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## Dissecting the chloride–nitrate anion transport assay†

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**A systematic study of chloride vs. nitrate selectivity across six anion transporters has revealed a good correlation between the selectivities of their anion binding and membrane transport properties. This work reveals the limitations of the chloride–nitrate exchange assay and shows how new approaches can be used to measure anion uniport.**

The design of synthetic anion receptors<sup>1</sup> that can carry biological anions, most importantly chloride, across phospholipid bilayers has been an active area of research in supramolecular chemistry.<sup>2</sup> These compounds have future therapeutic potential to replace the function of faulty anion channels in genetic diseases,<sup>3</sup> or to disrupt the ionic and pH gradients in cancer cells.<sup>4</sup> Efforts have been made to improve the potency of anion transporters, allowing them to function at low concentrations as required for therapeutic applications. Typically, chloride–nitrate exchange (antiport) assays have been used to evaluate the anion transport potency of the transporters.<sup>5</sup> In this assay, it is assumed that nitrate transport is unlikely to be the rate-limiting process due to the anion's high lipophilicity, and therefore this assay is assumed to indicate the chloride transport activity of the transporters.<sup>6</sup> However, in anion binding studies, chloride is almost always more strongly bound to hydrogen bond donor anion receptors than nitrate due to the higher charge density of Cl<sup>−</sup>.<sup>7</sup> This could result in faster chloride transport than nitrate transport. The study of Cl<sup>−</sup> vs. NO<sub>3</sub><sup>−</sup> selectivity in membrane transport and its correlation with binding selectivity is thus of fundamental importance in addressing the question whether anion transport selectivity is dominated by anion lipophilicity<sup>8</sup> or anion affinity.<sup>9</sup> This information is also practically useful to unravel the potential limitations of the chloride–nitrate exchange assay. Recently, we have developed membrane transport

assays that could measure the rate of anion uniport mediated by anion transporters without the need for an anion exchange process to occur.<sup>10</sup> We here make use of two complementary vesicle-based assays to determine anion transport selectivity, in particular Cl<sup>−</sup>/NO<sub>3</sub><sup>−</sup> selectivity, of a library of hydrogen bond-based anion transporters **1–6** that contain increasing numbers of hydrogen bond donors (Fig. 1). By comparing these results with association constants for anion complexation determined in acetonitrile, we demonstrate for the first time a strong correlation between binding selectivity and transport selectivity across a series of structurally diverse anion transporters.

Compounds **1**, **2**, **3**, **5**, and **6** are examples of anion transporters previously studied by our group.<sup>9,10b,11</sup> These compounds represent two distinct design approaches to highly effective anion transporters. Compounds **1–3** contain highly acidic NH groups, leading to high anion binding affinity and the ability to disperse the negative charge of the bound anion. Compounds **5** and **6** are without the electron-withdrawing CF<sub>3</sub> groups but contain more hydrogen bond donors that are favourable for binding and transport because of multivalency<sup>12</sup> and encapsulation.<sup>13</sup> Their protonated cationic forms do not participate in anion transport as demonstrated previously.<sup>10b</sup> We synthesised a new receptor **4** as a dipodal control of the tripodal thiourea **5**. The affinities of these receptors towards anions including Cl<sup>−</sup>, Br<sup>−</sup> and NO<sub>3</sub><sup>−</sup> in acetonitrile were determined by UV-vis absorption titrations using tetrabutylammonium (TBA<sup>+</sup>) salts of the anions

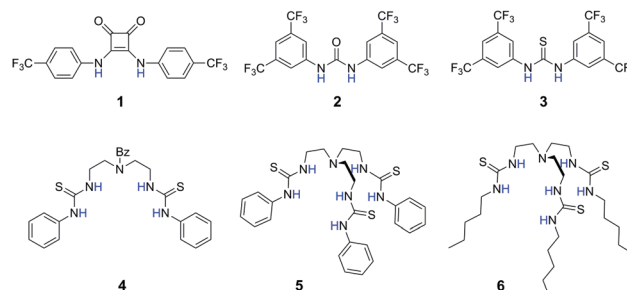


Fig. 1 Structures of anion transporters **1–6**.

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Table 1 Anion binding and transport data for compounds 1–6

Comp.	Anion binding in acetonitrile <sup>a</sup>			Anion transport in lipid bilayers						
	$K_a(\text{Cl}^-)/\text{M}^{-1}$	$K_a(\text{NO}_3^-)/\text{M}^{-1}$	$S_b(\text{Cl}^-/\text{NO}_3^-)$	ISE exchange assay <sup>b</sup>	HPTS assay <sup>e</sup>			Osmotic assay <sup>h</sup>		
				$\text{EC}_{50}^{\text{ISE}}(\text{Cl}^-/\text{NO}_3^-)$ exchange/mol%	$\text{EC}_{50}^{\text{HPTS}}(\text{Cl}^-)/\text{mol}\%$	$\text{EC}_{50}^{\text{HPTS}}(\text{NO}_3^-)/\text{mol}\%$	$S_t^{\text{HPTS}}(\text{Cl}^-/\text{NO}_3^-)$	$\text{EC}_{50}^{\text{Os}}(\text{Cl}^-)/\text{mol}\%$	$\text{EC}_{50}^{\text{Os}}(\text{NO}_3^-)/\text{mol}\%$	$S_t^{\text{Os}}(\text{Cl}^-/\text{NO}_3^-)$
1	$8.2 \times 10^5$	$2.5 \times 10^3$	330	0.060 <sup>c</sup>	0.0089 <sup>f</sup>	0.0073 <sup>f</sup>	0.82	0.093	0.078	0.85
2	$1.8 \times 10^4$	$1.1 \times 10^3$	17	0.30 <sup>c</sup>	0.043 <sup>f</sup>	0.0052 <sup>f</sup>	0.12	0.27	0.069	0.26
3	$2.3 \times 10^4$	$6.0 \times 10^2$	38	0.16 <sup>c</sup>	0.013 <sup>f</sup>	0.0014 <sup>f</sup>	0.11	0.12	0.031	0.26
4	$3.0 \times 10^4$	$2.1 \times 10^2$	140	> 5	0.49 <sup>g</sup>	0.045 <sup>g</sup>	0.091	3.3	0.92	0.28
5	$1.7 \times 10^6$	$2.7 \times 10^2$	6200	0.31 <sup>d</sup>	0.0044 <sup>g</sup>	0.036 <sup>g</sup>	8.1	0.055	0.56	10
6	$8.3 \times 10^5$	$4.9 \times 10^2$	1700	0.11	0.0034 <sup>g</sup>	0.010 <sup>g</sup>	3.0	0.037	0.18	4.9

<sup>a</sup> Association constants determined by UV-vis titration in acetonitrile at 298 K, using tetrabutylammonium anion salts. Errors were found to be <10%.  $\text{Cl}^-/\text{NO}_3^-$  binding selectivity  $S_b(\text{Cl}^-/\text{NO}_3^-) = K_a(\text{Cl}^-)/K_a(\text{NO}_3^-)$ . <sup>b</sup>  $\text{Cl}^-/\text{NO}_3^-$  exchange assay schematically shown in Fig. 2a, using POPC LUVs with a mean diameter of ~200 nm.  $\text{EC}_{50}$  calculated at 270 s. <sup>c</sup> Reported in ref. 9. <sup>d</sup> Reported in ref. 11a. <sup>e</sup>  $\text{H}^+/\text{Cl}^-$  cotransport assay schematically shown in Fig. 2b, using POPC LUVs with a mean diameter of ~200 nm.  $\text{EC}_{50}$  calculated at 200 s.  $\text{Cl}^-/\text{NO}_3^-$  transport selectivity in this assay:  $S_t^{\text{HPTS}}(\text{Cl}^-/\text{NO}_3^-) = \text{EC}_{50}^{\text{HPTS}}(\text{NO}_3^-)/\text{EC}_{50}^{\text{HPTS}}(\text{Cl}^-)$ . <sup>f</sup> Determined in the absence of proton channel gramicidin D. The addition of gramicidin D did not affect the overall transport rate because these compounds are by themselves good  $\text{H}^+/\text{OH}^-$  transporters. See ref. 10b. <sup>g</sup> Determined in the presence of gramicidin D to remove the need of anion transporters to facilitate  $\text{H}^+/\text{OH}^-$  transport. Without gramicidin,  $\text{H}^+/\text{OH}^-$  transport facilitated by these compounds is slow and would rate-limit the overall transport process. See ref. 10b. <sup>h</sup>  $\text{K}^+/\text{Cl}^-$  cotransport assay schematically shown in Fig. 2c, using POPC LUVs with a mean diameter of ~400 nm.  $\text{EC}_{50}$  calculated at 600 s.  $\text{Cl}^-/\text{NO}_3^-$  transport selectivity in this assay:  $S_t^{\text{Os}}(\text{Cl}^-/\text{NO}_3^-) = \text{EC}_{50}^{\text{Os}}(\text{NO}_3^-)/\text{EC}_{50}^{\text{Os}}(\text{Cl}^-)$ .

(see Table 1 for  $\text{Cl}^-$  and  $\text{NO}_3^-$  binding data, and Table S1 in ESI† for  $\text{Br}^-$  binding data). For all compounds, the anion binding selectivity trend was found to follow the anion charge density ( $\text{Cl}^- > \text{Br}^- > \text{NO}_3^-$ ). Despite the same anion binding selectivity sequence, the actual extent of  $\text{Cl}^-/\text{NO}_3^-$  binding selectivity, i.e. the  $S_b(\text{Cl}^-/\text{NO}_3^-)$  value, did show significant variation among different scaffolds. The two tripodal compounds 5 and 6 have very high (>1500-fold) binding selectivities for  $\text{Cl}^-$  over  $\text{NO}_3^-$ , whereas the other ureas and thioureas have a selectivity value of less than 150. A similar trend was found for the  $\text{Br}^-/\text{NO}_3^-$  binding selectivity (Table S1, ESI†). Compared with the dipodal thiourea 4, the additional thiourea arm in 5 dramatically enhances the binding of  $\text{Cl}^-$  and  $\text{Br}^-$  but does not improve  $\text{NO}_3^-$  binding as much. These binding results give a hint that the tripodal compounds could behave differently in anion transport selectivity compared with other compounds.

All compounds have been initially subject to a traditional  $\text{Cl}^-/\text{NO}_3^-$  exchange assay (Fig. 2a). Briefly, 1-palmitoyl-2-oleoyl-*sn*-glycero-3-phosphocholine (POPC) large unilamellar vesicles (LUVs) were loaded with NaCl (490 mM) buffered at pH 7.2, and suspended in  $\text{NaNO}_3$  (490 mM) buffered at pH 7.2. The anion transporter was added to the vesicle suspension as a DMSO solution. Chloride efflux (due to  $\text{Cl}^-/\text{NO}_3^-$  exchange induced by the anion transporters) was monitored *via* appearance of  $\text{Cl}^-$  in the external solution, measured by a chloride ion-selective

electrode (ISE). The activity of a transporter was quantified by an effective concentration of the transporter to reach 50% of ion transport ( $\text{EC}_{50}$  value) at 270 s. In this assay, the simple squaramide 1 was the champion while the two tripodal thioureas 5 and 6 significantly fell behind (Table 1).

As the above-mentioned exchange assay provides no information on  $\text{Cl}^-/\text{NO}_3^-$  selectivity, we conducted two additional assays to directly measure anion selectivity. In the first assay (an HPTS assay, Fig. 2b), POPC LUVs with a mean diameter of 200 nm were loaded with and suspended in a solution of the *N*-methyl-*D*-glucamine salt of the anion of interest (NMDG-X, 100 mM,  $\text{X}^- = \text{Cl}^-$  or  $\text{NO}_3^-$ ) buffered at pH 7.0 with HEPES. The external pH was brought to ~8 by addition of a base pulse (5 mM of *N*-methyl-*D*-glucamine), and then the transporter-induced dissipation of the pH gradient across vesicle membranes was measured using an intravesicular fluorescent pH indicator 8-hydroxypyrene-1,3,6-trisulfonic acid (HPTS). The overall process leading to pH gradient dissipation is  $\text{H}^+/\text{X}^-$  symport or  $\text{OH}^-/\text{X}^-$  antiport, which may be rate-limited by  $\text{H}^+$  or  $\text{OH}^-$  transport. However the need for an anion transporter to facilitate  $\text{H}^+$  or  $\text{OH}^-$  transport can be eliminated by using the proton channel gramicidin, <sup>10b</sup> allowing this assay to reveal the ability of an anion transporter to facilitate  $\text{X}^-$  uniport (Fig. 2b).<sup>14</sup> In the second assay (osmotic assay), larger POPC LUVs (mean diameter 400 nm) were loaded with a buffered KX (300 mM) solution

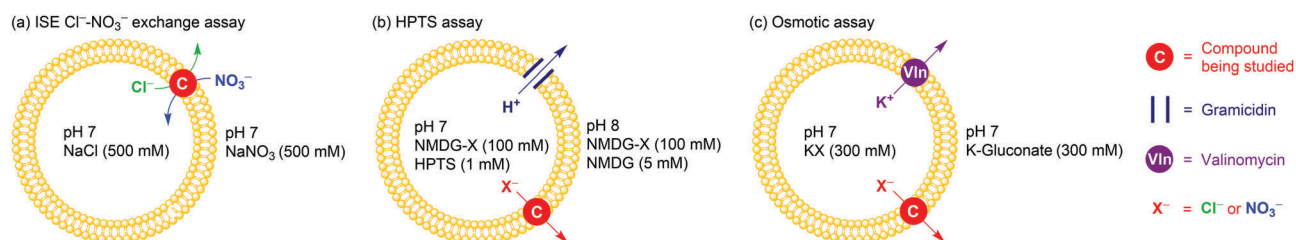


Fig. 2 Schematic representation of three membrane transport assays used. See ESI† for Experimental details.



and suspended in a buffered  $K^+$  gluconate (300 mM) solution. A combination of valinomycin (for transporting  $K^+$ ) and an anion transporter give overall  $K^+-X^-$  cotransport (Fig. 2c), leading to water efflux to balance the osmotic difference and the resultant shrinkage of vesicles, which can be monitored by following the light scattering of vesicles using a fluorometer.<sup>15</sup> This provides a method of quantifying the rate of anion uniport facilitated by anion transporters.<sup>14</sup> For all compounds, we determined the  $EC_{50}$  value at 200 s in the HPTS assay and at 600 s in the osmotic assay. In both assays, the  $Cl^-/NO_3^-$  transport selectivity  $S_t(Cl^-/NO_3^-)$  was quantified by the ratio between  $EC_{50}$  of  $NO_3^-$  transport and  $EC_{50}$  of  $Cl^-$  transport, where an  $S_t(Cl^-/NO_3^-)$  value  $> 1$  indicates  $Cl^-$  selectivity (Table 1).

The results from both HPTS and osmotic assays demonstrate that  $NO_3^-$  transport proceeds faster than  $Cl^-$  transport for monopodal transporters 1–3 and dipodal transporter 4. This is consistent with the higher lipophilicity of  $NO_3^-$  than  $Cl^-$  while contrary to the  $Cl^- > NO_3^-$  binding selectivity of the compounds in acetonitrile. The results suggest that in these cases transport selectivity is governed by the ease of anion dehydration. In contrast to 1–4, tripodal compounds 5 and 6 transported  $Cl^-$  faster than  $NO_3^-$ , with 5 showing an 8-fold  $Cl^-/NO_3^-$  selectivity in the HPTS assay and a 10-fold selectivity in the osmotic assay. Therefore, 5 and 6 are unusual examples of anion transporters that can overrule the normally observed  $NO_3^- > Cl^-$  lipophilicity bias. This cannot be interpreted as a general “anti-Hofmeister” selectivity as 5 and 6 transported the more lipophilic anion  $Br^-$  faster than  $Cl^-$  (Fig. S50 in ESI†). It is interesting to note that the transport selectivity  $S_t(Cl^-/NO_3^-)$  in general shows a good correlation with the binding selectivity  $S_b(Cl^-/NO_3^-)$  (Fig. 3 and Fig. S52 in ESI†). Both selectivities follow the trend of  $5 > 6 > 1 > 2-4$ . Chloride  $>$  nitrate selectivity is only observed for 5 and 6 that have the largest binding preference for  $Cl^-$  over  $NO_3^-$  among the library. Squaramide 1, despite being a  $NO_3^-$ -selective transporter, has a significantly larger  $S_t(Cl^-/NO_3^-)$  value compared with 2–4, consistent with 1 being the third most  $Cl^- > NO_3^-$  selective binder. The correlation, however, is not perfect with compounds 2–4. This may be rationalised by the idea that the rate of ionophore-catalysed ion transport depends not only on the ion binding affinity (which determines the amount of ion-ionophore complex with respect to the free

ionophore) but also on the ability of the complex to translocate across lipid bilayers.<sup>16</sup>

It is interesting to compare the activity determined in the three membrane transport assays. The data from HPTS and osmotic assays agreed well with each other, with the  $EC_{50}$  value in the osmotic assay consistently being about an order of magnitude higher than that in the HPTS assay for the same compound transporting the same anion.<sup>17</sup> Both assays have revealed the  $Cl^-$  transport activity in the sequence of  $6 > 5 > 1 > 3 > 2 > 4$ , whereas  $NO_3^-$  transport activity in the sequence of  $3 > 2 > 1 > 6 > 5 > 4$ . The  $Cl^-$ – $NO_3^-$  exchange assay showed the activity sequence of  $1 > 6 > 3 > 2 \approx 5 > 4$ , which reflected the activity of transporting the “slower” anion ( $Cl^-$  in the cases of 1–4, and  $NO_3^-$  in the cases of 5 and 6). By converting the  $EC_{50}$  values to normalised activities, a reasonable agreement between the three transport assays can be observed (Table S2 in ESI†). It is evident that the  $Cl^-$ – $NO_3^-$  exchange assay gives a fair assessment of  $Cl^-$  transport activity of compounds 1–4 because the overall process is rate-limited by  $Cl^-$  transport in these cases. However, the  $Cl^-$  transport activity of 5 and 6 is clearly underestimated in the exchange assay. Both compounds are in fact better  $Cl^-$  transporters than the best  $Cl^-$ – $NO_3^-$  exchanger 1. Compound 1 turned out to be neither the best  $Cl^-$  transporter nor the best  $NO_3^-$  transporter (Fig. 4).

For a better understanding of the  $Cl^- > NO_3^-$  selectivity of the tripodal scaffold, we performed PM6-optimisation of the structures of the  $Cl^-$  and  $NO_3^-$  complexes of 5. Fig. 5 (see Fig. S53 in ESI† for ball-and-stick models) shows that the spherical anion  $Cl^-$  fits well and is well-encapsulated inside the cavity of the tripodal scaffold, whereas binding of the planar anion  $NO_3^-$  forced the scaffold to adopt a more open conformation leaving a significant part of the bound  $NO_3^-$  exposed to the solvent. Encapsulation of the anion is crucial for effective membrane transport as demonstrated by previous work by Davis<sup>13</sup> and us,<sup>10b</sup> and therefore the poor encapsulation of  $NO_3^-$  could contribute to the  $Cl^- > NO_3^-$  transport selectivity of the tripodal compounds.

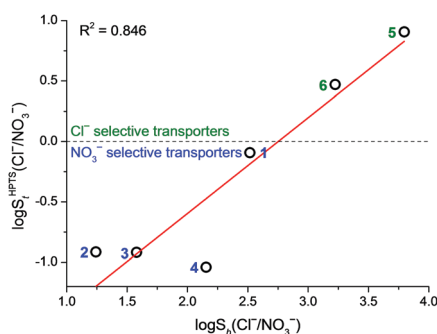


Fig. 3 Correlation between transport selectivity  $\log S_t^{\text{HPTS}}(Cl^-/NO_3^-)$  and binding selectivity  $\log S_b(Cl^-/NO_3^-)$ . Compound numbers are shown next to the data points.

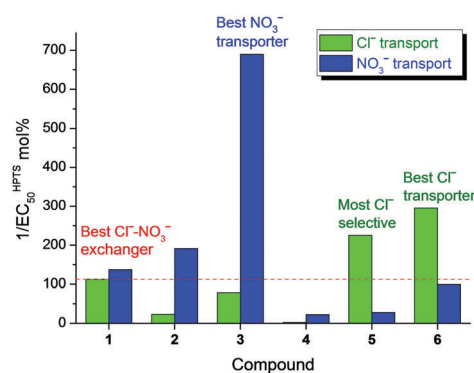


Fig. 4 Histogram showing activities of 1–6 in facilitating uniport of  $Cl^-$  (green bars) and  $NO_3^-$  (blue bars). Activities are expressed as reciprocal of  $EC_{50}$  values (at 200 s) determined in the HPTS assay. The red dashed line shows that compound 1 has the highest activity in facilitating the slower anion uniport process, which explains its highest activity in the  $Cl^-$ – $NO_3^-$  exchange assay.



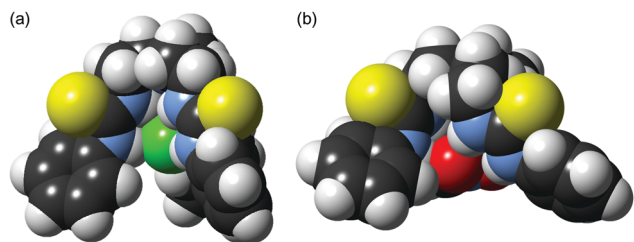


Fig. 5 PM6-optimised structures of  $\text{Cl}^-$  (a) and  $\text{NO}_3^-$  (b) complexes of **5** shown in space-filling models.

In summary, we have examined the chloride vs. nitrate selectivity of representative hydrogen bond-based anion transporters in both binding and transport. A strong correlation between transport selectivity and binding selectivity has been found. Only the tripodal transporters **5** and **6** that have a large  $\text{Cl}^-/\text{NO}_3^-$  binding preference can overcome the Hofmeister lipophilicity bias to transport  $\text{Cl}^-$  selectively over  $\text{NO}_3^-$ . We have compared the anion transport potency determined from different assays, showing that the commonly used  $\text{Cl}^-/\text{NO}_3^-$  exchange assay is valid in most cases for evaluating  $\text{Cl}^-$  transport activity but sometimes gives an underestimation in the case of a poor  $\text{NO}_3^-$  transporter. Importantly, we have also demonstrated that compound **5** functions as a highly  $\text{Cl}^-$ -selective transporter showing a  $\sim 10$ -fold  $\text{Cl}^-$  over  $\text{NO}_3^-$  selectivity in membrane transport, because of its complementary fit for  $\text{Cl}^-$ . We believe that the structure-selectivity and binding-transport relationships demonstrated here, and the different assays provided in this work will provide valuable tools for future development of highly potent and selective anion transporters for biomedical applications.

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- N.B. there are transporters that are incapable of (or have a poor activity in) mediating anion uniport. Examples include prodigiosin and an analogue of **5** with cyano substituents. In those cases, the HPTS assay showed no gramicidin-induced enhancement of transport rate and the valinomycin-coupled assay showed little or no activity. See ref. 10b.
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- The difference in  $\text{EC}_{50}$  between the two assays is mainly due to the fact that the HPTS assay measures a much smaller flux of  $\text{X}^-$  (5 mM) compared with the osmotic assay (300 mM) and also in part due to the different vesicle sizes. As a result, the transport process in the HPTS assay completes more quickly than in the osmotic assay. For a discussion of transport rate dependence on vesicle size, see H. Valkenier, N. L. Mora, A. Kros and A. P. Davis, *Angew. Chem., Int. Ed.*, 2015, **54**, 2137–2141.

