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## Ion pair extractant selective for LiCl and LiBr†

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Improved methods for achieving the selective extraction of lithium salts from lithium sources, including rocky ores, salt-lake brines, and end-of-life lithium-ion batteries, could help address projected increases in the demand for lithium. Here, we report an ion pair receptor (**2**) capable of extracting LiCl and LiBr into an organic receiving phase both from the solid state and from aqueous solutions. Ion pair receptor **2** consists of a calix[4]pyrrole framework, which acts as an anion binding site, linked to a phenanthroline cation binding motif *via* ether linkages. Receptor **2** binds MgBr<sub>2</sub> and CaCl<sub>2</sub> with high selectivity over the corresponding lithium salts in a nonpolar aprotic solvent. The preference for Mg<sup>2+</sup> and Ca<sup>2+</sup> salts is reversed in polar protic media, allowing receptor **2** to complex LiCl and LiBr with high selectivity and affinity in organic media containing methanol or water. The effectiveness of receptor **2** as an extractant for LiCl and LiBr under liquid–liquid extraction (LLE) conditions was found to be enhanced by the presence of other potentially competitive salts in the aqueous source phase.

## Introduction

The lithium ion (Li<sup>+</sup>) supports key features of modern life; it is an essential component in lithium ion batteries (LIBs), ceramics, and lubricating greases, and is used as a pharmaceutical agent in treating depressive illness.<sup>1–6</sup> Currently rechargeable LIBs account for approximately 80% of the end use of lithium.<sup>7</sup> According to the U.S. Geological Survey (USGS), global consumption of lithium rose from approximately 95 000 tons in 2021 to 134 000 tons in 2022.<sup>7</sup> While the global demand for lithium is increasing, the available lithium reserves remain limited. In addition, selective separation of the lithium ion from its sources requires an energy- and labor-intensive process, and can be time-consuming.<sup>8,9</sup> In principle, lithium may be separated from rocky ores, salt lakes, brines, and sea water.<sup>8–10</sup> When rocky ores or clays are used as the lithium source, roasting at a high temperature (1100 °C) is followed by baking at 250 °C in acid.<sup>8</sup> Undesired salts are then removed *via* several energy- and water-intensive steps that are a source of environmental concern.<sup>11–13</sup> The ocean is the largest source of lithium; however, selective extraction of lithium from seawater is challenging

because of the very low concentration of the lithium cation amounting to only 0.1–0.2 ppm while other potentially competitive ions exist at much higher concentrations.<sup>14–19</sup> At present, salt lake brines supply the majority of the commercial lithium. However, the extraction of lithium from brine reservoirs typically requires evaporation of residual water over a period of months to years.<sup>20–26</sup> A huge amount of water is also necessary to remove unwanted ions and contaminants. Because of the shortcomings of conventional extraction processes, efforts are being devoted increasingly to so-called direct lithium extraction (DLE) methods. In this context, the use of porous lithium sorbents and ion exchange materials has attracted attention because of their relative simplicity and potential to operate at relatively low levels of environmental stress.<sup>27,28</sup> Unfortunately, most DLE methods developed thus far require additional processing steps to free the lithium ions from the lithium adsorbing materials.<sup>27,28</sup> Therefore, so-called liquid–liquid extraction (LLE) and solid–liquid extraction (SLE), wherein an extractant capable of selectively complexing lithium salts is used to promote DLE, continue to attract interest. In principle, these approaches could enable the direct and selective extraction of lithium salts from mixed salt solid phases or brines. Nevertheless, it remains challenging to design and construct an extractant possessing the ability to extract selectively lithium salts due to the small size of the lithium cation and its relatively high hydration energy ( $\Delta_{\text{hyd}}G^* = -475 \text{ kJ mol}^{-1}$ ), as well as the interference of other competing cations, such as Na<sup>+</sup>, K<sup>+</sup>, Mg<sup>2+</sup>, and Ca<sup>2+</sup>.<sup>29–31</sup> For instance, Mg<sup>2+</sup> not only has a similar ionic radius (72 pm for Mg<sup>2+</sup> vs. 69 pm for Li<sup>+</sup>), it also has a higher net charge, and is typically present at  $\geq 8\times$  the concentration of Li<sup>+</sup> in most salt lake brines.<sup>23,29</sup> The likely presence of Mg<sup>2+</sup> and other potential interferants

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underscores the challenge associated with designing Li<sup>+</sup>-selective extractants. One way to meet this challenge could involve the use of ion pair receptors. Here we report a new phenanthroline-strapped calix[4]pyrrole (**2**) that acts as a selective receptor for LiCl and LiBr in polar media and which promotes lithium salt extraction under LLE conditions.

Ion pair receptors are systems with an ability to complex concurrently both a cation and an anion. Appropriately designed ion pair receptors display enhanced selectivity and affinity for target ions relative to cation or anion receptors, systems that bind either a cation or an anion, but not both.<sup>32–36</sup> Although numerous ion pair receptors capable of binding various alkali metal salts have been reported to date, only a very small number of receptors were found to bind lithium salts with sufficient affinity and selectivity to permit effective SLE or LLE of Li<sup>+</sup> salts.<sup>37–39</sup> Unfortunately, even in the most favorable cases the actual lithium extraction efficiencies proved very low. For instance, we reported ion pair receptors based on calix[4]pyrroles strapped with pyridine-fused multi-aromatic rings containing a methoxybenzene group. These heteroditopic ion pair receptors proved capable of extracting LiCl or LiNO<sub>3</sub> from the solid state into nitrobenzene-*d*<sub>5</sub> or dichloromethane-*d*<sub>2</sub> under solid–liquid extraction conditions provided the salts were present in excess.<sup>37,39</sup> The phenanthroline-strapped calix[4]pyrrole (**1**) having ester linkers was found to extract LiCl into chloroform from an aqueous solution containing near-saturated quantities of LiCl, but not at lower source phase concentrations.<sup>39</sup> In the case of receptor **1**, the presence of the electron-withdrawing ester groups directly linked to the phenanthroline cation binding site was thought to reduce the inherent Li<sup>+</sup> affinity accounting for the limits on its extraction ability. As inferred from a solid state X-ray crystal structure and solution phase <sup>1</sup>H NMR spectral studies, the ester carbonyl oxygen atoms present in receptor **1** do not participate in complexation with lithium.<sup>39</sup> We thus considered it likely that an analogue of **1** that incorporated oxygen donor sites within the tethering subunits that link the phenanthroline strap to the calix[4]pyrrole core would prove more effective as an ion pair receptor for lithium salts, such as LiCl and LiBr. With such considerations in mind, we designed a new ion pair receptor (**2**) wherein the ester groups present in **1** are replaced by phenoxy ether linkages (Fig. 1). The relatively rigid nature of the phenoxy

linkers was also expected to provide for increased structural preorganization within the receptor framework leading to improved LiCl and LiBr recognition. As detailed below, receptor **2** binds LiCl and LiBr with high selectivity and with an affinity ( $K_a > 10^5 \text{ M}^{-1}$  for LiCl in 10% methanol in chloroform-*d*) that exceeds that of receptor **1** ( $K_a = 180 \pm 7$ ), as well as all other lithium ion receptors of which we are aware. Receptor **2** also acts as an effective LLE extractant promoting the transfer of LiCl and LiBr from aqueous solutions into dichloromethane or nitrobenzene receiving phases.

## Results and discussion

We previously reported that the ion pair receptor **1**, a phenanthroline-strapped calix[4]pyrrole with ester linkers, was able to extract LiCl from LiCl-saturated aqueous solutions into a chloroform layer.<sup>39</sup> However, receptor **1** failed to extract LiCl from aqueous solutions containing relatively low concentrations of LiCl. Moreover, receptor **1** was not examined for its LiCl selectivity in the presence of MgX<sub>2</sub>, CaX<sub>2</sub>, NaX, or KX (X = Cl and Br), which coexist in most lithium sources, under LLE conditions.<sup>39</sup> As noted above, receptor **2**, wherein relatively rigid phenoxy groups serve to link the phenanthroline cation and calix[4]pyrrole anion recognition subunits, was prepared in an effort to address these shortcomings.

The synthesis of receptor **2** is summarized in Scheme 1. Briefly, phenanthroline ditosylate **3** and *cis*-bisphenolic calix[4]pyrrole **4** were prepared following known literature procedures (Scheme 1).<sup>40,41</sup> Reaction of compound **3** with calix[4]pyrrole **4** in the presence of K<sub>2</sub>CO<sub>3</sub> as a base in acetonitrile gave the desired ion pair receptor **2** in 11% yield. Receptor **2** was characterized by means of <sup>1</sup>H and <sup>13</sup>C NMR spectroscopies and high resolution QTOF (quadrupole time of flight) mass spectrometry, as well as a single crystal X-ray diffraction analysis of its LiBr complex.

Initially, we examined the ability of receptor **2** to bind various alkali and alkaline earth metal chloride salts, including LiCl, NaCl, KCl, MgCl<sub>2</sub>, and CaCl<sub>2</sub> in CH<sub>3</sub>OH/CDCl<sub>3</sub> (1 : 9, v/v) using <sup>1</sup>H NMR spectroscopy. This specific solvent system was chosen with consideration of the solubility of both the receptor and the test salts in mind. Upon exposure of receptor **2** to excess LiCl (*ca.* 100 equiv.), noticeable chemical shift changes were observed in the proton signals of both the calix[4]pyrrole subunit and the phenanthroline group, findings leading us to conclude that receptor **2** forms an ion pair complex with LiCl (Fig. S1 and S2, see the ESI† for details). By contrast, no chemical shift movements were seen in the presence of NaCl and KCl, which was taken as evidence for receptor **2** being incapable of complexing NaCl and KCl. In contrast, in the

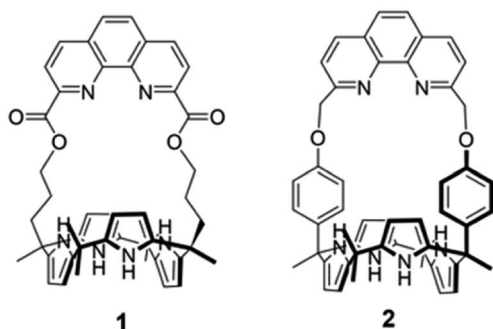
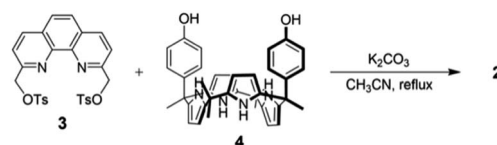


Fig. 1 Chemical structures of ion pair receptors **1** and **2**.



Scheme 1 Synthesis of receptor **2**.



presence of  $\text{MgCl}_2$  (ca. 100 equiv.), roughly 16% of receptor 2 forms a complex with  $\text{MgCl}_2$  under these conditions ( $\text{CH}_3\text{OH}/\text{CDCl}_3$ ; 1 : 9, v/v), which leads us to suggest that the affinity of receptor 2 for  $\text{MgCl}_2$  is very low ( $K_a < 5 \text{ M}^{-1}$ ) (Fig. S1†).<sup>42,43</sup> In analogy to what was seen with  $\text{LiCl}$ ,  $\text{Mg}^{2+}$  is presumed to be bound to the phenanthroline unit while one of two  $\text{Cl}^-$  anions is bound to the calix[4]pyrrole moiety *via* hydrogen bonds (Fig. S1 and S2; see the ESI† for details). Distinct chemical shift changes were also observed in the  $^1\text{H}$  NMR spectrum when receptor 2 was treated with  $\text{CaCl}_2$  in  $\text{CH}_3\text{OH}/\text{CDCl}_3$  (1 : 9, v/v) (Fig. S1†). These  $^1\text{H}$  NMR spectral changes are consistent with the  $\text{Ca}^{2+}$  cation only, and not the  $\text{Cl}^-$  anion, being complexed within receptor 2 (Fig. S2†). A follow up  $^1\text{H}$  NMR spectral titration with  $\text{CaCl}_2$  provided further support for this binding mode (Fig. S3; see the ESI† for details). This non-ion pair binding mode stands in contrast to what was seen in the cases of  $\text{LiCl}$  and  $\text{MgCl}_2$  (Fig. S2†). Taken in concert, these findings are thought to reflect a cation recognition site that is near-optimal for  $\text{Li}^+$  cation complexation, somewhat small for fully effective  $\text{Na}^+$  and  $\text{K}^+$  recognition, and essentially able to accommodate only  $\text{Ca}^{2+}$  without  $\text{Cl}^-$  being co-bound.

We also investigated the capability of receptor 2 to complex the bromide salts of alkali and alkaline earth metal cations, including  $\text{LiBr}$ ,  $\text{NaBr}$ ,  $\text{KBr}$ ,  $\text{MgBr}_2$  and  $\text{CaBr}_2$ , in 10% methanol in chloroform-*d*. In contrast to what was seen with the corresponding chloride salts, upon exposure of receptor 2 to the respective bromide salts, only  $\text{LiBr}$  gave rise to chemical shift movements attributable to ion pair complexation (Fig. S4†). This finding is taken as evidence that receptor 2 is capable of binding  $\text{LiBr}$  selectively over other test bromide salts. We also examined the binding selectivity of receptor 2 for other lithium halide salts, *i.e.*,  $\text{LiF}$ ,  $\text{LiCl}$ ,  $\text{LiBr}$ , and  $\text{LiI}$ , in  $\text{CH}_3\text{OH}/\text{CDCl}_3$  (1 : 9, v/v). Addition of  $\text{LiCl}$  and  $\text{LiBr}$  produced chemical shift changes for the signals of receptor 2 in the  $^1\text{H}$  NMR spectra ascribable to co-binding of the lithium cations and the halide anions while  $\text{LiF}$  caused no appreciable chemical shift changes (Fig. S5†). When receptor 2 was exposed to  $\text{LiI}$ , only the lithium cation was bound to the phenanthroline moiety without the iodide anion being co-complexed by the calix[4]pyrrole subunit (Fig. S5; see the ESI† for details).

To quantify the ability of receptor 2 to bind cations and anions as well as various test ion pairs, we performed  $^1\text{H}$  NMR spectroscopic titrations in  $\text{CH}_3\text{OH}/\text{CDCl}_3$  (1 : 9, v/v). For instance, upon the titration of receptor 2 with  $\text{LiCl}$ , the resulting  $^1\text{H}$  NMR spectral changes support the conclusion receptor 2 binds  $\text{LiCl}$  quantitatively with a 1 : 1 binding stoichiometry *via* a binding-release equilibrium that is slow on the NMR time scale (Fig. S6; see the ESI† for details). The association constant for  $\text{LiCl}$  approximated from this titration experiment was found to be  $K_a > 10^5 \text{ M}^{-1}$ . This value is at least three orders of magnitude larger than what was found in the case of receptor 1 (Table 1 and Fig. S7†).<sup>42</sup>

Receptor 2 was also found to bind  $\text{LiBr}$ , albeit with low affinity relative to  $\text{LiCl}$  and *via* an equilibrium process that is fast on the NMR timescale (Fig. S8, see the ESI† for details). The corresponding  $\text{LiBr}$  association constant was calculated to be  $K_a = 63 \pm 3 \text{ M}^{-1}$ .<sup>43</sup> The relatively low affinity for  $\text{LiBr}$  is presumably

**Table 1** Association constants ( $K_a$ ,  $\text{M}^{-1}$ )<sup>a</sup> corresponding to the interaction of receptor 2 with the lithium cation and selected halide anion salts as determined in  $\text{CH}_3\text{OH}/\text{CDCl}_3$  (1 : 9, v/v)

Host <sup>b</sup>	Guest	$K_a$ ( $\text{M}^{-1}$ )
2	$\text{Cl}^-$	No binding
2	$\text{Br}^-$	No binding
2	$\text{Li}^+$	$< 5^c$
1	$\text{LiCl}$	$180 \pm 7^c$
2	$\text{LiCl}$	$> 10^5$
1	$\text{LiBr}$	$24 \pm 1^c$
2	$\text{LiBr}$	$63 \pm 3^c$
2 + $\text{Li}^+$	$\text{Cl}^-$	$> 10^5$
2 + $\text{Li}^+$	$\text{Br}^-$	$148 \pm 5^c$
2 + $\text{Cl}^-$	$\text{Li}^+$	$> 10^5$
2 + $\text{Br}^-$	$\text{Li}^+$	$95 \pm 3^c$

<sup>a</sup> Values were obtained from  $^1\text{H}$  NMR spectroscopic titrations of 2.

<sup>b</sup> Unless otherwise indicated, the anions and lithium cations were used in the forms of their respective tetrabutylammonium and perchlorate salts. <sup>c</sup> The  $K_a$  value was approximated using BindFit v0.5 available from <https://app.supramolecular.org/bindfit>.

because the  $\text{Br}^-$  anion is too large to be co-complexed with the  $\text{Li}^+$  cation effectively within the receptor cavity. However, this association constant for  $\text{LiBr}$  is  $2.6 \times$  larger than that of receptor 1 (Table 1 and Fig. S9†). The association constants for  $\text{MgCl}_2$  and  $\text{CaCl}_2$  were likewise determined to be  $K_a < 5 \text{ M}^{-1}$  for both salts in 10%  $\text{CH}_3\text{OH}$  in  $\text{CDCl}_3$  (Fig. S1 and S3†).<sup>43</sup> Again, this finding is ascribed to a mismatch with the binding cavity present in receptor 2 as well as to larger solvation energies of the divalent cations (*vide infra*).

We also investigated the interactions of receptor 2 with  $\text{Cl}^-$ ,  $\text{Br}^-$ , and  $\text{Li}^+$  with non-coordinating counter ions in 10%  $\text{CH}_3\text{OH}$  in  $\text{CDCl}_3$ . When receptor 2 was treated with excess  $\text{Cl}^-$  and  $\text{Br}^-$  (as their TBA<sup>+</sup> (tetrabutylammonium) salts), no appreciable chemical shift changes were observed in the  $^1\text{H}$  NMR spectrum, a finding taken as evidence that receptor 2 fails to bind these halide anion salts in this protic solvent system (Fig. S10 and S11†). In contrast, in the presence of excess  $\text{Li}^+$  (as its  $\text{ClO}_4^-$  (perchlorate anion) salt), the CH proton signal ( $H_a$ ) of the phenanthroline subunit of receptor 2 underwent a slight downfield shift ( $\Delta\delta = 0.04$  ppm). Although modest, this change is thought to reflect  $\text{Li}^+$  cation complexation by the phenanthroline group (Fig. S10†). The association constant of receptor 2 for  $\text{LiClO}_4$  was calculated from this  $^1\text{H}$  NMR spectral titration to be  $< 5 \text{ M}^{-1}$  (Fig. S12†).<sup>43</sup> On the basis of these studies, we conclude that receptor 2 binds the  $\text{LiCl}$  and  $\text{LiBr}$  ion pairs far more effectively than the individual ions,  $\text{Cl}^-$ , and  $\text{Br}^-$ , and  $\text{Li}^+$ , when the latter were tested using a non-coordinating counter ion. These findings are rationalized in terms of the ion pair complexes with  $\text{LiCl}$  and  $\text{LiBr}$  being stabilized by electrostatic attractions between the co-bound anion and cation.

Consistent with the above conclusion, the ability of receptor 2 to bind the  $\text{Cl}^-$  and  $\text{Br}^-$  anions was markedly improved in the presence of the  $\text{Li}^+$  cation and *vice versa* (Fig. 2). For instance, when receptor 2 was titrated with  $\text{Cl}^-$  in the presence of  $\text{Li}^+$  in 10%  $\text{CH}_3\text{OH}$  in  $\text{CDCl}_3$ , a new set of proton signals attributable to the  $\text{LiCl}$  complex of receptor 2 emerged with saturation being



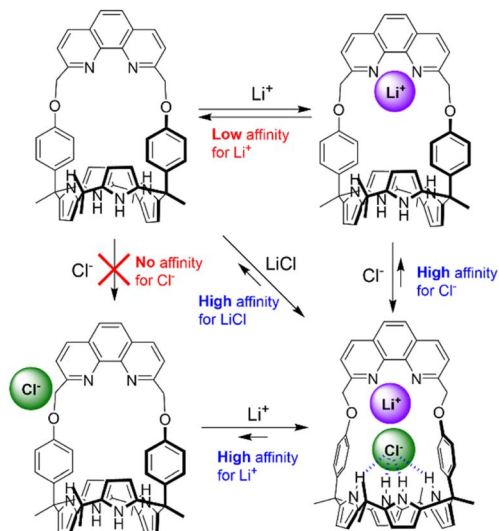


Fig. 2 Proposed ion binding modes of receptor 2 in the presence of separately  $\text{Li}^+$  and  $\text{Cl}^-$  (added with a non-coordinating counter ion) and in the presence of both  $\text{Li}^+$  and  $\text{Cl}^-$  in  $\text{CH}_3\text{OH}/\text{CDCl}_3$  (1 : 9, v/v) as inferred from  $^1\text{H}$  NMR spectroscopic studies.

achieved upon the addition of 1.0 equiv. of  $\text{Cl}^-$  (Fig. S13<sup>†</sup>). The association constant of receptor 2 for  $\text{Cl}^-$  in the presence of  $\text{Li}^+$  was found to be  $>10^5 \text{ M}^{-1}$  on the basis of this  $^1\text{H}$  NMR spectral titration experiment.<sup>42</sup> Receptor 2 was also found to bind the  $\text{Br}^-$  anion in the presence of the  $\text{Li}^+$  cation (*ca.* 50 equiv.) with an association constant of  $K_a = 148 \pm 5 \text{ M}^{-1}$  (Fig. S14<sup>†</sup>).<sup>43</sup> This value is considerably larger than what was seen in its absence (no affinity for  $\text{Br}^-$ ; *vide supra*). We thus conclude that co-complexation of  $\text{Cl}^-$  and  $\text{Br}^-$  with  $\text{Li}^+$  significantly enhances the affinity of 2 for these two halide anions, which are otherwise incapable of binding to the receptor (Fig. 2).

The affinity of receptor 2 for  $\text{Li}^+$  was also found to be significantly enhanced in the presence of  $\text{Cl}^-$  and  $\text{Br}^-$ . On the basis of the  $^1\text{H}$  NMR spectral titration of receptor 2 with  $\text{LiClO}_4$  in the presence of TBACl (5.0 equiv.), the association constant of receptor 2 for  $\text{Li}^+$  was estimated to be  $>10^5 \text{ M}^{-1}$  (Fig. S15; see the ESI<sup>†</sup> for details), a value that is higher than that measured in the absence of  $\text{Cl}^-$  by  $>100\,000$ -fold (Table 1).<sup>42</sup> In contrast, the affinity of receptor 2 for  $\text{Li}^+$  was enhanced in the presence of  $\text{Br}^-$  by only approx. 20-fold (Fig. S16<sup>†</sup> and Table 1). We thus conclude that cooperativity between the cation and anion binding sites within receptor 2 plays a crucial role in regulating the ion binding affinity of the receptor for appropriately chosen lithium halide ion pairs, with the effect being particularly dramatic in the case of  $\text{LiCl}$ , but also noteworthy for  $\text{LiBr}$ .

Further evidence that receptor 2 forms an ion pair complex with  $\text{LiBr}$  came from a single crystal X-ray diffraction analysis. Single crystals of the complex  $2 \cdot \text{LiBr}$  appropriate for an X-ray diffraction analysis were grown by allowing a mixture of  $\text{CHCl}_3$  and  $\text{CH}_3\text{OH}$  containing 2 and an excess  $\text{LiBr}$  to evaporate slowly. The resulting crystal structure revealed that the  $\text{Br}^-$  anion is bound to the calix[4]pyrrole moiety *via* hydrogen bonds with  $\text{N}-\text{H} \cdots \text{Br}^-$  distances of 2.50–2.51 Å while the  $\text{Li}^+$  cation is coordinated not only to the phenanthroline nitrogen atoms but

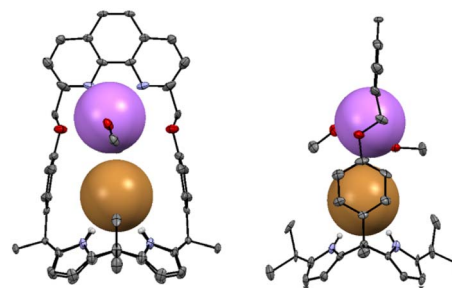


Fig. 3 Two different views of the X-ray crystal structure of the  $2 \cdot \text{LiBr}$ . Thermal ellipsoids are scaled to the 50% probability level. Most hydrogen atoms are omitted for clarity.

also to the ether oxygen atoms with  $\text{N} \cdots \text{Li}^+$  distances of 1.91 Å and 2.34 Å and  $\text{O} \cdots \text{Li}^+$  distances of 2.627 Å and 3.269 Å, respectively (Fig. 3). The  $\text{Li}^+$  cation is further stabilized by the coordination of two methanol molecules. The  $\text{Li}^+$  cation was also found to interact directly with the co-bound  $\text{Br}^-$  anion at a distance of 4.02 Å.

We also tested receptor 2 for its ability to solubilize and extract the solid  $\text{LiCl}$  and  $\text{LiBr}$  into dichloromethane- $d_2$  ( $\text{CD}_2\text{Cl}_2$ ). When receptor 2 along with  $\text{LiCl}$  or  $\text{LiBr}$  (*ca.* 100 equiv.) was subjected to sonication for 1 hour in  $\text{CD}_2\text{Cl}_2$ , distinct  $^1\text{H}$  NMR spectral changes attributable to the formation of ion pair complexes  $2 \cdot \text{LiCl}$  and  $2 \cdot \text{LiBr}$  were seen (Fig. S17<sup>†</sup>). These ion pair complexes proved soluble in  $\text{CD}_2\text{Cl}_2$  enabling receptor 2 to extract  $\text{LiCl}$  and  $\text{LiBr}$  into this organic phase under solid–liquid extraction conditions. When an equimolar ratio of  $\text{LiCl}$  and  $\text{LiBr}$  (*ca.* 100 equiv. each) was used, receptor 2 was found to complex both  $\text{LiCl}$  and  $\text{LiBr}$  at a nearly 1 : 1 ratio in  $\text{CD}_2\text{Cl}_2$ . This proved true in spite of the greater affinity displayed for  $\text{LiCl}$  over  $\text{LiBr}$  in  $\text{CH}_3\text{OH}/\text{CDCl}_3$ ; 1/9, v/v (Table 1). This seeming disparity is rationalized in terms of the relatively small lattice energy of  $\text{LiBr}$  in  $\text{CD}_2\text{Cl}_2$ . On the other hand, release of the lithium cation from  $2 \cdot \text{LiCl}$  could be triggered in  $\text{CD}_2\text{Cl}_2$  *via* the addition of the fluoride anion (as its tetrabutylammonium ( $\text{TBA}^+$ ) salt). This treatment enables recovery of  $\text{Li}^+$  in the form of its insoluble  $\text{LiF}$  salt by filtration (Fig. S18<sup>†</sup>). The process could be followed by  $^1\text{H}$  NMR spectroscopy (Fig. S18; see the ESI<sup>†</sup> for details).

The selectivity of 2 for the chloride and bromide salts of alkali and alkaline earth cations was also evaluated under SLE conditions using dichloromethane- $d_2$  as the receiving phase. Receptor 2 is soluble in this solvent, whereas the test salts are insoluble. In this aprotic solvent, receptor 2 exhibited different binding behavior as compared to 10%  $\text{CH}_3\text{OH}$  in  $\text{CDCl}_3$ . Upon treatment of receptor 2 with approx. 500 equiv. each of  $\text{LiCl}$ ,  $\text{NaCl}$ ,  $\text{KCl}$ ,  $\text{MgCl}_2$ , and  $\text{CaCl}_2$  in  $\text{CD}_2\text{Cl}_2$  (separate studies), only  $\text{LiCl}$  induced chemical shift changes in the spectrum that could be readily interpreted in terms of  $\text{LiCl}$  complexation (Fig. S19<sup>†</sup>). In contrast,  $\text{NaCl}$ ,  $\text{KCl}$ , and  $\text{CaCl}_2$  gave rise to no appreciable chemical shift changes in the  $^1\text{H}$  NMR spectrum of 2. Meanwhile, treatment with  $\text{MgCl}_2$  produced an orange precipitate (Fig. S19<sup>†</sup>). This latter finding is interpreted in terms of receptor 2 forming an ion pair complex with  $\text{MgCl}_2$  that is insoluble in



CD<sub>2</sub>Cl<sub>2</sub>. We next monitored the changes in the <sup>1</sup>H NMR spectrum of **2** when treated with a mixture of LiCl, NaCl, KCl, MgCl<sub>2</sub>, and CaCl<sub>2</sub> salts (*ca.* 100 equiv. each) in CD<sub>2</sub>Cl<sub>2</sub>. Under these conditions, an orange precipitate was formed with no discernable proton signals being seen in the <sup>1</sup>H NMR spectrum of the residual CD<sub>2</sub>Cl<sub>2</sub> layer. Given the analogy to what was seen when receptor **2** was treated with MgCl<sub>2</sub> alone, we interpret these findings in terms of receptor **2** capturing MgCl<sub>2</sub> in preference to the other chloride salts with this nominal selectivity being driven in part by solubility considerations (Fig. S20†).

In case of the corresponding bromide salts, receptor **2** was found to extract LiBr and MgBr<sub>2</sub> in the solid state into CD<sub>2</sub>Cl<sub>2</sub> but *via* different apparent binding modes. For instance, when exposed to LiBr in CD<sub>2</sub>Cl<sub>2</sub>, receptor **2** exhibited <sup>1</sup>H NMR spectral changes consistent with the formation of 2·LiBr where the Li<sup>+</sup> and Br<sup>−</sup> are bound to the phenanthroline nitrogen atoms and the calix[4]pyrrole NH protons, respectively (Fig. S20†). In contrast, exposure of **2** to MgBr<sub>2</sub> led the signal assignable to the calix[4]pyrrole NH protons to undergo a relatively small downfield shift ( $\Delta\delta = 0.26$  ppm for MgBr<sub>2</sub> vs.  $\Delta\delta = 3.31$  ppm for LiBr, respectively) while the proton signals of the phenanthroline and the phenoxy CH hydrogens were seen to undergo noticeable downfield shifts (Fig. S20†). These <sup>1</sup>H NMR spectral changes provide support for the notion that only the Mg<sup>2+</sup> cation is bound to the receptor with the two Br<sup>−</sup> counter anions being located outside the receptor cavity. In contrast, treatment of receptor **2** with CaBr<sub>2</sub> led to formation of a white solid with no observable proton signals appearing in the <sup>1</sup>H NMR spectrum of **2**. Again, this finding is consistent with the formation of a strong, insoluble complex with CaBr<sub>2</sub> (Fig. S20†). A similar phenomenon took place when receptor **2** in CD<sub>2</sub>Cl<sub>2</sub> was treated with an equimolar mixture of LiBr, NaBr, KBr, MgBr<sub>2</sub>, and CaBr<sub>2</sub> (*ca.* 100 equiv. each) (Fig. S20†). We thus conclude that receptor **2** complexes CaBr<sub>2</sub> with high selectivity among the various test bromide salts.

The high selectivity of receptor **2** for MgCl<sub>2</sub> over LiCl and for CaBr<sub>2</sub> over LiBr and MgBr<sub>2</sub> observed in CD<sub>2</sub>Cl<sub>2</sub> was rationalized in terms of the associated binding energies and geometries of the resulting complexes obtained *via* density functional theory (DFT) calculations carried out in the gas phase at the X3LYP/6-31g\* level (Fig. 4; see the ESI† for details). For instance, the complexation energies of receptor **2** for LiCl and MgCl<sub>2</sub> were computed to be  $-57.17$  kcal mol<sup>−1</sup> and  $-71.23$  kcal mol<sup>−1</sup>,

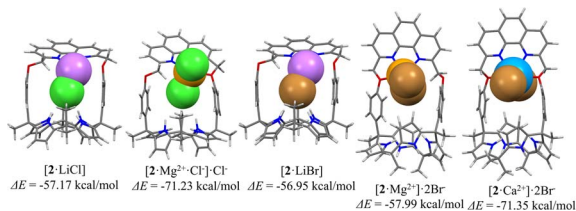


Fig. 4 Optimized structures of the complexes of receptor **2** with LiCl, MgCl<sub>2</sub>, LiBr, MgBr<sub>2</sub>, and CaBr<sub>2</sub> as calculated in the gas phase and the corresponding computed complexation energies for the interaction of receptor **2** with the ion pairs in question.

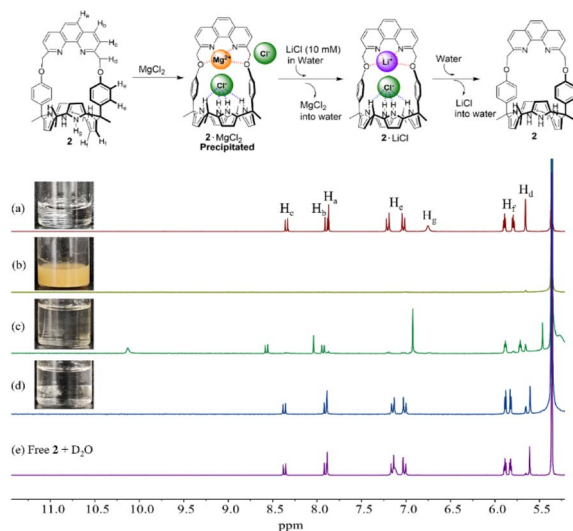


Fig. 5 Partial <sup>1</sup>H NMR spectra of CD<sub>2</sub>Cl<sub>2</sub> solutions of (a) **2** (3 mM) only, (b) **2** + MgCl<sub>2</sub> (500 equiv.), (c) solution (b) after contacting with 10 M LiCl aqueous solution, (d) solution (c) after removing an aqueous phase and then contacting with an ion-free aqueous D<sub>2</sub>O solution, and (e) a CD<sub>2</sub>Cl<sub>2</sub> solution of free **2** after contacting with an ion-free aqueous D<sub>2</sub>O solution. The asterisk (\*) denotes the residual CH<sub>2</sub>Cl<sub>2</sub> peak in the NMR solvent.

respectively. In case of the optimized structure of the MgCl<sub>2</sub> complex, one of two chloride anions was bound to the calix[4]pyrrole subunit of receptor **2** interacting directly with the magnesium cation coordinated by the phenanthroline nitrogen atoms as well as one ether oxygen atom (Fig. 4; *cf.* Fig. S1 and S2†). In contrast, the other chloride anion was located outside the receptor cavity forming a contact ion pair with the magnesium cation. The complexation energy of receptor **2** for MgCl<sub>2</sub> proved larger than that for LiCl by  $-14.06$  and could account for the selective binding of the receptor for MgCl<sub>2</sub> over LiCl; however, solubility considerations are ignored in this analysis. The stabilization energies of receptor **2** upon complexing various bromide salts were similarly calculated to be  $-56.95$  kcal mol<sup>−1</sup> for LiBr,  $-57.99$  kcal mol<sup>−1</sup> for MgBr<sub>2</sub>, and  $-71.35$  kcal mol<sup>−1</sup> for CaBr<sub>2</sub>, respectively (Fig. 5). These computed values could account for the selectivity seen for CaBr<sub>2</sub> over LiBr or MgBr<sub>2</sub>; however, as above, effects such as solvation and solubility are not considered.

The selectivity of receptor **2** seen in 10% CH<sub>3</sub>OH in CDCl<sub>3</sub> is considered to reflect solvation effects. Based on their respective hydration energies ( $\Delta_{\text{hyd}}G^* = -1830$  kJ mol<sup>−1</sup> for Mg<sup>2+</sup>,  $\Delta_{\text{hyd}}G^* = -1505$  kJ mol<sup>−1</sup> for Ca<sup>2+</sup>, and  $\Delta_{\text{hyd}}G^* = -475$  kJ mol<sup>−1</sup> for Li<sup>+</sup>), Mg<sup>2+</sup> and Ca<sup>2+</sup> are presumed to be more strongly solvated by methanol molecules than Li<sup>+</sup>.<sup>29</sup> This strong solvation is thought to reduce the binding interactions with receptor **2** leading to the observed selective binding of LiCl and LiBr in CH<sub>3</sub>OH/CDCl<sub>3</sub> (1/9, v/v). Support for this presumption came from a <sup>1</sup>H NMR spectroscopic analysis. For instance, when receptor **2** was treated with an equimolar mixture of LiCl and MgCl<sub>2</sub> (*ca.* 100 equiv. each) in CD<sub>2</sub>Cl<sub>2</sub>, orange solids corresponding to 2·MgCl<sub>2</sub> precipitated out with no proton signals of the receptor



appearing in the  $^1\text{H}$  NMR spectrum of the residual solvent phase (Fig. S21†). Adding methanol (10% by volume relative to  $\text{CD}_2\text{Cl}_2$ ) to the sample caused the precipitates to dissolve and produced a spectrum analogous to that of  $2 \cdot \text{LiCl}$  (Fig. S21†). The reversal in selectivity seen for receptor 2 from  $\text{MgCl}_2$  to  $\text{LiCl}$  upon moving to a more polar medium is rationalized in terms of  $\text{MgCl}_2$  being more strongly solvated than  $\text{LiCl}$  by methanol. Similarly, the apparent high selectivity of receptor 2 for  $\text{CaBr}_2$  over  $\text{LiBr}$  in  $\text{CD}_2\text{Cl}_2$  was also reversed when methanol (5% by volume) was added to a  $\text{CD}_2\text{Cl}_2$  solution containing precipitated  $2 \cdot \text{CaBr}_2$  (Fig. S22†).

The solvation effects on the receptor selectivity were further supported by two-phase liquid–liquid extraction experiments using  $\text{CD}_2\text{Cl}_2$  as the organic receiving phase and an aqueous  $\text{D}_2\text{O}$  solution as the lithium salt source. For instance, when  $\text{CD}_2\text{Cl}_2$  solutions containing precipitates of the respective  $2 \cdot \text{MgCl}_2$  and  $2 \cdot \text{CaBr}_2$  complexes were contacted with aqueous solutions of  $\text{LiCl}$  (10 M) and  $\text{LiBr}$  (13 M), respectively, complete dissolution ensued (Fig. 5 and S23†). The  $^1\text{H}$  NMR spectra of the organic phases exhibited proton signals consistent with those of the  $\text{LiCl}$  and  $\text{LiBr}$  complexes, respectively (Fig. 5 and S23†). These findings lead us to suggest that  $\text{MgCl}_2$  and  $\text{CaBr}_2$  are released from receptor 2 in the organic phase into the aqueous phases while  $\text{LiCl}$  and  $\text{LiBr}$  initially present in the aqueous phase form complexes with receptor 2 that are soluble in the organic phase. This permits the selective extraction of these two lithium salts from an aqueous source phase into a  $\text{CD}_2\text{Cl}_2$  receiving phase. When the resulting organic phases containing the respective  $\text{LiCl}$  and  $\text{LiBr}$  complexes of receptor 2 were further contacted with ion-free  $\text{D}_2\text{O}$ , release of  $\text{LiCl}$  and  $\text{LiBr}$  into the aqueous  $\text{D}_2\text{O}$  phase occurs. This produces receptor 2 in its ion-free form (Fig. 5 and S23†).

In order to obtain further insights into the ability of receptor 2 to extract the above metal chloride and bromide salts from an aqueous phase, we carried out liquid–liquid extraction experiments using  $\text{CD}_2\text{Cl}_2/\text{D}_2\text{O}$ . For instance, when an  $\text{CD}_2\text{Cl}_2$  layer containing receptor 2 (3 mM) was contacted with aqueous solution layers containing excess  $\text{LiCl}$ ,  $\text{NaCl}$ ,  $\text{KCl}$ ,  $\text{MgCl}_2$ , and  $\text{CaCl}_2$ , respectively, only in the case of  $\text{LiCl}$  were chemical shift changes seen in the  $^1\text{H}$  NMR spectrum of the organic layer consistent with effective extraction (Fig. S24†). The resulting  $^1\text{H}$  NMR spectrum was almost identical to that seen upon complexation of receptor 2 with  $\text{LiCl}$  (cf. Fig. S19†). This finding was taken as evidence that receptor 2 is able to extract  $\text{LiCl}$  with high selectivity from an aqueous source phase into a  $\text{CD}_2\text{Cl}_2$  organic layer. Upon exposure of a  $\text{CD}_2\text{Cl}_2$  solution of receptor 2 (3 mM) to aqueous solutions containing various respective bromide salts, as above, only the lithium salt ( $\text{LiBr}$ ) induced chemical shift changes attributable to  $\text{LiBr}$  complex formation (Fig. S25†).

To evaluate further the extraction capacity of receptor 2 for  $\text{LiCl}$  and  $\text{LiBr}$ , we determined the percentages of the receptor loaded with  $\text{LiCl}$  and  $\text{LiBr}$  in the  $\text{CD}_2\text{Cl}_2$  organic phase after contacting with aqueous solutions containing different concentrations of  $\text{LiCl}$  and  $\text{LiBr}$ , respectively. When receptor 2 was contacted with an aqueous  $\text{D}_2\text{O}$  solution containing 5 M of  $\text{LiCl}$ , two sets of proton signals were visible in the  $^1\text{H}$  NMR spectrum of the organic phase (Fig. S26†). Based on integrations, ca. 24% of

the receptor was presumed to participate in the  $\text{LiCl}$  extraction (Fig. S26†). In contrast, the  $\text{LiCl}$  loading percentage of the receptor from an aqueous solution containing 10 M of  $\text{LiCl}$  was calculated to be ca.  $\approx 100\%$  (Fig. S26†). Further evidence for the ability of receptor 2 to extract  $\text{LiCl}$  came from a high resolution ESI mass spectrometric analysis. A major peak at  $m/z = 795.4105$ , a value corresponding to  $[\text{M} + \text{Li}]^+$ , was seen (Fig. S27†). On the other hand, the corresponding receptor loading levels for  $\text{LiBr}$  were estimated to be ca.  $(70 \pm 10)\%$  and ca. 100% when a  $\text{CD}_2\text{Cl}_2$  solution containing receptor 2 (3 mM) was contacted with aqueous solutions containing 10 M and 15 M concentrations of  $\text{LiBr}$ , respectively (Fig. S28†). These findings stand in sharp contrast to what was seen with receptor 1 that fails to extract  $\text{LiCl}$  or  $\text{LiBr}$  under the same LLE conditions (Fig. S29 and S30†). Both the  $\text{LiCl}$  and  $\text{LiBr}$  complexes of receptor 2 could be separated off and washed with  $\text{D}_2\text{O}$  to release the bound salts into an aqueous  $\text{D}_2\text{O}$  layer (Fig. 5 and S23†) while freeing up receptor 2 for possible reuse.

The extraction efficiency of receptor 2 for  $\text{LiCl}$  and  $\text{LiBr}$  could be improved by employing relatively polar nitrobenzene- $d_5$  as the organic receiving phase instead of  $\text{CD}_2\text{Cl}_2$ . For instance, after a nitrobenzene- $d_5$  solution containing receptor 2 was contacted with an aqueous source phase containing an excess of  $\text{LiCl}$ ,  $\text{NaCl}$ ,  $\text{KCl}$ ,  $\text{MgCl}_2$ , and  $\text{CaCl}_2$ , it was found that 100% of the receptor in the organic phase existed in the form of the  $\text{LiCl}$  complex (Fig. S31†). In the case of  $\text{NaCl}$  and  $\text{KCl}$ , less than 30% of the receptor was loaded with these salts in the organic phase while no evidence for  $\text{MgCl}_2$  and  $\text{CaCl}_2$  extraction by receptor 2 was found (Fig. S31†). In contrast, when a nitrobenzene- $d_5$  organic layer of receptor 2 (3 mM) was contacted with an aqueous layer containing all five test chloride anion salts ( $\text{LiCl}$ ,  $\text{NaCl}$ ,  $\text{KCl}$ ,  $\text{MgCl}_2$ , and  $\text{CaCl}_2$ ) at a concentration of  $\approx 5.0$  M each, the resulting  $^1\text{H}$  NMR spectrum of the nitrobenzene layer was consistent with that recorded after receptor 2 was treated with  $\text{LiCl}$  only (Fig. S31†). This finding was taken as evidence that receptor 2 is capable of extracting  $\text{LiCl}$  with high selectivity from a mixed salt aqueous solution. In analogy to what was seen in the case of the metal chloride salts, receptor 2 was found to extract  $\text{LiBr}$  selectively from an aqueous solution over other test bromide salts. For instance, upon subjecting the nitrobenzene- $d_5$  layer containing receptor 2 to contact with a  $\text{D}_2\text{O}$  layer containing an excess amount of  $\text{LiBr}$ ,  $\text{NaBr}$ ,  $\text{KBr}$ ,  $\text{MgBr}_2$ , and  $\text{CaBr}_2$ , respectively, only  $\text{LiBr}$  gave rise to a  $^1\text{H}$  NMR spectrum of the organic layer consistent with formation of an ion pair complex (Fig. S32†).

We also examined the capacity of receptor 2 to extract  $\text{LiCl}$  and  $\text{LiBr}$  into nitrobenzene- $d_5$  from an aqueous  $\text{D}_2\text{O}$  solution containing various concentrations of  $\text{LiCl}$  and  $\text{LiBr}$ , respectively. For instance, when a nitrobenzene- $d_5$  phase containing receptor 2 (3 mM) was contacted with aqueous  $\text{D}_2\text{O}$  layers containing 0.5–5.0 M of  $\text{LiCl}$ , the resulting  $^1\text{H}$  NMR spectra of the nitrobenzene- $d_5$  phase exhibited two distinguishable sets of proton signals that could be assigned to the ion-free form and the  $\text{LiCl}$  complex of receptor 2, respectively. The  $\text{LiCl}$  loading percentages of the receptor in the organic layer were determined to be <10%, 39%, 63%, 74%, 93%, and 100% when the nitrobenzene- $d_5$  phases containing the receptor were contacted



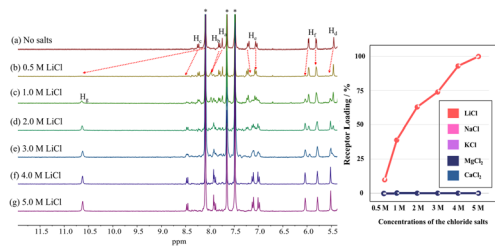


Fig. 6 Left: partial  $^1\text{H}$  NMR spectra of nitrobenzene- $d_5$  layers of **2** (3 mM) recorded upon exposure to aqueous  $\text{D}_2\text{O}$  phases at concentrations of 0.0 M, 0.5 M, 1 M, 2 M, 3 M, 4 M, and 5 M of LiCl, respectively. Percentages of the receptor loaded with LiCl are estimated to be <10%, 39%, 63%, 74%, 93% and 100% respectively. The asterisks denote the residual nitrobenzene peak in the NMR solvent. Right: plots showing loading percentages of receptor **2** for the metal chloride salts after liquid–liquid extraction.

with  $\text{D}_2\text{O}$  solutions containing 0.5 M, 1.0 M, 2.0 M, 3.0 M, 4.0 M, and 5.0 M of LiCl, respectively (Fig. 6). The LiCl loading levels for receptor **2** were compared to those achieved by receptor **1**. For instance, the LiCl loading percentage of receptor **1** from a 10 M LiCl aqueous solution was 15%.<sup>39</sup> In the case of LiBr, when nitrobenzene- $d_5$  solutions of receptor **2** were treated with  $\text{D}_2\text{O}$  aqueous solutions containing LiBr at concentrations of 3.0 M, 5.0 M, and 8.0 M, respectively, 21%, 53%, and 100% of the receptor molecules formed a LiBr complex within the organic phase (Fig. S33<sup>†</sup>).

The ability of receptor **2** to extract LiCl selectively was evaluated under liquid–liquid extraction conditions using aqueous solutions containing other potentially competitive chloride salts including NaCl, KCl,  $\text{MgCl}_2$ , and  $\text{CaCl}_2$ . In the case of the individual salts, receptor **2** failed to extract chloride salts other than LiCl under otherwise identical LLE conditions when aqueous solutions containing 5.0 M of the metal cation chloride anion salts in question were tested (Fig. 6). This selectivity was retained when mixtures of salts were tested. However, unexpectedly, the extraction efficiency of the receptor for LiCl was boosted in the presence of the other test chloride salts. For instance, the proton integration ratios corresponding to the

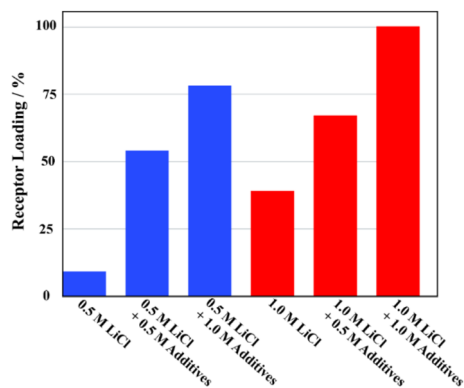


Fig. 7 LiCl loading percentages for receptor **2** after nitrobenzene- $d_5$  organic phases were contacted with aqueous  $\text{D}_2\text{O}$  phases containing 0.5 M and 1.0 M LiCl in the absence and presence of (concurrently) NaCl, KCl,  $\text{MgCl}_2$ , and  $\text{CaCl}_2$  at concentrations of 0.5 M and 1.0 M in each additive.

$2\cdot\text{LiCl}$  complex in the  $^1\text{H}$  NMR spectra of the nitrobenzene- $d_5$  phases increased as the concentration of the other chloride salts increased (Fig. 7 and S34<sup>†</sup>). By way of a specific example, when a nitrobenzene- $d_5$  phase containing receptor **2** (3 mM) was contacted with an aqueous solution containing 0.5 M of LiCl in the absence of the other salts, <10% of the receptor was loaded with LiCl in the organic layer (Fig. 7 and S34<sup>†</sup>). By contrast, the LiCl loading percentages of receptor **2** were enhanced up to 54% and 78% when the aqueous LiCl source phase (0.5 M) consisted of a mixture of NaCl, KCl,  $\text{MgCl}_2$ , and  $\text{CaCl}_2$  at concentrations of 0.5 M each and 1.0 M each, respectively. This loading increased to nearly 100% when each of the salts, including LiCl, was present at 1.0 M (Fig. 7 and S32<sup>†</sup>). Similar results were observed in liquid–liquid extraction studies involving dichloromethane- $d_2$  as the receiving phase. For instance, upon contacting an organic dichloromethane- $d_2$  phase containing receptor **2** with an aqueous  $\text{D}_2\text{O}$  phase containing 5.0 M LiCl, 24% of the receptor in the organic phase was loaded with LiCl. In the presence of excess NaCl and NaBr along with 5.0 M LiCl in the aqueous phase, the LiCl loading percentages improved to 36% and 42%, respectively (Fig. S35 and S36<sup>†</sup>). These findings, which appear analogous to a classic salting-out effect, lead us to suggest that the ion extraction efficiency of other LLE extractants could be improved by employing aqueous source phases containing non-competing salts that are intrinsically insoluble in the organic receiving phase.

## Conclusions

The new ion pair receptor **2** reported here acts as a selective extractant for LiCl and LiBr under solid–liquid and liquid–liquid extraction conditions. Ion pair receptor **2** is composed of a phenanthroline unit of a cation binding site connected *via* ether linkages to a calix[4]pyrrole anion binding site. In dichloromethane- $d_2$ , receptor **2** proved able to complex LiCl, LiBr,  $\text{MgCl}_2$ ,  $\text{MgBr}_2$ , and  $\text{CaBr}_2$ . When treated with the salts in question (*i.e.*,  $\text{LiX}$ ,  $\text{NaX}$ ,  $\text{KX}$ ,  $\text{MgX}_2$ , and  $\text{CaX}_2$  where  $\text{X} = \text{Cl}$  or  $\text{Br}$ ), receptor **2** forms insoluble complexes selectively with  $\text{MgCl}_2$  and  $\text{CaBr}_2$ . This operational selectivity is reversed in more polar protic solvents, such as water or methanol. For instance, upon the *in situ* addition of methanol to  $\text{CD}_2\text{Cl}_2$  solutions containing the presumed  $\text{MgCl}_2$  and  $\text{CaBr}_2$  complexes of receptor **2** along with the other chloride and bromide salts, the bound  $\text{MgCl}_2$  and  $\text{CaBr}_2$  are released from the receptor giving rise to the corresponding LiCl and LiBr complexes. This finding is ascribed to the  $\text{Mg}^{2+}$  and  $\text{Ca}^{2+}$  cations being strongly solvated by polar protic media as compared to the  $\text{Li}^+$  cation. In a moderately polar medium consisting of 10%  $\text{CH}_3\text{OH}$  in  $\text{CDCl}_3$ , receptor **2** binds LiCl and LiBr ion with significantly higher affinity and selectivity than it does other competitive chloride and bromide salts. Receptor **2** was also found to be able to extract LiCl and LiBr efficiently into an organic phase from aqueous sources under two-phase liquid–liquid extraction conditions. This efficiency could be further boosted by adding other chloride salts, including NaCl, KCl,  $\text{MgCl}_2$ , and  $\text{CaCl}_2$  to the aqueous source phase. We thus suggest that receptor **2** may have a role to play in lithium cation separation scenarios.



## Data availability

All data, including synthetic details,  $^1\text{H}$  NMR spectroscopic analyses, DFT calculations, and a single crystal X-ray diffraction analysis of  $2 \cdot \text{LiBr}$  (CCDC 2356046), are available in the ESI.† All data are included either in the main text, the ESI,† or (X-ray work only) uploaded with the Cambridge Crystallographic Data Centre.

## Author contributions

SKK and JLS conceived and supervised the project. NJH synthesized the compounds. NJH and JHO performed ion binding studies using  $^1\text{H}$  NMR spectroscopy. NJH and KL carried out the X-ray diffraction analysis. AL and QH carried out DFT calculations. SKK and JLS wrote the manuscript. All authors contributed to the editing of the manuscript.

## Conflicts of interest

There are no conflicts to declare.

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## Notes and references

- M. Armand and J.-M. Tarascon, *Nature*, 2008, **451**, 652–657.
- R. Oruch, M. A. Elderbi, H. A. Khattab, I. F. Pryme and A. Lund, *Eur. J. Pharmacol.*, 2014, **740**, 464–473.
- E. C. Evarts, *Nature*, 2015, **526**, S93–S95.
- G. Martin, L. Rentsch, M. Höck and M. Bertau, *Energy Storage Mater.*, 2017, **6**, 171–179.
- L. Kavanagh, J. Keohane, G. G. Cabellos, A. Lloyd and J. Cleary, *Resources*, 2018, **7**, 57.
- S. K. Sarai, H. M. Madurui and S. Lippmann, *Innov. Clin. Neurosci.*, 2018, **15**, 30–32.
- U.S. Geological Survey, *Mineral Commodity Summaries 2022*, U.S. Geological Survey, Reston, VA, 2022.
- B. Swain, *Sep. Purif. Technol.*, 2017, **172**, 388–403.
- A. Z. Haddad, L. Hackl, B. Akuzum, G. Pohlman, J.-F. Magnan and R. Kostecki, *Nature*, 2023, **616**, 245–248.
- P. Meshram, B. D. Pandey and T. R. Mankhand, *Hydrometallurgy*, 2014, **150**, 192–208.
- N. Vieceli, C. A. Nogueira, M. F. C. Pereira, A. P. S. Dias, F. O. Durao, C. Guimaraes and F. Margarido, *Miner. Eng.*, 2017, **102**, 1–14.
- D. Yelatontsev and A. Mukhachev, *Hydrometallurgy*, 2021, **201**, 105578.
- T. Gao, N. Fan, W. Chen and T. Dai, *Chin. Geol.*, 2023, **6**, 137–153.
- S. Nishihama, K. Onishi and K. Yoshizuka, *Solvent Extr. Ion Exch.*, 2011, **29**, 421–431.
- P. Loganathan, G. Naidu and S. Vigneswaran, *Environ. Sci.: Water Res. Technol.*, 2017, **3**, 37–53.
- S. Yang, F. Zhang, H. Ding, P. He and H. Zhou, *Joule*, 2018, **2**, 1648–1651.
- G. R. Harvianto, S.-H. Kim and C.-S. Ju, *Rare Met.*, 2016, **35**, 948–953.
- C. Liu, Y. Li, D. Lin, P.-C. Hsu, B. Liu, G. Yan, T. Wu, Y. Cui and S. Chu, *Joule*, 2020, **4**, 1459–1469.
- X. He, S. Kaur and R. Kostecki, *Joule*, 2020, **4**, 1357–1368.
- V. Flexer, C. F. Baspineiro and C. I. Galli, *Sci. Total Environ.*, 2018, **639**, 1188–1204.
- G. Liu, Z. Zhao and A. Ghahreman, *Hydrometallurgy*, 2019, **187**, 81–100.
- P. Xu, J. Hong, X. Qian, Z. Xu, H. Xia, X. Tao, Z. Xu and Q.-Q. Ni, *J. Mater. Sci.*, 2021, **56**, 16–63.
- Y. Sun, Q. Wang, Y. Wang, R. Yun and X. Xiang, *Sep. Purif. Technol.*, 2021, **256**, 117807.
- X. Zhao, M. Feng, Y. Jiao, Y. Zhang, Y. Wang and Z. Sha, *Desalination*, 2020, **481**, 114360.
- A. Khalil, S. Mohammed, R. Hashaikeh and N. Hilal, *Desalination*, 2022, **528**, 115611.
- J. Ying, Y. Lin, Y. Zhang and J. Yu, *ACS ES&T Water*, 2023, **3**, 1720–1739.
- W. T. Stringfellow and P. F. Dobson, *Energies*, 2021, **14**, 6805.
- M. L. Vera, W. R. Torres, C. I. Galli, A. Chagnes and V. Flexer, *Nat. Rev. Earth Environ.*, 2023, **4**, 149–165.
- Y. Marcus, *J. Chem. Soc., Faraday Trans.*, 1991, **87**, 2995–2999.
- P. Xu, J. Hong, X. Qian, Z. Xu, H. Xia, X. Tao, Z. Xu and Q.-Q. Ni, *J. Mater. Sci.*, 2021, **56**, 16–63.
- J. Zhang, N. Tanjedrew, M. Wenzel, P. Royola, H. Du, S. Kiatischevi, L. F. Lindoy and J. J. Weigand, *Angew. Chem., Int. Ed.*, 2023, **62**, e202216011.
- S. K. Kim and J. L. Sessler, *Chem. Soc. Rev.*, 2010, **39**, 3784–3809.
- S. K. Kim and J. L. Sessler, *Acc. Chem. Res.*, 2014, **47**, 2525–2536.
- Q. He, G. I. Vargas-Zuniga, S. H. Kim, S. K. Kim and J. L. Sessler, *Chem. Rev.*, 2019, **119**, 9753–9835.
- A. J. McConnell and P. D. Beer, *Angew. Chem., Int. Ed.*, 2012, **51**, 5052–5061.
- A. J. McConnell, A. Docker and P. D. Beer, *ChemPlusChem*, 2020, **85**, 1824–1841.
- Q. He, Z. Zhang, J. T. Brewster, V. M. Lynch, S. K. Kim and J. L. Sessler, *J. Am. Chem. Soc.*, 2016, **138**, 9779–9782.
- K. I. Hong, H. Kim, Y. Kim, M. G. Choi and W. D. Jang, *Chem. Commun.*, 2020, **56**, 10541–10544.
- Q. He, N. J. Williams, J. H. Oh, V. M. Lynch, S. K. Kim, B. A. Moyer and J. L. Sessler, *Angew. Chem., Int. Ed.*, 2018, **57**, 11924–11928.
- K. Wang, C.-C. Yee and H. Y. Au-Yeung, *Chem. Sci.*, 2016, **7**, 2787–2792.
- J. Yoo, I.-W. Park, T.-Y. Kim and C.-H. Lee, *Bull. Korean Chem. Soc.*, 2010, **31**, 630–634.
- K. A. Connors, *Binding Constants: The Measurement of Molecular Complex Stability*, Wiley-Interscience, New York, 1987.
- Association constants ( $K_a$ ) were evaluated using BindFit v0.5, available from, <https://app.supramolecular.org/bindfit/>.

