



Cite this: *Dalton Trans.*, 2016, **45**, 11892

Received 22nd May 2016,

Accepted 30th June 2016

DOI: 10.1039/c6dt02046a

www.rsc.org/dalton

Tris-ureas as transmembrane anion transporters†

Martina Olivari,^a Riccardo Montis,^a Stuart N. Berry,^b Louise E. Karagiannidis,^b Simon J. Coles,^b Peter N. Horton,^b Lucy K. Mapp,^b Philip A. Gale*^b and Claudia Caltagirone*^a

Nine tris-urea receptors (L^1 – L^9) have been synthesised and shown to coordinate to a range of anionic guests both by ^1H NMR titration techniques and single crystal X-ray structural analysis. The compounds have been shown to be capable of mediating the exchange of chloride and nitrate and also chloride and bicarbonate across POPC or POPC : cholesterol 7 : 3 vesicle bilayer membranes at low transporter loadings. An interesting dependency of anion transport on the nature of the cation is evidence to suggest that a M^+/Cl^- cotransport process may also contribute to the release of chloride from the vesicles.

Introduction

Transmembrane transport of anions across lipid bilayers is an important biological process that is normally regulated by complex membrane spanning proteins. A range of diseases, known as “channelopathies”, including cystic fibrosis, are caused by malfunctioning ion channels.¹ There is currently interest in the design of synthetic membrane transporters for anions that can act as potential future therapeutic substitutes for these malfunctioning proteins and have other biological applications.^{2–4}

Gale and co-workers have recently described the anion binding properties of a series of *ortho*-phenylenediamine-based bis-ureas.^{5,6} These compounds are highly effective anion transporters that function by an anion antiport, and in some cases a HCl symport carrier mechanism. Addition of electron-withdrawing groups to either the central core or the peripheral phenyl groups improved the anion transport ability: the transporter activity increased with the electron withdrawing strength of the substituent with the trend $\text{H} < \text{F} \approx \text{Cl} < \text{CF}_3 < \text{CN} < \text{NO}_2$. Indeed, the *p*-nitro functionalised compound was shown to possess a very high transport activity, facilitating chloride efflux at concentrations as low as 1 : 1 000 000 transporter to lipid molar ratio.

^aDipartimento di Scienze Chimiche e Geologiche, Università degli Studi di Cagliari, S.S. 554 Bivio per Sestu, 09042 Monserrato (CA), Italy. E-mail: ccaltagirone@unica.it

^bChemistry, University of Southampton, Southampton, SO17 1BJ, UK.

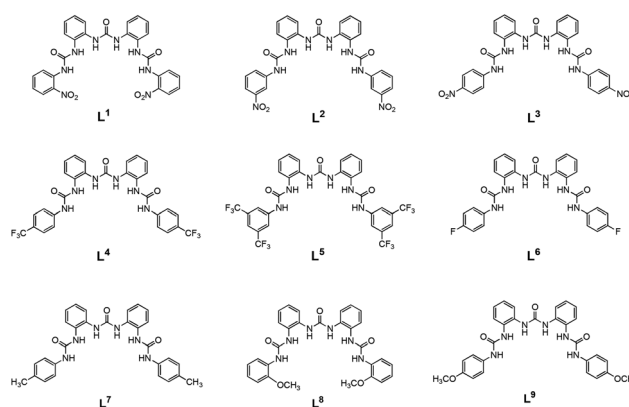
E-mail: philip.gale@soton.ac.uk

† Electronic supplementary information (ESI) available: Additional information as noted in the text including synthetic details for the preparation of L^1 – L^9 , fittings of ^1H -NMR titrations, crystallographic tables, transport studies. CCDC 1481148–1481150 for [$L^5(\text{Cl}^-)$](TBA⁺), [$L^4(\text{Cl}^-)_2$](TBA⁺)₂, [$L^5(\text{AcO}^-)$](TBA⁺). For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c6dt02046a

Gale *et al.* have also reported the transmembrane anion transport of phosphoric triamide and thiophosphoric triamide-based receptors,⁷ and tris-urea tripodal receptors.^{8–10}

Tris-urea receptors can be divided in two main families: tripodal receptors based on flexible linkers such as TREN (tris-(2-aminoethyl)amine) that are able to preferentially bind oxoanions^{5,11–19} and to work as organogelators,²⁰ or rigid spacers such as cyanuric acid,²¹ benzene,²² or trindane.²³

Recently Wu and co-workers designed and synthesised a new family of tris-ureas^{24,25} and tris-thioureas²⁶ developed mimicking the scaffold of terpyridine as efficient receptors for phosphate and sulfate. Starting from the interesting results obtained by Gale with the *ortho*-phenylenediamine-based bis-ureas transporters we decided to expand the family of tris-ureas reported by Wu and therefore we synthesised nine receptors (L^1 – L^9 in Scheme 1). We investigated the anion binding properties both in solution and in the solid state of the nine receptors and their ability to transport anions across lipid bilayers.



Scheme 1 Representation of receptors L^1 – L^9 . Receptor L^3 has already been published.²⁵



Results and discussion

The synthesis of receptor **L**³ has previously been reported by Wu.²⁵ **L**¹–**L**⁹ were synthesized *via* different reaction steps. Firstly 1,3-bis(2-aminophenyl)urea was prepared by reaction of *ortho*-phenylenediamine with *ortho*-nitro-phenylisocyanate in a mixed solvent THF/toluene at 0 °C and subsequent reduction of the 1-(2-nitrophenyl)-3-(2-aminophenyl)urea obtained by hydrazine and Pd/C 10%. 1,3-Bis(2-aminophenyl)urea was the reacted with the appropriate isocyanate (4-(trifluoromethyl) phenyl isocyanate; 3,5-bis(trifluoromethyl)phenyl isocyanate; 4-fluorophenyl isocyanate; *p*-tolyl isocyanate; 2-methoxyphenyl isocyanate; 4-methoxyphenyl isocyanate) in refluxing dichloromethane (DCM) under a N₂ atmosphere to obtain **L**¹–**L**⁹ in 60–90% yield (see ESI† for synthetic details).

Anion-binding studies were performed by means of ¹H-NMR titrations in DMSO-*d*₆. Stability constants from the ¹H-NMR titration curves obtained (see ESI Fig. S1–S13†) were calculated by fitting the data to a 1:1 binding model using EQNMR²⁷ as shown in Table 1.

Under the conditions of these experiments, the receptors did not interact with nitrate (*i.e.* no shift of the NH proton resonances occurred upon addition of tetrabutylammonium nitrate). Interestingly, receptors **L**¹–**L**³ which contain nitro electron-withdrawing groups showed little interaction with chloride, while addition of bicarbonate caused the disappearance of the signals attributed to the urea NH groups, evidence in support of a deprotonation or exchange process. Receptor **L**⁴, bearing one CF₃ substituent on the peripheral phenyl ring showed some affinity for chloride. On the other hand, receptors **L**⁶, **L**⁷, and **L**⁹ were able to bind both chloride and bicarbonate with comparable stability constants, while **L**⁵ and **L**⁸ bound bicarbonate preferentially.

A series of crystallization experiments of receptors **L**¹–**L**⁹ in presence of an excess of anions such as acetate, chloride, bicarbonate and nitrate were carried out with the aim of investigating the anion-binding properties of the receptors in the

solid state. In order to be consistent with solution studies, all the crystallization experiments were conducted in DMSO. However, (compatibly with the low solubility of receptors **L**¹–**L**⁹ in common solvents) with the aim of investigating the possible influence of less polar solvents in the anion-binding process, other solvents (AcOEt, MeOH, EtOH, THF, MeNO₂, MeCN) and mixture of solvents (MeOH/MeNO₂ and THF/DMF) were employed. Details of the crystallization experiments are reported in ESI (Table S1†).

As shown in Table S1 (see ESI†) only a limited number of crystallizations were successful in producing single crystals. In particular, for a total of sixty-six crystallization experiments, only seven gave samples suitable for X-ray investigation. Within these, three gave the crystal structure of the simple tetrabutylammonium salt and the remaining four resulted in crystal structures [L⁴(Cl[−])₂](TBA⁺)₂**a**, [L⁴(Cl[−])₂](TBA⁺)₂**b** (isostructures obtained from MeOH/MeNO₂ and MeCN respectively; for the structural description the code [L⁴(Cl[−])₂](TBA⁺)₂ is used to identify both), [L⁵(Cl[−])](TBA⁺) and [L⁵(AcO[−])](TBA⁺) (both obtained from DMSO).

In general, these results seem to be consistent with the low anion-binding affinity observed in solution studies (Table 1). This is particularly evident for receptors **L**¹–**L**³, where the unsuccessful crystallization experiments agree with the negligible interactions observed in solution (Table 1). In the case of receptor **L**⁴, though the compound unexpectedly forms a 1:2 [L⁴(Cl[−])₂](TBA⁺)₂ complex, the affinity toward Cl[−] is confirmed. For the remaining receptors **L**⁵–**L**⁹, only in the case of receptor **L**⁵ it was possible to isolate single crystals of anion complexes with chloride and acetate (not investigated in solution).

[L⁴(Cl[−])₂](TBA⁺)₂ crystallised in the triclinic crystal system (space group *P* $\bar{1}$) with an asymmetric unit consisting of one receptor **L**⁴, two chloride anions and two tetrabutylammonium counterions, resulting in a 1:2 complex. **L**⁴ adopts a closed conformation, with the urea NHs all oriented toward the centre of a pseudo-cavity. The three urea groups are slightly tilted to interact *via* N–H...Cl hydrogen bonds (N–H...Cl distances are in the range 2.36(2)–2.51(2) Å, average 2.43 Å) with the two Cl[−] anions which respectively lie below (Cl1) and above (Cl2) the pseudo-cavity (Fig. 1a and b). In particular it is worth noticing that the shortest distances are observed for Cl2 (N1–H1...Cl2 2.36(2) Å, N1–H1...Cl2 2.34(2) Å).

[L⁵(Cl[−])](TBA⁺) also adopts a triclinic crystal system (space group *P* $\bar{1}$). The asymmetric unit consists of one independent receptor **L**⁵, one independent chloride and one independent tetrabutylammonium resulting in a 1:1 complex. Similarly to **L**⁴ the receptor **L**⁵ shows a closed conformation with the two peripheral urea NHs (Fig. 2a) pointing at the centre of the pseudo-cavity and interacting with the chloride *via* four N–H...Cl hydrogen bonds (N–H...Cl) distances are in the range 2.32(2)–2.75(3) Å (average distance 2.53 Å). The central urea NHs are tilted to interact with an adjacent receptor-chloride unit *via* a second set of N–H...Cl hydrogen bonds (N–H...Cl distances are 2.44(3) Å and 2.74(3) Å respectively) forming a centrosymmetric dimer (Fig. 2b).

Table 1 Association constants (K_a/M^{-1}) for the formation of complexes of **L**¹–**L**⁹ with anions added as tetrabutylammonium salts (or tetraethylammonium in the case of hydrogencarbonate) in DMSO-*d*₆ at 300 K. All errors estimated to be <14% (see ESI)

Receptors	Anions		
	Cl [−]	HCO ₃ [−]	NO ₃ [−]
L ¹	<10	Deprotonation ^a	No interaction
L ²	<10	Deprotonation ^a	No interaction
L ³	<10	Deprotonation ^a	No interaction
L ⁴	262	Deprotonation ^a	No interaction
L ⁵	<10	226	No interaction
L ⁶	205	203	No interaction
L ⁷	226	221	No interaction
L ⁸	28	802	No interaction
L ⁹	225	240	No interaction

^a The NHs signals disappeared after the addition of one equivalent of anion.



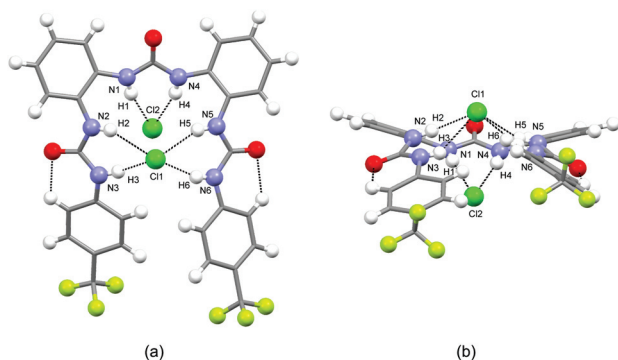


Fig. 1 Pseudo-cavity and main intermolecular interactions observed in structure $L^4(Cl^-)$, viewed along two orthogonal projections. TBA^+ counter cations are omitted for clarity. $N-H\cdots Cl$ hydrogen bonds are indicated as black dashed lines.

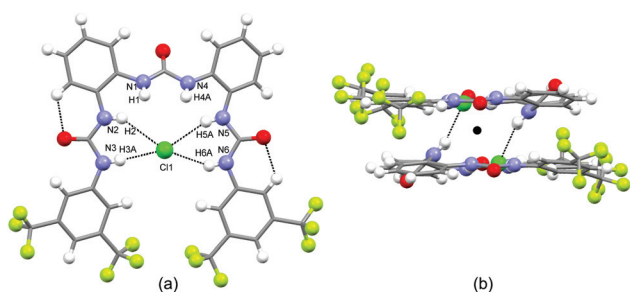


Fig. 2 Pseudo-cavity and main intermolecular interactions observed in structure $L^5(Cl^-)$, viewed along two orthogonal projections. TBA^+ counter cations and positional disorder in the CF_3 groups are omitted for clarity. $N-H\cdots O$ hydrogen bonds are indicated as black dashed lines, centre of inversion as black circle.

Similarly to the previous two structures, $[L^5(AcO^-)](TBA^+)$ crystallises in the triclinic crystal system (space group $P\bar{1}$). The asymmetric unit consists of one independent receptor L^5 , one independent acetate and one independent tetrabutylammonium. The receptor molecule adopts a closed conformation (Fig. 3a) to form a pseudo-cavity similar to those observed for $[L^4(Cl^-)]_2(TBA^+)_2$ and $[L^5(Cl^-)](TBA^+)$. The peripheral NHs are oriented toward the centre of the pseudo-cavity interacting with the AcO^- via $N-H\cdots O$ hydrogen bonds ($N-H\cdots O$ distances are in the range 1.89(2)–2.34(2) Å, average distance 2.13 Å). This is oriented perpendicularly to the plane of the pseudo-cavity and interacts via two $N-H\cdots O$ hydrogen bonds ($N-H\cdots O$ distances are 1.97(2) Å and 1.99(2) Å respectively) with the central urea NHs of an adjacent receptor molecule (Fig. 3b) to form a centrosymmetric dimer similar to that observed for $[L^5(Cl^-)](TBA^+)$.

The anion transport properties of receptors L^1-L^9 were studied using vesicle-based methods.²⁸ A sample of unilamellar POPC vesicles was prepared containing 489 mM NaCl buffered to pH 7.2 with 5 mM sodium phosphate salts. The

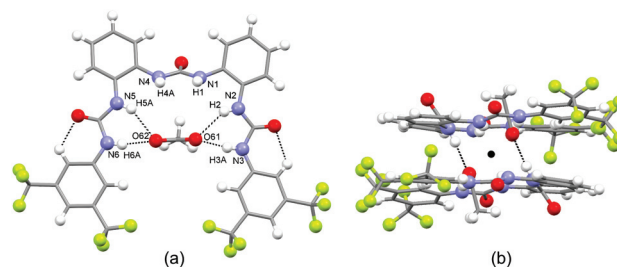


Fig. 3 Pseudo-cavity and main intermolecular interactions observed in structure $[L^5(AcO^-)](TBA^+)$; (a) $N-H\cdots O$ hydrogen bonds involving peripheral ureas and AcO^- . (b) $N-H\cdots O$ hydrogen bonds involving central ureas and AcO^- and centro-symmetric dimer. The molecules are oriented to best show the intermolecular interactions. TBA^+ counter cations are omitted for clarity. $N-H\cdots O$ hydrogen bonds are indicated as black dashed lines, centre of inversion as black circle.

vesicles were suspended at a lipid concentration of 1 mM in 489 mM $NaNO_3$ buffered to pH 7.2 with 5 mM sodium phosphate salts.

A small amount of DMSO solution of the receptor (0.02–2 mol% with respect to lipid) was added to the vesicles suspension, and the resulting chloride efflux was monitored using a chloride ion selective electrode (ISE) for 300 s. At the end of the experiment, the vesicles were lysed by the addition of detergent, and the final electrode reading was used to calibrate 100% chloride release. We found that all the compounds except L^1 (i.e. L^2-L^9) (at 2 mol% with respect to lipid) were capable of mediating chloride transport.

Under these experimental conditions the most active compounds are to be L^2 , L^3 , L^4 and L^6 (Fig. 4 and S14–S29 in ESI†) In order to determine the mechanism of chloride release by receptors L^2-L^9 , the transport assays were repeated suspending

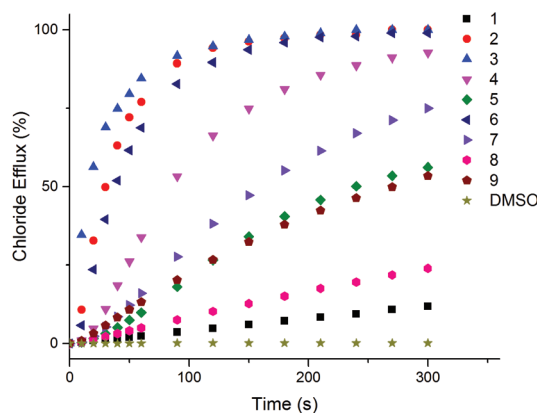


Fig. 4 Chloride efflux promoted by a DMSO solution of compounds L^1-L^9 (2 mol% carrier to lipid) from unilamellar POPC vesicles loaded with 489 mM NaCl buffered to pH 7.2 with 5 mM sodium phosphate salts. The vesicles were dispersed in 489 mM $NaNO_3$ buffered to pH 7.2 with 5 mM sodium phosphate salts. At the end of the experiment detergent was added to lyse the vesicles and calibrate the ISE to 100% chloride efflux. Each point represents an average of three trials. DMSO was used as a control.



POPC vesicles loaded with NaCl (451 mM with 20 mM phosphate buffer at pH 7.2) in a solution of Na_2SO_4 (150 mM with 20 mM phosphate buffer at pH 7.2). DMSO suspensions of compounds L^2 – L^9 were then added to the suspension. Usually during this experiment the antiport mechanism ($2\text{Cl}^-/\text{SO}_4^{2-}$ exchange) would not be expected to be observed due to the high hydrophilicity of the SO_4^{2-} anion²⁹ which would inhibit the chloride efflux from the liposomes.⁹ However, as shown in Fig. 5, during the first two minutes of the experiment significant chloride release was observed for compounds L^3 , L^2 and L^6 . There is also some chloride efflux mediated by the other compounds except for L^4 and L^5 under these conditions. After 120 s, a pulse of NaHCO_3 solution was added, and we observed an increase of chloride efflux (Fig. 5 and S31–S46 in ESI†), evidence in support of a chloride/bicarbonate exchange process.

A lucigenin assay for chloride/sulfate exchange was run with compounds L^2 – L^9 which showed that these compounds do not mediate sulfate transport (see ESI, Fig. S56† in the case of L^6 , as a representative compound).¹⁰

Although it has been recently reported that ureas and thioureas can facilitate proton or hydroxide transport,³⁰ a HPTS assay to test for H^+/Cl^- co-transport resulted in inconclusive results for the class of molecules presented herein (see ESI, Fig. S57† in the case of L^6 as a representative compound).

To further examine the origin of the chloride transport during the first two minutes of this assay, we examined the possibility of a mechanism involving sodium/chloride co-transport. The $\text{Cl}^-/\text{NO}_3^-$ transport assays were repeated using vesicles containing caesium chloride (489 mM NaCl buffered to pH 7.2 with 5 mM sodium phosphate salts) instead of sodium chloride, suspended in an isotonic sodium nitrate solution. In the event of NaCl co-transport we would expect the

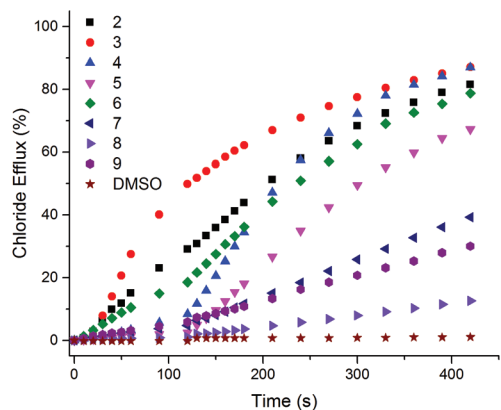


Fig. 5 Chloride efflux promoted by a DMSO solution of compounds L^2 – L^9 (2 mol% carrier to lipid) from unilamellar POPC vesicles loaded with 451 mM NaCl buffered to pH 7.2 with 20 mM sodium phosphate salts. The vesicles were dispersed in 150 mM Na_2SO_4 buffered to pH 7.2 with 20 mM sodium phosphate salts. At $t = 120$ s a solution of sodium bicarbonate was added such that the external concentration of bicarbonate was 40 mM. At the end of the experiment, detergent was added to lyse the vesicles and calibrate the ISE to 100% chloride efflux. Each point represents an average of three trials. DMSO was used as a control.

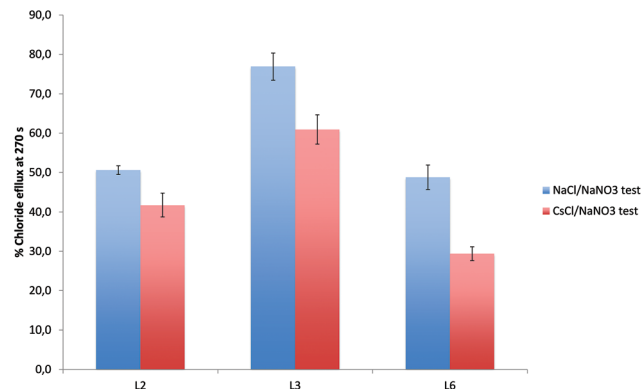


Fig. 6 Percentage chloride efflux at 270 s mediated by L^2 , L^3 , and L^6 (0.1 mol% carrier to lipid) from unilamellar POPC vesicles loaded with either 489 mM NaCl (red) or 489 mM CsCl (blue) buffered to pH 7.2 with 5 mM sodium phosphate salts. The vesicles were dispersed in 489 mM NaNO_3 buffered to pH 7.2 with 5 mM sodium phosphate salts.

rate of chloride release to be different in the presence of CsCl. As shown in Fig. 6 in the case of L^2 , L^3 , and L^6 (and Fig. S47–S55 in the ESI†) the release of chloride is dependent on the nature of metal cation and it is reduced when the internal solution was replaced by CsCl.

The results described above suggest, from the first assay, that the receptors can facilitate $\text{Cl}^-/\text{NO}_3^-$ exchange; the second assay, supports the hypothesis of $\text{Cl}^-/\text{HCO}_3^-$ antiport mechanism although in the first two minutes in the presence of sulfate for all the compounds except L^4 and L^5 a small amount of chloride efflux was observed. The caesium chloride assay demonstrates that for L^2 , L^3 , L^6 – L^9 the counter cation is involved in transport possibly *via* a M^+/Cl^- co-transport process.

We tested the $\text{Cl}^-/\text{NO}_3^-$ antiport activity of L^2 – L^9 in vesicles composed of POPC–cholesterol (7 : 3). This mixture is a closer mimic of biological membranes than pure POPC lipid bilayers. The presence of cholesterol is known to increase the order in the bilayer and its viscosity. All the receptors tested showed a reduced rate of transport in the POPC–cholesterol system (see Fig. 7 for L^2 and Fig. S58–S64 in ESI† for L^3 – L^9), providing evidence that for this class of receptor diffusion through the interior of the bilayer may be the rate determining step.³¹

To quantify the transport activity of compounds L^2 – L^9 Hill analyses^{32,33} for the chloride/nitrate and chloride/bicarbonate antiport assays were performed (see Fig. S14–S29, S31–S46 in ESI†). Hill analysis allows the calculation of the $\text{EC}_{50,270\text{ s}}$ which is a measure of transporter efficiency, defined as the required receptor concentration to mediate 50% of the total chloride efflux 270 s after the addition of the carrier (or after the bicarbonate ‘pulse’). This allows us to compare the transport activity of the compounds. These values are summarised in Table 2, together with the Hill coefficients, which can be correlated to the number of transporter molecules required to transport a single anion and can provide further evidence for a mobile carrier mechanism.



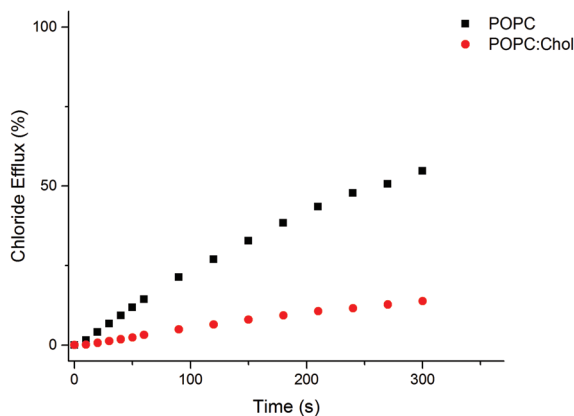


Fig. 7 Chloride efflux promoted by a DMSO solution of compound L^2 (0.2 mol% carrier to lipid) from unilamellar vesicles comprising of either POPC or POPC–cholesterol (7 : 3 molar ratio, POPC : Chol), loaded with 489 mM NaCl buffered to pH 7.2 with 5 mM sodium phosphate salts. The vesicles were dispersed in 489 mM NaNO₃ buffered to pH 7.2 with 5 mM sodium phosphate salts. At the end of the experiment detergent was added to lyse the vesicles and calibrate the ISE to 100% chloride efflux. Each point represents an average of three trials. DMSO was used as a control.

Table 2 The EC₅₀ is the concentration (mol% carrier to lipid) needed to obtain 50% efflux after 270 s and n is the Hill coefficient that represents an estimate of the number of transporter molecules required to transport a single anion

Compound	EC ₅₀ at 270 s (Cl ⁻ /NO ₃ ⁻)	n	EC ₅₀ at 270 s (Cl ⁻ /HCO ₃ ⁻)	n
L^2	0.095	1.207	0.50	0.76
L^3	0.047	1.67	0.21	0.77
L^4	0.039	1.07	0.14	0.98
L^5	0.066	0.88	0.25	0.97
L^6	0.10	1.39	0.76	0.80
L^7	0.6	0.96	4.27	0.88
L^8	4.98	1.29	21.93	0.86
L^9	1.96	0.88	8.43	0.84

From the EC_{50,270 s} values reported in Table 2 the most active transporter among the series appears to be L^4 (EC_{50,270 s} 0.039 mol% and 0.14 mol% with respect to lipid for nitrate and bicarbonate antiport, respectively).

Further, Hill coefficients of ~ 1 provides further evidence that these compounds are functioning *via* a mobile carrier mechanism and are not forming membrane spanning channels in which we would expect the Hill coefficient to be higher.

Conclusions

In conclusion we have synthesised nine tris-urea receptors bearing a range of substituents attached to the pendant arms. We have studied the anion binding properties of the receptors both in solution and in the solid state and we have tested their ability to mediate chloride transport through membranes.

Solution studies demonstrate that these receptors bind anions with moderate stability constants (as reported in Table 1).

Solid state studies confirm the results observed in solution and indeed, despite many attempts, only three crystal structures were obtained. In particular, the crystal structure of L^4 in the presence of chloride suggests that the receptor has a good degree of pre-organization and binds chloride *via* the urea NH groups with a 1 : 2 stoichiometry. We also demonstrated that these systems are able to mediate transmembrane chloride transport as mobile carriers with different mechanisms, Cl⁻/NO₃⁻ and Cl⁻/HCO₃⁻ antiport, and metal-dependent cation co-transport. Hill-plot analysis demonstrates that the most active compound of the series is L^4 .³⁴

Acknowledgements

CC would like to thank Fondazione Banco di Sardegna for financial support. PAG thanks the University of Southampton and the A*STAR ARAP programme for a studentship (SNB), the Royal Society and the Wolfson Foundation for a Research Merit Award and the EPSRC (EP/K039466/1) (Core Capability for Chemistry Research in Southampton). We also thank the EPSRC for access to the crystallographic facilities at the University of Southampton.³⁵

Notes and references

- F. M. Ashcroft, *Ion Channels and Disease*, Academic Press, San Diego, 2000.
- P. A. Gale, R. Pérez-Tomás and R. Quesada, *Acc. Chem. Res.*, 2013, **46**, 2801–2813; N. Busschaert and P. A. Gale, *Angew. Chem., Int. Ed.*, 2013, **52**, 1374–1382.
- S. Matile, A. Vargas Jentzsch, J. Montenegro and A. Fin, *Chem. Soc. Rev.*, 2011, **40**, 2453–2474.
- (a) V. Soto-Cerrato, P. Manuel-Manresa, E. Hernando, S. Calabuig-Fariñas, A. Martínez-Romero, V. Fernández-Dueñas, K. Sahlholm, T. Knöpfel, M. García-Valverde, A. M. Rodilla, E. Jantus-Lewintre, R. Farràs, F. Ciruela, R. Pérez-Tomás and R. Quesada, *J. Am. Chem. Soc.*, 2015, **137**, 15892–15898; (b) S. N. Berry, V. Soto-Cerrato, E. N. W. Howe, H. J. Clarke, I. Mistry, A. Tavassoli, Y.-T. Chang, R. Pérez-Tomás and P. A. Gale, *Chem. Sci.*, 2016, DOI: 10.1039/c6sc01643j; (c) W. Van Rossom, D. J. Asby, A. Tavassoli and P. A. Gale, *Org. Biomol. Chem.*, 2016, **14**, 2645–2650; (d) A. I. Share, K. Patel, C. Nativi, E. J. Cho, O. Francesconi, N. Busschaert, P. A. Gale, S. Roelens and J. L. Sessler, *Chem. Commun.*, 2016, **52**, 7560.
- S. J. Moore, C. J. E. Haynes, J. Gonzalez, J. L. Sutton, S. J. Brooks, M. E. Light, J. Herniman, G. J. Langley, V. Soto-Cerrato, R. Perez-Tomas, I. Marques, P. J. Costa, V. Felix and P. A. Gale, *Chem. Sci.*, 2013, **4**, 103–117.
- L. E. Karagiannidis, C. J. E. Haynes, K. J. Holder, I. L. Kirby, S. J. Moore, N. J. Wells and P. A. Gale, *Chem. Commun.*, 2014, **50**, 12050–12053.



- 7 P. B. Cranwell, J. R. Hiscock, C. J. E. Haynes, M. E. Light, N. J. Wells and P. A. Gale, *Chem. Commun.*, 2013, **49**, 874–876.
- 8 N. Busschaert, P. A. Gale, C. J. E. Haynes, M. E. Light, S. J. Moore, C. C. Tong, J. T. Davis and J. W. A. Harrell, *Chem. Commun.*, 2010, **46**, 6252–6254.
- 9 N. Busschaert, L. E. Karagiannidis, M. Wenzel, C. J. E. Haynes, N. J. Wells, P. G. Young, D. Makuc, J. Plavec, K. A. Jolliffe and P. A. Gale, *Chem. Sci.*, 2014, **5**, 1118–1127.
- 10 N. Busschaert, M. Wenzel, M. E. Light, P. Iglesias-Hernández, R. Pérez-Tomás and P. A. Gale, *J. Am. Chem. Soc.*, 2011, **133**, 14136–14148.
- 11 B. Akhuli, I. Ravikumar and P. Ghosh, *Chem. Sci.*, 2012, **3**, 1522–1530.
- 12 P. Bose, R. Dutta and P. Ghosh, *Org. Biomol. Chem.*, 2013, **11**, 4581–4584.
- 13 R. Custelcean, *Chem. Commun.*, 2013, **49**, 2173–2182.
- 14 S. K. Dey, R. Chutia and G. Das, *Inorg. Chem.*, 2012, **51**, 1727–1738.
- 15 M. Emami Khansari, C. R. Johnson, I. Basaran, A. Nafis, J. Wang, J. Leszczynski and M. A. Hossain, *RSC Adv.*, 2015, **5**, 17606–17614.
- 16 Y. Hao, C. Jia, S. Li, X. Huang, X. J. Yang, C. Janiak and B. Wu, *Supramol. Chem.*, 2012, **24**, 88–94.
- 17 J. R. Hiscock, P. A. Gale and M. J. Hynes, *Supramol. Chem.*, 2012, **24**, 355–360.
- 18 M. Li, Y. Hao, B. Wu, C. Jia, X. Huang and X. J. Yang, *Org. Biomol. Chem.*, 2011, **9**, 5637–5640.
- 19 M. Li, B. Wu, C. Jia, X. Huang, Q. Zhao, S. Shao and X. J. Yang, *Chem. – Eur. J.*, 2011, **17**, 2272–2280.
- 20 M. Yamanaka, *J. Inclusion Phenom. Macrocyclic Chem.*, 2013, **77**, 33–48.
- 21 R. Dutta and P. Ghosh, *Eur. J. Inorg. Chem.*, 2013, 2673–2681.
- 22 V. K. Bhardwaj, S. Sharma, N. Singh, M. S. Hundal and G. Hundal, *Supramol. Chem.*, 2011, **23**, 790–800.
- 23 W. Kim, S. K. Sahoo, G. D. Kim and H. J. Choi, *Tetrahedron*, 2015, **71**, 8111–8116.
- 24 C. Jia, B. Wu, S. Li, Z. Yang, Q. Zhao, J. Liang, Q. S. Li and X. J. Yang, *Chem. Commun.*, 2010, **46**, 5376–5378.
- 25 R. Li, Y. Zhao, S. Li, P. Yang, X. Huang, X. J. Yang and B. Wu, *Inorg. Chem.*, 2013, **52**, 5851–5860.
- 26 Y. Zhang, R. Zhang, Y. Zhao, L. Ji, C. Jia and B. Wu, *New J. Chem.*, 2013, **37**, 2266–2270.
- 27 M. J. Hynes, *J. Chem. Soc., Dalton Trans.*, 1993, 311–312.
- 28 B. D. Smith and T. N. Lambert, *Chem. Commun.*, 2003, 2261–2268.
- 29 Y. Marcus, *J. Chem. Soc., Faraday Trans.*, 1991, **87**, 2995–2999.
- 30 X. Wu, L. W. Judd, E. N. W. Howe, A. M. Withecombe, V. Soto-Cerrato, H. Li, N. Busschaert, H. Valkenier, R. Perez-Tomas, D. N. Sheppard, Y.-B. Jiang, A. P. Davis and P. A. Gale, *Chem*, 2016, DOI: 10.1016/j.chempr.2016.04.002.
- 31 W. F. D. Bennett, J. L. MacCallum and D. P. Tieleman, *J. Am. Chem. Soc.*, 2009, **131**, 1972–1978.
- 32 A. V. Hill, *Biochem. J.*, 1913, **7**, 471.
- 33 S. Bhosale and S. Matile, *Chirality*, 2006, **18**, 849–856.
- 34 N. Busschaert, S. J. Bradberry, M. Wenzel, C. J. E. Haynes, J. R. Hiscock, I. L. Kirby, L. E. Karagiannidis, S. J. Moore, N. J. Wells, J. Herniman, G. J. Langley, P. N. Horton, M. E. Light, I. Marques, P. J. Costa, V. Félix, J. G. Frey and P. A. Gale, *Chem. Sci.*, 2013, **4**, 3036–3045; N. J. Knight, E. Hernando, C. J. E. Haynes, N. Busschaert, H. J. Clarke, K. Takimoto, M. García-Valverde, J. G. Frey, R. Quesada and P. A. Gale, *Chem. Sci.*, 2016, **7**, 1600–1608.
- 35 S. J. Coles and P. A. Gale, *Chem. Sci.*, 2012, **3**, 683–689.

