502) formation seen in Figures 1 and 2. The presence of intact BMIM$^+$dca$^-$ ion pairs in the gas phase is less direct and indicated by the formation of the intact BMIM$^+$ cation (mass 139) seen in Figure 3. This experimental technique enables the examination of the influence of various energetic factors such as proton affinity, electron affinity, and ionization potential of the reactants on the outcome of the reactivity of vaporized RTIL ion-pairs with reagent ions. While dissociation to the cation or anion, ion exchange or ion addition dominate for the EMIM$^+$NTf$_2^-$-ion reactions, more complex reactivity is observed for the BMIM$^+$dca$^-$-ion reactions. More experimental and theoretical work is needed to evaluate these influences in RTIL ion pair-ion reactions.

**Experimental**

The measurements were carried out using the Air Force Research Laboratory Selected Ion Flow Tube (SIFT), which has been well described in detail previously.$^{19}$ Ions are made in a high pressure ion source by electron impact, mass selected in a quadrupole mass filter, and injected into a helium buffered flow tube through a Venturi inlet. Source gases were O$_2$ for O$_2^-$, HNO$_3$ for NO$_3^-$, NO or NO$_2$ for NO$^+$, and NH$_3$ for NH$_4^+$. In all cases, the primary ion accounted for >95% of all signal in the absence of the added ionic liquid. The helium carries the primary ions past a neutral inlet and the core of the flow is sampled into a quadrupole mass spectrometer for ion identification. The pressure inside the flow tube is 0.5 Torr. Since one of the main aims of the present experiments was to observe intact ion pairs whose ionized forms can readily
dissociate, we ran the flow tube at 170 K to minimize thermal dissociation of the ions. The mass spectrometer was calibrated with C\textsubscript{7}F\textsubscript{4} and the masses are accurate to 1 amu.

What is described above is quite standard and straightforward for the SIFT. What is more challenging is to introduce ionic liquids into the flow tube since their vapor pressure is low. The inlet is similar to one we have used for sulfuric acid.21 A ¼” glass tube is wrapped with insulated heating wire in two zones, a short one near the tip and a longer one covering most of the inlet. Resistance temperature detectors measure the temperature of the glass tube in each zone. The inlet extends into the center of the flow tube. Since the SIFT has a unipolar plasma that charges insulating surfaces, the whole inlet is wrapped by a stainless steel shim connected to ground. Glass wool is inserted inside the length of the glass tube and a small amount of ionic liquid is distributed throughout the glass wool by syringe injection. Helium flush gas of up to 145 sccm is used to transport the ionic liquid to the flow tube, and a He flow of 10,000 sccm was present in the SIFT.

Before introducing a new RTIL sample into the SIFT apparatus, methanol was used to thoroughly clean the inlet, the inlet was dried and new glass wool was inserted. The temperature of the inlet was kept at approximately 433 K and 453 K for BMIM\textsuperscript{+}dca\textsuperscript{-} and EMIM\textsuperscript{+}NTf\textsubscript{2}\textsuperscript{-} respectively. Preliminary measurements indicated it was easy to overheat and pyrolyze the RTIL so heating was done slowly.

Two types of data were taken: 1) mass spectra with a constant helium flow and 2) peak stepping through all relevant peaks (minor peaks ~ 1-2% were excluded) as a function of the helium flow. The latter data were taken to look for secondary reactions. As the flow of helium increased, the flow of ionic liquid also increased. In order to account for potential secondary reactions, we used the standard SIFT method\textsuperscript{22} of plotting the fraction of each product vs. neutral
flow rate (in this case, the He flush gas – plots are standard and not shown). Extrapolations of the branching to zero flow yield primary branching fractions, negative slopes indicate products that further react with the RTIL and positive slopes indicate products formed by secondary chemistry.

Although the vapor pressures of the ILs can be estimated\textsuperscript{20} and are thought to be in the range of 0.01 to 0.1 mTorr, we did not attempt to measure rate constants. However, for all species, the product abundances appeared roughly constant for the same He flow through the inlet indicating that the rates were similar and fast, i.e. near the collision rate.

The \textit{ab initio} calculations were performed at the MP2/6-311++G(d,p) and B3LYP/6-311++G(d,p) levels of theory,\textsuperscript{23-35} using the GAMESS\textsuperscript{36,37} quantum chemistry code. Reaction enthalpies and free energies (MP2) were computed using the ideal gas + rigid rotor + harmonic oscillator models\textsuperscript{32} with the computed vibrational frequencies scaled by a factor of 0.9748.\textsuperscript{35} The uncertainty assigned to the calculated reaction enthalpies in Table 1 is ± 10 kcal/mol, which was obtained by the procedure described in the Supplementary Information.

Acknowledgements

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Supporting Information Available: In order to estimate the uncertainty in the MP2 calculations in Table 1, reaction energies using MP2/aug-cc-pvtz and CCSD(T)/6-311++G(d,p) single point energies were computed and are available in the supplementary material. This material is available free of charge via the Internet at http://pubs.acs.org.

References


Table 1. Enthalpies and free energies of reaction of the RTIL EMIM$^+$NTf$_2^-$ with various ions at 298 K (MP2/6-311++G(d,p)). The EMIM carbene species in reaction 10 is denoted as EMIM:.

<table>
<thead>
<tr>
<th>Reaction</th>
<th>$\Delta H$ (298 K) (kcal/mol)</th>
<th>$\Delta G$ (298 K) (kcal/mol)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1) EMIM$^+$NTf$_2^-$ + NO$^+$ $\rightarrow$ EMIM$^+$ + NTf$_2^-$ + NO</td>
<td>3.6</td>
<td>-11.0</td>
</tr>
<tr>
<td>2) EMIM$^+$NTf$_2^-$ + NO$^+$ $\rightarrow$ EMIM$^+$ + NO$^+$NTf$_2^-$</td>
<td>-32.8</td>
<td>-35.9</td>
</tr>
<tr>
<td>3) EMIM$^+$NTf$_2^-$ + NH$_4^+$ $\rightarrow$ EMIM$^+$ + HNTf$_2$ + NH$_3$</td>
<td>-3.3</td>
<td>-17.4</td>
</tr>
<tr>
<td>4) EMIM$^+$NTf$_2^-$ + NH$_4^+$ $\rightarrow$ EMIM$^+$ + NH$_3$HNTf$_2$</td>
<td>-19.8</td>
<td>-25.6</td>
</tr>
<tr>
<td>5) EMIM$^+$NTf$_2^-$ + NH$_4^+$ $\rightarrow$ EMIM$^+$ + NTf$_2$ + NH$_4$</td>
<td>117.5</td>
<td>103.0</td>
</tr>
<tr>
<td>6) EMIM$^+$NTf$_2^-$ + NH$_4^+$ $\rightarrow$ EMIM$^+$NTf$_2$NH$_4^+$</td>
<td>-47.7</td>
<td>-40.8</td>
</tr>
<tr>
<td>7) EMIM$^+$NTf$_2^-$ + NO$_3^-$ $\rightarrow$ NO$_3^-$EMIM$^+$NTf$_2^-$</td>
<td>-36.3</td>
<td>-28.0</td>
</tr>
<tr>
<td>8) EMIM$^+$NTf$_2^-$ + O$_2^-$ $\rightarrow$ NTf$_2^-$ + EMIM:HO$_2$</td>
<td>8.9</td>
<td>3.9</td>
</tr>
<tr>
<td>9) EMIM$^+$NTf$_2^-$ + O$_2^-$ $\rightarrow$ NTf$_2^-$ + EMIM$^+$O$_2^-$</td>
<td>13.4</td>
<td>10.3</td>
</tr>
<tr>
<td>10) EMIM$^+$NTf$_2^-$ + O$_2^-$ $\rightarrow$ NTf$_2^-$ + EMIM: + HO$_2$</td>
<td>27.1</td>
<td>13.0</td>
</tr>
<tr>
<td>11) EMIM$^+$NTf$_2^-$ + O$_2^-$ $\rightarrow$ NTf$_2^-$ + EMIM + O$_2$</td>
<td>24.7</td>
<td>12.1</td>
</tr>
<tr>
<td>12) EMIM$^+$NTf$_2^-$ + EMIM$^+$ $\rightarrow$ EMIM$^+$NTf$_2$EMIM$^+$</td>
<td>-32.4</td>
<td>-21.2</td>
</tr>
</tbody>
</table>
Figure 1. Positive ion mass spectra for the reactions (a) EMIM$^+$NTf$_2^-$ + NO$^+$ and (b) EMIM$^+$NTf$_2^-$ + NH$_4^+$

Figure 2. Negative ion mass spectra for the reactions (a) EMIM$^+$NTf$_2^-$ + NO$_3^-$ and (b) EMIM$^+$NTf$_2^-$ + O$_2^-$
Figure 3. Positive ion mass spectra for the reactions of (a) BMIM$^+$dca$^-$ + NH$_4^+$ and (b) BMIM$^+$dca$^-$ + NO$^+$

Figure 4. NH$_4^+$NTf$_2^-$EMIM$^+$ ion cluster geometry.
Reactions of ions with ionic liquid vapors by selected-ion flow tube mass spectrometry

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Supplementary Information

The uncertainties in the computed reaction enthalpies were estimated by performing single point energy calculations (a) at the MP2 level using the augmented correlation consistent polarized valence triple-zeta basis sets of Dunning and Woon,\(^1,2\) denoted as MP2/aug-ccpvtz//MP2/6-311++G(d,p), and (b) at the coupled cluster level using single and double excitations with perturbative estimates of triples (CCSD(T))\(^3-5\) using the 6-311++G(d,p) basis set, denoted as CCSD(T)/6-311++G(d,p)/MP2/6-311++G(d,p), for the two reactions listed in Table S1. The reactions in Table S1 are similar to those in Table 1, except that the alkyl ligands on the ethylmethylimidazolium ring have been replaced with hydrogen (denoted as IMID in Table S1) in order to make the CCSD(T) calculations tractable.

For each reaction, the sum of the absolute differences in reaction energies relative to MP2/6-311++G(d,p) is taken as a lower bound to the uncertainty in the reaction enthalpy. As shown in Table S1, the larger of the uncertainties for the two reactions is 8.2 kcal/mol, which is
the basis for the estimated uncertainty for the reaction enthalpies listed in Table 1 of ±10 kcal/mol.

**Table S1.** MP2/aug-ccpvtz//MP2-6-311++G(d,p) and CCSD(T)/6-311++G(d,p)//MP2-6-311++G(d,p) reaction energies at 0 K, excluding zero point vibrational energy corrections, in kcal/mol.

<table>
<thead>
<tr>
<th>Reaction</th>
<th>∆E elec (MP2/6-311++G(d,p))</th>
<th>∆E elec (MP2/aug-ccpvtz//MP2/6-311G++(d,p))</th>
<th>∆E elec (CCSD(T)/6-311++G(d,p)//MP2/6-311G++(d,p))</th>
<th>∆MP2</th>
<th>∆CCSD(T)</th>
<th>uncertainty ≤ ∆MP2 + ∆CCSD(T)</th>
</tr>
</thead>
<tbody>
<tr>
<td>IMID⁺NTf₂⁻ + NO₂⁺ → IMID⁺ + NO₂⁺NTf₂</td>
<td>-27.2</td>
<td>-26.8</td>
<td>-35.0</td>
<td>0.4</td>
<td>7.8</td>
<td>8.2</td>
</tr>
<tr>
<td>IMID⁺NTf₂⁻ + NH₄⁺ → IMID⁺NTf₂NH₄⁺</td>
<td>-45.1</td>
<td>-45.1</td>
<td>-45.2</td>
<td>0.0</td>
<td>0.1</td>
<td>0.1</td>
</tr>
</tbody>
</table>

**References**