Transmembrane Fluoride Transport: Direct Measurement and Selectivity Studies

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ABSTRACT: Fluoride has been overlooked as a target in the development of synthetic anion transporters despite natural fluoride transport channels being recently discovered. In this paper we report the direct measurement of fluoride transport across lipid bilayers facilitated by a series of strapped calix[4]pyrroles and show that these compounds facilitate transport via an electrogenic mechanism (determined using valinomycin and monensin coupled transport assays and an additional osmotic response assay). An HPTS transport assay was used to quantify this electrogenic process and assess the interference of naturally occurring fatty acids with the transport process and Cl̅ over H+/OH̅ transport selectivity.

INTRODUCTION

Transmembrane ion transport facilitated by protein channels is essential in maintaining ion homeostasis. In the past decade, the development of synthetic anion transporters has attracted much attention, with a variety of small molecules being developed to facilitate the transport of biologically relevant anions such as chloride, bicarbonate, sulfate and sulfate. These compounds may find future application in the treatment of diseases caused by faulty ion channels, such as cystic fibrosis.

Fluoride is used as an additive in water and oral hygiene products and is found at levels of 10–100 μM in the aqueous environment. Exposure to high levels of fluoride (above 1.5 mg/L) can be detrimental to health, leading to fluorosis, which affects teeth and bones. To the best of our knowledge, no synthetic compound has been shown to facilitate fluoride transport across lipid bilayers. Fluoride is small, hard, and basic, with a higher hydration energy than chloride and bicarbonate, making fluoride complexation in water a challenging problem. In organic environments, deprotonation of anion receptors by fluoride-driven by the formation of bifluoride, can be a competing process to fluoride complexation. The use of Lewis acidic metal centers by Gabbai and co-workers in fluoride sensors has allowed fluoride detection in water below the maximum contaminant level (4 ppm) set by the Environment Protection Agency. Additionally, some Lewis acid-based compounds have been found to act as fluoride anionophores across artificial PVC-DOS membranes within the field of ion-selective electrodes (ISEs) and fluoride sensing using boronic acid derivatives has been shown to be a successful strategy by Shinkai, James, and others. In early work by Sessler and co-workers, fluoride transport through a liquid membrane system was facilitated by a protonated sapphyrin within a U-tube. The transmembrane exchange of fluoride facilitated by the protein capnophorin in red blood cells has also been investigated utilizing 19F NMR techniques. Other interesting work by Matile has shown promising indirect results for fluoride as a potential target for lipid bilayer transport and provides a strong precedent that this transport is achievable.

Over the past 4 years, seminal work by Miller et al. has revealed the existence of two naturally occurring, phylogenetically unrelated families of F̅ exporter anion channels, CLC F̅/H+ antiporters, and small membrane channel “Fluc” proteins, the former found solely in bacterial cells, while the Fluc proteins are widespread among unicellular organisms as well as eukaryotes. These channels possess strong selectivity for F̅ over Cl̅, and crystal structures of a Fluc channel from Bordetella pertussis reveal a surprising “double-barreled” channel architecture in which selectivity is achieved by the small bore of the channel.

These channels reduce bacterial fluoride toxicity, which occurs due to the weak acid accumulation effect, by exporting fluoride to maintain the cytoplasmic fluoride concentration below enzyme inhibitory levels.

We therefore decided to investigate whether it was possible to facilitate the transport of fluoride through lipid bilayers using...
small-molecule anion carriers and determine whether such transport could be achieved selectively over chloride transport. Herein we report the first instance of directly measured small-molecule transmembrane fluoride transport with a series of strapped calix[4]pyrroles 2−4.49 We have recently found that anion transporters can bind and facilitate the flip-flop of fatty acids across lipid bilayer membranes.50 We therefore also decided to investigate the interaction of these compounds both with acetate in solution and with fatty acids in lipid bilayers.

**RESULTS AND DISCUSSION**

We investigated whether strapped calix[4]pyrroles could be used to mediate fluoride transport. Calix[4]pyrroles are less acidic than the urea and thiourea transporter motifs that our group, and others, have recently explored and are therefore less likely to be deprotonated within the lipid bilayer.23 Calix[4]pyrroles 1−4 (Figure 1) were synthesized following literature procedures,49 and their F$^−$ binding affinity was evaluated using $^1$H NMR titration methods and isothermal titration calorimetry (ITC). A variety of anion transport assays were employed to fully characterize the fluoride transport processes facilitated by 1−4.51

**Anion Binding Studies and Solid-State Analysis.**

Proton NMR titration studies in DMSO-$d_6$/0.5% H$_2$O with tetrabutylammonium fluoride (TBAF) and compounds 1−4 resulted in slow exchange of the fluoride complex and the free calix[4]pyrrole on the NMR time scale. A downfield shift of the pyrrole NH resonances and the triazole CH resonance and coupling of the NH resonance to the $^{19}$F nucleus of the fluoride anion were observed, which are usually indicative of strong hydrogen bonding. Slow exchange was observed previously with tetrabutylammonium chloride (TBACl) and compounds 2−4.49 Slow exchange was also observed upon addition of tetrabutylammonium acetate with compounds 2−4, and a stability constant of 6060 M$^{-1}$ was derived from global fitting analysis using bindfit52,53 with compound 1 in DMSO-$d_6$/0.5% H$_2$O. An interesting splitting effect for the pyrrole NH was observed for the titration with 2 with acetate, suggesting two environments, which we attribute to the OAc$^−$ not binding symmetrically within the smaller cavity of compound 2. NMR spectra and species ratio plots can be found in the Supporting Information. Quantification of the binding affinity was performed using ITC in acetonitrile. The results, summarized in Table 1, show that compounds 1−4 exhibit exothermic complexation of both salts with slightly unfavorable entropic contributions for all, except 1 and 4, with tetrabutylammonium fluoride. Receptors 1−4 bind strongly to fluoride, with $K_a \geq 10^6$ M$^{-1}$ with no indications of deprotonation, supporting their suitability for fluoride transport. This strong binding for 1−4 is similar to that seen for tetrabutylammonium chloride binding, with previous literature

Table 1. Overview of the Association Constants and Thermodynamic State Functions for the 1:1 Complexation of Tetrabutylammonium Salts with 1−4 As Measured in Acetonitrile by ITC at 303 K

| receptor | $K_a$ (M$^{-1}$) | $\Delta G$ (kJ mol$^{-1}$) | $\Delta H$ (kJ mol$^{-1}$) | $T\Delta S$ (kJ mol$^{-1}$) | $K_a$ (M$^{-1}$) | $\Delta G$ (kJ mol$^{-1}$) | $\Delta H$ (kJ mol$^{-1}$) | $T\Delta S$ (kJ mol$^{-1}$) |
|----------|----------------|-----------------------------|-----------------------------|-----------------------------|----------------|-----------------------------|-----------------------------|-----------------------------|
| 1        | 3.5 $\times$ 10$^6$ | −38.0                      | −30.1                      | 7.9                        | 5.2 $\times$ 10$^5$ | −33.2                      | −45.3                      | −12.1                      |
| 2        | 1.3 $\times$ 10$^6$ | −35.5                      | −44.5                      | −9.0                       | 1.3 $\times$ 10$^5$ | −29.7                      | −41.4                      | −11.7                      |
| 3        | 1.1 $\times$ 10$^7$ | −40.8                      | −44.6                      | −3.8                       | 1.7 $\times$ 10$^6$ | −36.3                      | −51.1                      | −14.8                      |
| 4        | 1.9 $\times$ 10$^7$ | −42.2                      | −34.5                      | 7.7                        | 1.2 $\times$ 10$^6$ | −35.4                      | −42.7                      | −7.3                       |

Figure 1. Structures of the calix[4]pyrroles 1−4 used in this study.

Figure 2. (a) Single-crystal X-ray structure of 2·TBAF complex (TBA not shown for clarity). (b) Single-crystal X-ray structure of 2·TBACl complex (TBA not shown for clarity). (c) Single-crystal X-ray structure of 3·CsF complex (ethanol solvent not shown for clarity). N···F, N···Cl, C···F, and C···Cl distances for all structures are displayed as red dashed lines, with measurements (Å) in black. Full crystallographic data can be found in the Supporting Information.
reporting a $K_a$ range of $10^5$–$10^6$ M$^{-1}$ in acetonitrile. Acetate binding was slightly weaker, with a $K_a$ range of $10^5$–$10^6$ M$^{-1}$ (Table 1). As expected, acetate binding was stronger for 1, 3, and 4 compared to 2, which may be due to the short strap constraining the binding cavity size. ITC binding isotherms can be found in the Supporting Information. 

Solid-state structures of 2 complexed with $F^-$ and $Cl^-$ (Figure 2a,b) show the anion bound within the strapped cavity, with shorter N···F distances of 2.747(3)–2.787(3) Å observed compared to N···Cl (3.247(2)–3.271(2) Å) and an additional stabilizing effect from the triazole CH for 2-TBAF. Hence, the fluoride appears to be bound more strongly, at least
in the solid state. Figure 2c shows the crystal structure of the cesium fluoride complex of 3, which is similar to the cesium chloride complex of the parent macrocycle 1, with N–F distances of 2.775(3)–2.828(4) Å and the cesium sitting in the calixpyrrole cup-shaped cavity formed by the pyrrole rings.

**Anion Transport Studies.** The fluoride transport activities of 1–4 were probed using the vesicle-based ISE assay shown in Figure 3a(i), which was adapted from the literature. Large unilamellar vesicles (LUVs, 200 nm) composed of 1-palmitoyl-2-oleoyl-sn-glycero-3-phosphocholine (POPC) were loaded with KF (300 mM) and suspended in potassium gluconate (KGlc, 300 mM). All solutions were buffered to pH 7.2 with HEPES (10 mM). Transport was initiated by addition of valinomycin (Vln), followed by the anionophore 30 s later, and fluoride efflux was measured directly using a chloride-selective electrode. The driving force in this assay is the large fluoride gradient. In the absence of valinomycin and an anion transporter, fluoride efflux is minimal. Fluoride efflux can occur only in the presence of valinomycin, which allows for the dissipation of the membrane potential that would occur if the fluoride transport facilitated by an anion transporter occurred alone.

The results for this assay are shown in Figure 3b. Compounds 2–4 were all found to function as fluoride transporters at 20 μM, 2 mol% (with respect to lipid concentration), with 3 being the most effective, while compound 1 showed no transport activity under these conditions. Following this encouraging result, the assay was repeated in the absence of Vln with POPC vesicles loaded with MF (M = Na, K, Cs, Rb) to assess the result, the assay was repeated in the absence of Vln with POPC vesicles loaded with MF (M = Na, K, Cs, Rb) to assess the transport experiment, all the receptors function as CsF co-transporters (and not NaF, KF, RbF), with compound 4 performing best.

Previous studies have shown that calix[4]pyrroles have high affinity for cesium halides. Small halide ions have been shown to organize 1 into a cone conformation, providing an electron-rich cup which binds the large charge-dia
ced receptor F−/OH− from 30% (as illustrated in Figure 3e for 4) (the data for the other compounds are shown in the Supporting Information). These results support the hypothesis that 2–4 function via an electrogenic transport mechanism. An osmotic response assay was performed to provide further evidence in support of the electrogenic transport mechanism, and to provide supporting evidence to rule out H+/OH− transport, which could cause background HF transport, a control assay was performed with KOAc-loaded vesicles. The acetate ion alone is membrane impermeable (unless transported), however, protonation of acetate resulting from transport of H+ or OH−, potentially facilitated by 2–4, may form membrane-permeable acetic acid. Figure 4b shows the results for 3 and 4. No permeabilization of acetate was seen for 1–3, which is evidence in support of the absence of H+ and OH− transport. Conversely, 4 shows a positive result for the acetate control assay, which was attributed to the larger strapped cavity allowing for acetate binding and transport, and not H+ or OH− transport. This is supported by the strong affinity of this receptor for OAc− from the ITC results and the fact the transport mechanism cannot couple to monensin.

Additionally, we quantified the Cl−/H+/OH− selectivity of the strapped calixpyrroles using a modified NMDG-Cl assay, which involves the use of gramicidin D (Gra), a naturally occurring proton channel, and oleic acid (OA), a naturally occurring monounsaturated omega-9 fatty acid. POPC LUVs (200 nm) were loaded with 8-hydroxyquinoline-1,3,6-trisulfonic acid (HPTS, a pH-sensitive dye, 1 mM) and N-methyl-D-glucamine

**Table 2. Transport Activity from the Fluoride and Chloride ISE Assays with Fluoride Selectivity Factors (Sf), from Hill Analyses**

| receptor | EC50 μM (mol%) | Sf |
|----------|---------------|---|
| F− | Cl− | F− | Cl− |
| 1 | 1 (1.3) | 24 (2.4) | 1.8 |
| 2 | 9 (0.9) | 16 (1.6) | 1.8 |
| 3 | 12 (1.2) | 5 (0.5) | 0.4 |

EC50 at 270 s of receptor with respect to lipid. Fluoride selectivity determined from EC50 for chloride transport divided by EC50 for fluoride transport. Sf > 1 indicates fluoride selectivity. No transport ability in this assay; no Hill analysis results or selectivity factors.
chloride (NMDG-Cl, 100 mM) and suspended in NMDG-Cl (100 mM). All solutions were buffered to pH 7.0 with HEPES (10 mM). The anionophore was added, followed by a base pulse (NMDG 5 mM) to generate a pH gradient, and the dissipation of this gradient by the anionophore was monitored by following the HPTS fluorescence ratio. This assay measures the receptor’s electroneutral H+/Cl\(^{-}\) co-transport (Cl\(^{-}\)/OH\(^{-}\) antiport) ability. If the receptor functions as a selective electrogenic Cl\(^{-}\) transporter, as found for 2–4, the H\(^{+}\)/OH\(^{-}\) transport will be the rate-limiting factor for transport. Addition of the proton channel Gra should accelerate the H\(^{+}\) transport, removing this rate-limiting effect.

Quantification of the transport in both cases (with and without the proton channel) permits selectivity for chloride over proton or hydroxide transport to be calculated. Addition of oleic acid allows evaluation of the chloride selectivity of the receptors over the ability to contribute to the naturally occurring fatty acid proton transport pathway by assisting flip-flop of fatty acid anions described in Figure 5a.b. Figure 5a shows how a receptor capable of complexation to the deprotonated headgroup of a fatty acid can artificially enhance the natural proton shuttling seen with fatty acids in nature. Receptor masking of the charged headgroup accelerates movement of the charged form of the headgroup across the membrane. Figure 5b(iii) shows this effect within the NMDG assay, with overall H+/Cl\(^{-}\) symport being observed. Experimentally, this effect can artificially accelerate the pH gradient dissipation with the fatty acid flip-flop process, hence, the Cl\(^{-}\) > H\(^{+}\)/OH\(^{-}\) selectivity will decrease in the presence of fatty acids within a membrane if the receptor has some affinity for the fatty acid headgroup. Since the presence of free (unesterified) fatty acids is ubiquitous in cellular membranes, this is important/undesirable in our ongoing effort in the development of Cl\(^{-}\) > H\(^{+}\)/OH\(^{-}\)-selective synthetic anionophores for potential therapeutic application against cystic fibrosis.

Dose–response experiments and Hill analyses were performed in the absence and presence of gramicidin and oleic acid to gain EC\(_{50}\) values. The ratio of these EC\(_{50}\) values was...
used to quantify the chloride selectivity, giving selectivity factors ($S_C$ and $S_{OA/G}$) presented in Table 3. It is important to note that these selectivity factors are relative to the $\text{Cl}^-$ concentration used within the assay.51 Receptors 2–4 show good selectivity for chloride along the series, with an $S_C$ of 18.2, 10.3, and 13.5, respectively; 2, the shortest strapped, smallest cavity receptor possessed the highest selectivity for chloride, which supports previous findings that a higher degree of anion encapsulation enforces a higher degree of anion desolvation and hence better selectivity for chloride transport.51 Figure 6 shows the $EC_{50}$ values in a bar chart to help visualize this selectivity. In the presence of oleic acid, the effect of artificially enhanced $H^+/\text{Cl}^-$ symport due to the fatty acid flip-flop side process is negligible in 2 and 3; however, there is a mild increase in the transport ability of 4 by a factor of 1.7. This result correlates to the findings in the osmotic assay and ITC study that 4 is likely to bind and transport acetate, as it is more inclined to bind the oxoanionic oleic acid headgroup (Figure S) to enhance its proton shuttle effect and therefore accelerate the dissipation of the pH gradient.63 This resulted in a loss of $\text{Cl}^- > H^+/\text{OH}^-$ selectivity ($S_{OA/G}$) for 4 by a difference of 5.5; while there is a small loss for 3, the selectivity of the shortest strapped 2 was not compromised in the presence of fatty acid. These results reiterated the advantageous effect of encapsulation and enforced steric hindrance in the design of synthetic anion transporters for overall $\text{Cl}^->H^+/\text{OH}^-$ selectivity.

**CONCLUSIONS**

Using a fluoride ion-selective electrode assay, we have shown the first example of a direct measurement of small synthetic molecule facilitated, transmembrane fluoride transport via an electrogenic anion transport mechanism, mimicking an important natural transport process that occurs in bacteria. The shorter strapped receptors 2 and 3 are moderately selective for fluoride over chloride in the Vln coupled assay. Additionally, we have quantified the $\text{Cl}^- > H^+/\text{OH}^-$ selectivity of 2–4, finding them highly selective for $\text{Cl}^-$ over $H^+/\text{OH}^-$, which supports previous findings that encapsulation is the key to selectivity.53 We also evaluated how the presence of a naturally occurring fatty acid interferes with the receptor’s transport mechanism and compromises the overall $\text{Cl}^- > H^+/\text{OH}^-$ selectivity, confirming the need for consideration of fatty acid affinity when designing transporters for physiological purposes. Remarkably, the shortest strapped calixpyrrole 2 not only demonstrated the best $\text{Cl}^- > H^+/\text{OH}^-$ selectivity but also fully maintained the selectivity in the presence of fatty acid. These results illustrate the importance of encapsulation and point the way forward in the design of halide-selective transporters.

**ASSOCIATED CONTENT**

* Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/jacs.6b10694.

Details of general procedures, NMR binding studies, ITC, single-crystal X-ray studies for 2 and 3, and all anion transport experimental details for all the vesicle-based assays used in this study (PDF)

Crystallographic data for 2 (CIF)

Crystallographic data for 2-TBACl (CIF)

Crystallographic data for 2-TBAF (CIF)

Crystallographic data for 3-CsF (CIF)

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Notes

The authors declare no competing financial interest. The data from this study have been made available at http://dx.doi.org/10.5258/SOTON/403025 to comply with the EPSRC open data policy.

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