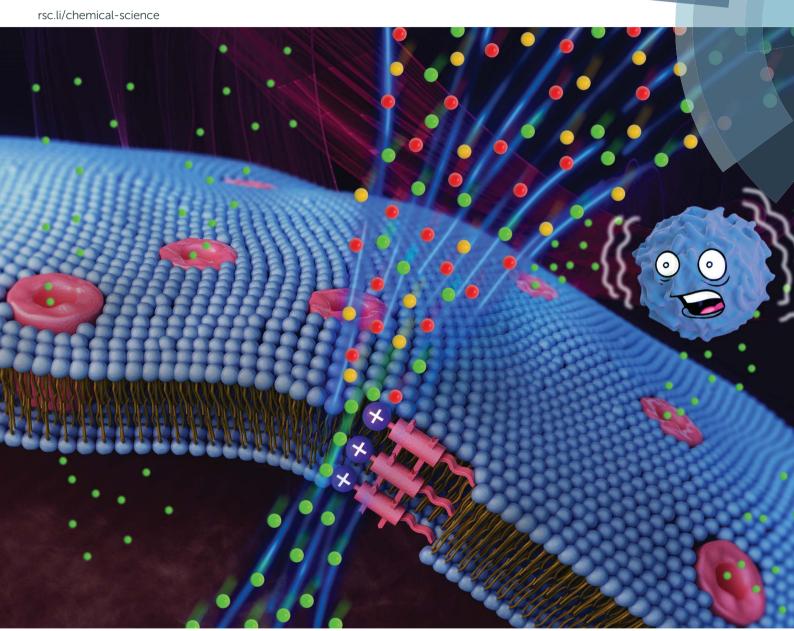
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A halogen bond-mediated highly active artificial chloride channel with high anticancer activity†

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Chloride-selective transmembrane carriers or channels might have possible uses in treating channelopathies or cancers. While chloride carriers have been extensively investigated, the corresponding chloride channels have remained limitedly studied. Moreover, all hitherto reported channel systems lack clearly definable and readily modifiable positions in their structures for the reliable construction and combinatorial optimization of their ion transport properties. As a result, the existing channels are limited by their large molecular weight, weak activity or low anion selectivity. In this report, we describe a readily accessible and robust monopeptide-based scaffold for the reliable construction of halogen bond-mediated artificial anion channels via directional assembly of electron-deficient iodine atoms, which create a transmembrane pathway for facilitating anion transport. The high intrinsic modularity of the backbone of the scaffold, which enables the rapid and combinatorial optimization of the transport activity and selectivity of channels, effectively delivers a highly active chloride channel A10. Such high activity in chloride transport subsequently leads to an excellent IC_{50} value of 20 μ M toward inhibiting the growth of human breast cancer cells (BT-474), an anticancer activity that is even higher than that of the well-known anticancer agent cisplatin.

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Introduction

Tightly regulated by varying families of ion channels, precise maintenance of ion concentration gradients across biological membranes is crucial for many biological and cellular processes. Chloride is the most abundant anion in the human body due to its important roles in controlling the membrane potential, cell volume, cellular pH balance, secretion of transepithelial fluid and electrolytes, etc. 1 Misregulation of chloride ions could lead to a variety of life-shortening 'channelopathies' including cystic fibrosis, Bartter syndrome and Dent's disease. Accordingly, significant research efforts have been made recently to coherently develop artificial chloride carriers2 or channels³ as possible replacements for natural chloride channels for channelopathies.4 On the other hand, artificial anion transporters might also disrupt pH gradients4f,5a,b or ion homeostasis^{5c} that may induce cell death, and therefore may have medical applications as anticancer agents. 4f,5

Recently, Matile et al. reported self-assembled chloride carriers2b,c or channels,3i elegantly exploiting halogen bonding interaction as the driving force for anion transport. Halogen bonds offer good bond strength and also good anion-induced directionality.6 Therefore, artificial anion transporters comprising simple structural motifs based on halogen-anion interactions may indeed offer a new horizon of prospects to comprehend complex chloride transport phenomena, often manifested in protein channels. Nevertheless, most documented artificial chloride channels purposefully harness various non-covalent interactions such as electrostatic, Hbonding, anion- π and halogen bonding forces. Channels that primarily rely on halogen bonds for facilitating anion transport still remain limitedly investigated with just one precedent.31 Such a rare occurrence presumably results from the lack of suitable molecular scaffolds for aligning electron-deficient iodine atoms into a well-defined one-dimensional (1D) array. Moreover, the majority of existing chloride channels^{3a-m} suffer from relatively complex molecular structures and/or low selectivity in anion recognition, greatly restricting their application in drug discovery.

We recently described a novel class of readily accessible monopeptide-based scaffolds capable of self-assembling into 1D columnar structures through highly directional intermolecular H-bonding forces, subsequently enabling rapid room-temperature gelation of diverse types of crude oils (Fig. 1a).^{7a-c}

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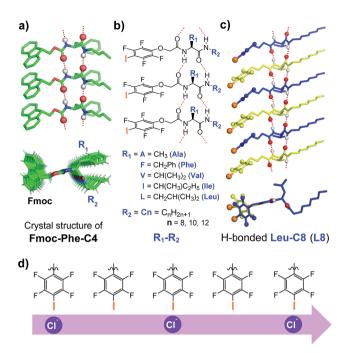


Fig. 1 (a) Crystal structure of F-Phe-C4, illustrating the formation of the 1D columnar stack with directionally assembled side chains. (b) Molecular design of the chloride-transporting channel library with systematically tunable $\rm R_1$ and $\rm R_2$ groups. (c) Side and top views of the computationally optimized H-bonded 1D columnar ensemble formed from L8. (d) A possible multi-ion jumping mode responsible for chloride transport.

Revealed by the crystal structure of **F-Phe-C4** (Fig. 1a), one unique feature of this monopeptide-based scaffold is that the same type of side chains such as Fmoc, R_1 and R_2 are always aligned to the same side as a result of the high directionality of H bonds (Fig. 1a).^{7 α} On this basis, we envisioned that replacing the Fmoc group with a chloride-binding unit such as the tetrafluoroiodobenzyl group may lead to an interesting class of artificial anion channels with anion transport mediated by

halogen bonds (Fig. 1b-d).^{3i,7d} A high intrinsic modularity involving R₁ and R₂ groups in the backbone should allow for rapid and combinatorial optimization of transport activity and selectivity of channels. In this report, we demonstrate that this indeed is the case, with the identified chloride channel exhibiting a high activity in chloride transport with respect to many other types of anions. We further demonstrate that human breast cancer cells (BT-474) are sensitive to a sodium chloride concentration gradient across the membrane, and this feature can be utilized by artificial chloride channels to inhibit cancer cell growth for their potential uses in cancer chemotherapy.^{4f,5}

Results and discussion

Combinatorial identification of highly active aniontransporting artificial channels

We synthesized a total of 15 possible channel molecules (five amino acids × three alkyl chains of different lengths, Fig. 1b) with the tetrafluoroiodobenzyl group as the chloride-binding and -transporting unit. Corroborated by the crystal structure of F-Phe-C4 (Fig. 1a) and the computationally optimized structure of L8 (Fig. 1c), the ability of these 15 molecules to pile up to form a H-bonded 1D structure can be convincingly demonstrated by the ability of A10, L8 and L10 to efficiently congeal in n-hexane via the formation of a 3D entangled fibrous network (ESI, Fig. S1†). The ion transport activities of the synthesized channels were then evaluated using the pH-sensitive HPTS (8-hydroxypyrene-1,3,6-trisulfonic acid) assay (Fig. 2 and S2†).8 In a typical experiment, large unilamellar vesicles (LUVs), encapsulating HPTS (100 μ M) and NaCl (100 mM) at pH = 7.0, were diluted into the same buffer at pH 8.0 to generate a pH gradient across LUVs. After the addition of channel molecules, changes in the fluorescence intensity of HPTS were monitored over 5 min. Fig. 2b shows that, among all 15 channels tested at 10 μM, A10, L8 and L10 display the highest ion transport capacities with fractional activities (R_{Cl}) of 60%, 68% and 59%, respectively. The corresponding EC₅₀ values, which are the

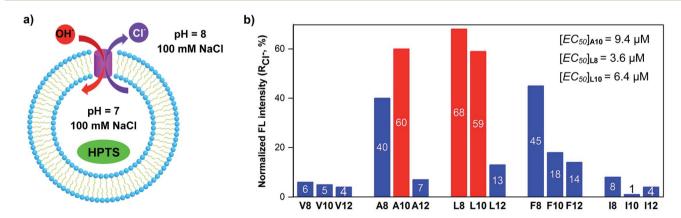


Fig. 2 (a) Schematic illustration of the HPTS assay for the Cl⁻ transport study conducted using a pH gradient of 7 to 8 and the pH-sensitive dye HPTS entrapped inside LUVs. (b) Normalized transport activities (R_{Cl^-}) obtained over 5 min at 10 μ M for all channel molecules. $R_{\text{Cl}^-} = (I_{\text{Cl}^-} - I_0)/(I_{\text{Triton}} - I_0)$ wherein I_{Cl^-} and I_0 are the ratiometric values of I_{460}/I_{403} before the addition of triton at t = 300 s, and I_{Triton} is the ratiometric value of I_{460}/I_{403} at t = 300 s right after the addition of triton with internal/external buffers containing 100 mM NaCl. All the data were averaged over two measurements.

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concentrations required to reach 50% transmembrane activity, were determined via Hill analyses to be 9.4, 3.6 and 6.4 μM, respectively (Fig. S3†).

Cl⁻/OH⁻ as the major transport species

Except for the presence of arrayed electron-deficient iodine atoms that can bind the Cl⁻ anion, ^{2b,c,3i} all peptide molecules carry no other readily accessible functional groups for strongly binding either cationic or anionic species. Further, varying extravesicular metal chloride salts from LiCl to CsCl produces near-identical changes in fluorescence intensity (Fig. 3a and S4†), suggesting the inability of L8 or A10 to transport any of the five alkali metal ions. These combined structural features and experimental evidence led us to believe that, rather than the H⁺/ M⁺ antiport mechanism, either the OH⁻/Cl⁻ antiport (Fig. 2a) or H⁺/Cl⁻ symport mechanism may largely account for the observed increases in the fluorescence intensity of HPTS.

To deduce the most likely transport species, we first carried out the SPQ assay9 using a chloride-sensitive SPQ dye (6-methoxy-N-(3-sulfopropyl)quinolinium). As illustrated in Fig. 3b and S5,† the addition of L8 or A10 results in a rapid quenching of SPQ fluorescence in a concentration-dependent manner. These results suggest that the influx of Cl⁻ increases with increasing channel concentrations from 0.32 to 5 μ M for L8 and from 2.5 to 20 μM for A10, thereby establishing the Cl⁻ anion as one of the molecular species transported by both L8 and A10.

Next, carbonyl cyanide 4-(trifluoromethoxy)phenylhydrazone (FCCP, a well-known proton carrier) was employed (Fig. 3c). Compared to the transport efficiencies of 11% for FCCP alone and of 57% for L8 alone, an increased transport efficiency of 13% (e.g., 78-57% - (11-3%)) for L8 in the presence of FCCP indicates a cooperative action between L8 and FCCP. In the case of A10, an even larger enhancement of 25% was observed (Fig. S6†). These results suggest that the transport rate of Cl⁻ is faster than that of H⁺.

The HPTS assay was then performed in the presence of valinomycin (VA, a K⁺-selective carrier, Fig. 3d) to compare the transport rate between OH- and Cl-. For this assay carried out under iso-osmolar K⁺ (extravesicular) vs. Na⁺ (intravesicular) and pH gradient, the VA-mediated influx of K⁺ ions will induce an anion channel-mediated influx of either OH or Cl in order to maintain overall charge equality. If the influx of OH is faster than that of Cl⁻, an increase in fluorescence intensity is anticipated. Experimentally, valinomycin at 25 pM produces a small increment of fluorescence intensity (7%) compared to the blank (3%). In a similar way, the ion transport activities of L8 (3.6 μ M) in the presence and absence of valinomycin were found to be near-identical (58% vs. 59%), which is indicative of a preferential transport of Cl $^-$ over OH $^-$ (e.g., Cl $^-$ > OH $^-$). A similar result was also seen for A10 (Fig. S7†). Consistent with transport activities presented in Fig. 3a, these data also suggest that the H⁺/M⁺ antiport mechanism is unlikely for L8-mediated enhancement in fluorescence intensity.

A mere increase of 1.3% (4.6-3.3%, Fig. 3e) in fluorescence intensity at 3.6 µM for molecule 1, carrying a simple benzene group, strongly suggests a lack of readily accessible functional

groups in the H-bonded peptidic backbone for interacting with either anions or protons, and thus a minor role played by the backbone in mediating chloride transport. Instead, it is halogen bonds formed between anions (e.g., Cl or OH) and electrondeficient iodine atoms that cause efficient exchanges between Cl⁻ and OH⁻ anions across LUVs as observed for L8 (57%), pointing to a high unlikelihood of having H⁺/Cl⁻ as the transport species. The herein assumed formation of halogen bonds I···Cl and I···OH was recently observed in their corresponding crystal structures^{2c} and can be further supported by ¹⁹F NMR titration experiments involving titrating 0-20 equiv. of tetrabutylammonium chloride into a D2O-saturated CDCl3 solution containing L8 at 1 mM (Fig. S8†). Relative to the internal standard (1,4-difluorobenzene, -120.50 ppm), increasing additions of up to 20 equiv. of TBACl led to increasing upfield shifts of up to 0.61 ppm in the chemical shift of ¹⁹F of L8, a fact that is consistent with earlier observations^{2b} and indicates binding between the chloride anion and the acidic iodine atom.

Additional comparisons among L8 and 2-4 convincingly demonstrate the importance of (1) the co-existence of two amide bonds that likely allow for a tighter and more directional stacking of channel molecules (L8 vs. 2), (2) the side chain of leucine for more efficient stacking (2 vs. 3) and (3) the amide bond in forming a H-bonded structure for ordered spanning of the hydrophobic membrane region (3 vs. 4). These summative findings are also in accordance with the molecular dynamics (MD) simulation results (Fig. 4).

Anion selectivity in chloride transport

With high activity exhibited by these chloride transporters, our subsequent investigations focused on evaluating and comparing anion selectivity for the most efficient anion channels formed by A10, L8 and L10. For this purpose, both intraand extravesicular anions were kept the same (100 mM NaX, $X^- = Cl^-$, Br^- , I^- , NO_3^- and ClO_4^-) with a proton gradient of pH 7 (inside) to pH 8 (outside). Data compiled in Fig. 3f, S9 and S10† show that A10 with an EC₅₀ value of 9.4 μ M exhibits good anion selectivity in chloride transport, and channels L8 and L10 likely selectively transport non-chloride anions (Fig. S10†).

Chloride transport through a channel mechanism

Single channel current traces for chloride transport, recorded in a planar lipid bilayer at various voltages including -100 mv in symmetric baths (cis chamber = trans chamber = 1 M KCl, Fig. 3g and S11†), unambiguously confirm that L8-mediated chloride transport occurs via a channel, rather than a carrier mechanism. On the basis of the fitted linear current-voltage (I-V) plot (Fig. 3h and S10†), the Cl⁻ conductance (γ_{Cl^-}) of L8 was found to be 586 \pm 11 fS.

To shed some light on the possible structural features of onedimensionally aligned channel molecules in the lipid membrane, MD simulation using the CHARMM program, 10a-e PME method10f and SHAKE algorithm10g was performed on L8. In particular, the H-bonded 1D structure, which consists of eight molecules of L8 (528 atoms), was first computationally optimized using the COMPASS force field10h and then was embedded

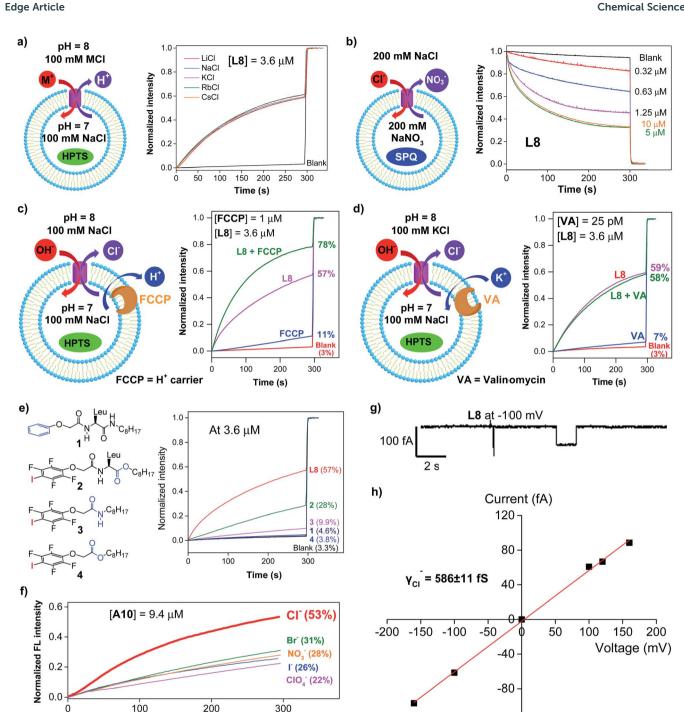


Fig. 3 (a) Transport selectivity of alkali metal ions by L8 (3.6 μ M) obtained from the HPTS assay by varying the extravesicular MCl (M⁺ = Li⁺, Na⁺, K^+ , Rb^+ and Cs^+). (b) Concentration-dependent changes in the fluorescence intensity of the chloride-sensitive SPQ dye ($\lambda_{ex} = 360$ nm, $\lambda_{em} = 360$ nm 430 nm) after the addition of L8 at different concentrations. Inside LUV: 200 mM NaNO₃ and 0.5 mM SPQ. Outside LUV: 200 mM NaCl. (c) Ion transport activities of L8 (3.6 μ M) determined in the absence or presence of proton transporter FCCP (1 μ M), indicating a preferential transport of Cl^- over H^+ . (d) Ion transport activities of L8 (3.6 μ M) determined in the absence or presence of a potassium carrier, valinomycin (VA, 25 μ M), indicating a preferential transport of Cl^- over OH^- . (e) Ion transport activities of four control compounds 1–4 at 3.6 μ M, suggesting a critical role of halogen bonds in mediating the transport of Cl⁻ and OH⁻. (f) Anion selectivity assayed for A10 (For L8 and L10, see Fig. S10†). (g) Single current trace of L8 recorded at -100 mv with symmetric baths (cis chamber = trans chamber = 1 M KCI). (h) I-V curve for obtaining the Cl⁻ conductance (γ_{Cl}) of **L8**. All the data were averaged over two measurements.

in a bilayer of 128 POPC molecules (17 152 atoms) solvated on two sides by 2×2397 water molecules (Fig. 4a). This leads to a simulation system of 32 062 atoms with a dimension of 70 Å

Time (s)

(w) \times 70 Å (w) \times 74 Å (h). After equilibration steps, the production run was carried out for 30 ns. The last 20 ns trajectories with 1000 structural snapshots were used for analysing

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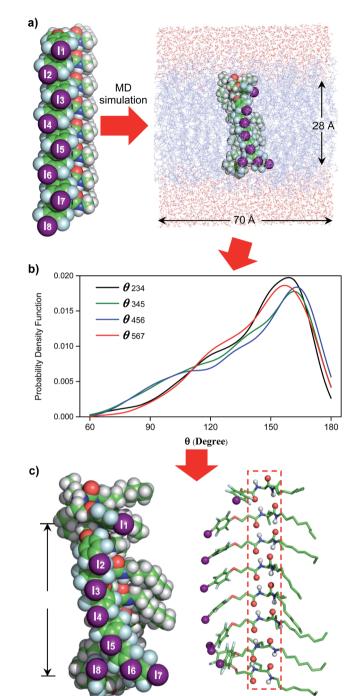


Fig. 4 (a) Embedding computationally optimized H-bonded L8 inside a simulation box of 70 Å (w) \times 70 Å (w) \times 74 Å (h), comprising 128 POPC molecules and 4794 water molecules. (b) Probability distribution patterns of $\theta_{234},\,\theta_{345},\,\theta_{456}$ and θ_{567} obtained after analysing 1000 structures. (c) A representative highly populated structure with the four θ angles approaching those having the highest probabilities shown in (b). POPC = 1-palmitoyl-2-oleoyl-sn-glycero-3-phosphocholine.

both the hydrophobic thickness of the POPC membrane and the distribution probability of angle θ (*e.g.*, the angle formed by the three immediately adjacent iodine atoms such as iodine atoms 2–4, Fig. 4a and b) using a probability density function.¹⁰ⁱ

To estimate the hydrophobic membrane thickness, the *Z*-coordinates along the membrane normal of all 254 ester O-atoms from 64 POPC molecules located either at the top or on the bottom layers were averaged. The separation distance between the two averaged *Z*-coordinates of O-atoms from the top and bottom layers was calculated with deduction of a van der Waals diameter of 3.1 Å for the O-atom. The same calculation was performed for all 1000 structural snapshots to derive the probability distribution of the hydrophobic membrane thickness (Fig. S12†). From these computations, 633 structures have a hydrophobic thickness of 27 to 29 Å, and the average thickness over 1000 structures is 28.1 Å, a value that could be corroborated by the experimentally and computationally determined values of 27.1 and 27.8 Å, respectively, for the POPC membrane.¹¹

Embedding eight molecules of **L8** in the lipid membrane shows that the first and eighth molecules are not aligned well with the central six molecules (Fig. 4a and c). The most highly populated angles for θ_{234} , θ_{345} , θ_{456} and θ_{567} are 159° , 162° , 163° and 157° , respectively, with a representative structure, closely capturing these angles, shown in Fig. 4c. In this highly populated structure, the largest intermolecular separation distance along the *Z* axis among the eight iodine atoms occurs between the $1^{\rm st}$ and $7^{\rm th}$ iodine atoms (25 Å). Consistent with the intermolecular separation of 5.0 Å in **Fmoc-Phe-C4** in the solid state (Fig. 1a)^{7a} and of 4.9 Å in the computationally optimized H-bonded **L8** (Fig. 1c), this separation distance of 25 Å suggests that six or seven molecules might be sufficient to span the hydrophobic core distance of 28.1 Å.

Through these MD simulations, the most important point to note is that, regardless of angle θ of varying magnitudes, all eight molecules of **L8** remain H-bonded to each other in all 1000 structures surveyed (Fig. 4c). This demonstrates the reliability of intermolecular H-bonds in linking six or seven anion-binding molecules together to form a self-assembled 1D pathway, which spans the hydrophobic membrane region to facilitate highly efficient anion transport across the membrane.

High activity in chloride transport

The use of halogen bonds to mediate anion transport was first explored by Matile, ^{2b,c,3i} with compounds 5 (EC₅₀ = $3.1 \, \mu M$)^{2c} and 8 (EC₅₀ = $0.88 \, \mu M$ in terms of chloride binding units)³ⁱ as the most active carrier and channel molecules among their own

Table 1 EC_{50} values of channels A10, L8 and L10 as well as other halogen bond-mediated chloride carriers (5–7) or channels (8) a

	A10	L8	L10	5^{b}	6 ^c	7 ^d	8 ^e
	2.37	0.39	0.93	3.6	260	68	0.88
n value ^f	3.5	3.6	3.1	3.3	3.6	1.9	0.8

 $[^]a$ Determined using Matile's conditions (100 mM NaCl, 10 mM HEPES, pH gradient from 7 to 8, and the total lipid concentration without cholesterol = 31.25 μ M; see ref. 2b, c and 3i). b EC $_{50}$ value from ref. 2c is 3.1 μ M with a Hill coefficient of 3.3. c From ref. 2c. d From ref. 2b. e From ref. 3i with the value normalized by the authors on the basis of eight halogen-binding units contained in channel 8. f Hill coefficient.

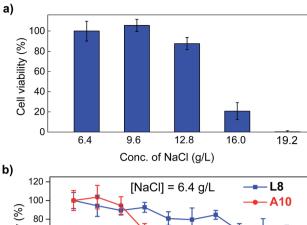
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categories, respectively. These EC_{50} values, including those of 6 (260 μ M) and 7 (68 μ M) presented in Table 1, were determined using cholesterol-free LUVs with a total lipid concentration of 31.3 μ M, while our assay conditions contain lipid and cholesterol at concentrations of 73.9 and 36.9 μ M, respectively. For a fair comparison, we have therefore re-determined the EC_{50} values of **A10**, **L8** and **L10** using Matile's conditions (100 mM NaCl, 10 mM HEPES, pH gradient from 7 to 8, and the total lipid concentration without cholesterol = 31.3 μ M).

To confirm that we are able to produce LUVs with properties similar to those prepared by Matile and his co-workers under the same assay conditions, we have performed Hill analysis on commercially available compound 5, and obtained an EC₅₀ value of 3.6 μ M with a Hill coefficient of 3.3 (Fig. S13†). These values are similar or identical to the reported values of EC₅₀ (3.1 μ M) and the Hill coefficient (3.3). ^{2c} Under these assay conditions, the EC₅₀ values of **A10**, **L8** and **L10** were

determined to be 2.37, 0.39 and 0.93 μ M (Table 1 and Fig. S13†), respectively. The most active L8 is 8.2 and 1.2 times more active than carrier 5 (3.6 μ M) and unimolecular channel molecule 8 (0.88 μ M in terms of effective chloride-binding units). When normalized based on the molecular weight, the ion transport activity of L8 is about 6.5 and 1.6 times those of 5 and 8, respectively. That L8 is more potent than unimolecular channel 8 is a clear indication of the excellent ability of the noncovalently associated peptidic scaffold in inducing linearly arrayed iodine atoms into a conformation that is more conducible to chloride transport than the rod-like scaffold in 8.

In addition, chloride carrier 5, when evaluated under our assay conditions containing 33 mol% cholesterol that makes membrane less fluid, 12 shows a moderate transport activity of 38% at 40 μM , and such moderate activity remains unchanged even when the concentration increases to 100 μM (Fig. S14†). These values suggest the ion transport activity of 5 to be at least 10 times lower than that of L8 having an EC $_{50}$ value of 3.6 μM in the cholesterol-containing environment.



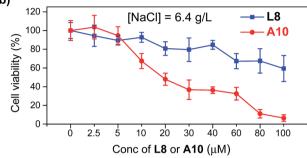


Fig. 5 (a) Sensitivity of human breast cancer cells BT-474 toward five different concentrations of NaCl from 6.4 to 19.2 g mL $^{-1}$ after 24 h. (b) Viabilities of BT-474 cells in the presence of various concentrations of L8 and A10 at [NaCl] = 6.4 g L $^{-1}$.

High anticancer activity

The above verified highly efficient chloride transport through a channel mechanism displayed by these artificial anion channels prompted us to examine the possibility of their uses in cancer chemotherapy.4f,5 Human breast cancer cells (BT-474, obtained from the American Type Culture Collection, USA) were cultured in Dulbecco's Modified Eagle Medium in the presence of up to 19.2 g L⁻¹ NaCl (6.4 g L⁻¹ is the typical concentration for cell growth) with concentrations of L8 or A10 varying from 0 to 100 μM (Fig. 5 and S15†). The viabilities of cells, determined after culturing the cells for 1 day at 37 °C with 5% CO₂ in the absence of channel molecules, show that BT-474 cells are sensitive to the sodium chloride concentration gradient (Fig. 5a). That is, compared to cell viability at [NaCl] = 6.4 g L^{-1} , 12, 79 and 99% fewer cells survive at [NaCl] of 12.8, 16.0 and 19.2 g L^{-1} , respectively. Continued testing shows that A10 is considerably more potent than L8 in inhibiting cell growth across all three different concentrations of NaCl (e.g., 6.4, 9.6 and 12.8 g L^{-1}) with an IC₅₀ value of 20 μ M for **A10** at [NaCl] = 6.4 g L⁻¹ (>100 μ M for L8, Fig. 5b and S15†). For comparison, highly effective anticancer agent cisplatin has an IC₅₀ value of 37 μM against the same BT-474 cells. Assuming all A10 molecules associate to form channels with each channel comprising six or more such molecules, the IC50 value in terms of effective channel concentration is lower than 3.3 μM.

Conclusions

In conclusion, we have developed a class of artificial chloride channels with high activity and good selectivity *via* directional self-assembly of polarized electron-deficient iodine atoms. While these linearly arrayed acidic iodine atoms, which form a series of chloride-binding and -transporting sites, are responsible for facilitating chloride transport likely via a multiion jumping mode, the high modularity of the monopeptide backbone enables rapid optimization of the transport activity and selectivity of channels in a combinatorial format. The identified channels A10, L8 and L10 all turn out to be very active with the best EC₅₀ values for chloride transport reaching 0.39 μM (1.2 mol% relative to lipid) and 3.6 μM (3.2 mol% relative to lipid/cholesterol) in cholesterol-free and -containing LUVs. In particular, the highly active A10 exhibits not only a fractional transport activity for chloride much better than other monovalent anions including bromide, but also an excellent inhibitory activity toward human breast cancer cells with an IC₅₀ value of 20 μM. Further refinement to enhance the anion selectivity and cytotoxicity of channels toward cancer cells

Conflicts of interest

is possible and expected.

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There are no conflicts to declare.

Acknowledgements

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