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Anion recognition using *meta*-substituted ureidocalix[4]arene receptors†

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Calix[4]arenes bearing urea units at the *meta* position(s) of the upper rim of the macrocyclic skeleton were prepared by the reaction of the corresponding amines with aryl isocyanates. As shown by the ¹H NMR and UV/vis titration experiments, these systems are capable of effectively complexing selected anions even in a highly competitive environment (such as DMSO-*d*₆). While the monoureido derivatives showed approximately the same complexation ability irrespective of the substitution (*para* vs. *meta* isomers), the bisureas at the upper rim demonstrated interesting differences in complexation. The *meta*, *meta* and *para,para* isomers were shown to prefer 2 : 1 complexes (anion : receptor) regardless of the anion tested, while the analogous *meta,para* isomer formed 1 : 1 complexes with strongly coordinated anions (e.g. H₂PO₄[−]) based on synchronous complexation by both ureido groups. This suggests that the regioselective introduction of urea units into the upper rim of calix[4]arene brings with it the possibility of “tuning” the complexation properties depending on the substitution pattern of the functional groups.

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Introduction

Recognition of anions, their targeted complexation and transport have become one of the cornerstones of modern supramolecular chemistry.¹ Selective complexation of anions is a topic with enormous application potential, ranging from the removal of toxic anions from the environment, through materials chemistry and catalysis to targeted complexation/decomplexation of anionic species in medicine. It is therefore not surprising that the research and development of new receptors for anions is becoming increasingly important, as evidenced by the huge number of recent articles and reviews² dealing with this topic.

Anion complexation uses many different basic strategies and approaches, but generally all receptors can be divided into charged and neutral systems. The charged ones usually rely on electrostatic interactions, which, however, are in principle not

very directional. These positively charged molecules can be exemplified by quaternary ammonium cages, protonated or alkylated azacrowns, cryptands and azamacrocycles, amidinium- and guanidinium-based receptors, *etc.*^{2,3}

The opposite of these substances are neutral receptors, based on highly directional interactions, such as hydrogen bonds (HBs) or halogen bonding.^{2d} Examples include receptors using amides/thioamides,^{2f,4} sulfonamides, ureas/thioureas,⁵ pyrroles,⁶ triazoles^{2a} or molecules with acidic CH bonds (e.g. bambusurils).⁷

The effectiveness of these highly directional interactions, particularly HBs, can be further enhanced by the thoughtful design of receptors, using several preorganized functional groups. Although there are a number of such molecules, a completely irreplaceable role is played by calix[*n*]arene derivatives, especially systems based on calix[4]arene.⁸ The possibility of tuning their 3D structures by simple alkylations of the lower rim of the macrocycle (phenolic hydroxyls) makes these molecules the preferred choice in the design and synthesis of new receptors.⁹ The existence of four basic conformations/atropisomers (*cone*, *partial cone*, *1,2-alternate*, and *1,3-alternate*) together with well-established derivatization procedures makes calix[4]arene an ideal molecular scaffold for the design of more complex supramolecular systems, including anion receptors.¹⁰

Anion receptors based on ureido/thioureido-calixarenes are very popular because these compounds are synthetically readily available. In particular, calix[4]arene-based receptors immobilised in the *cone* and *1,3-alternate* conformations have been documented many times in the literature¹¹ and some of

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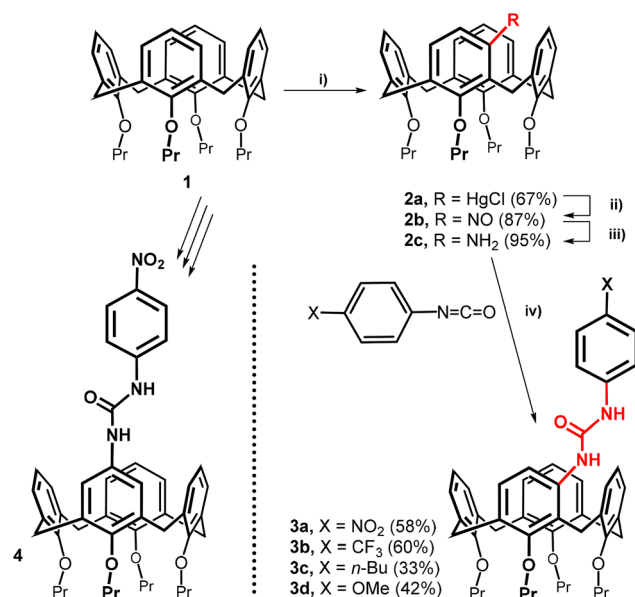
these systems show interesting complexation properties towards selected anions. All receptors described so far have one common feature – they use *para*-substituted phenolic sub-units of calixarenes, which is related to the methods of their preparation and the general reactivity of calixarenes. Only recently have direct procedures¹² for the *meta* substitution of the upper rim of calix[4]arene, allowing the introduction of various functional groups, been described.¹³

In this context, we realized that no anion receptors bearing urea groups at the *meta* position of the calixarene skeleton have been prepared so far (Fig. 1). It would therefore be interesting to find out whether this type of substitution exhibits the same behaviour as the already described *para*-isomers or whether the *meta* isomers lead to some new properties. In this paper, we report the first preparation of *meta*-ureido calix[4]arenes and the study of their complexation abilities towards selected anions in comparison with their *para*-substituted analogues.

Results and discussion

The synthesis of the corresponding *meta*-substituted aminocalix[4]arene **2c** was accomplished according to published procedures. Thus, direct monomercuration^{12a} (1 equiv. Hg(TFA)₂ in CHCl₃) of the starting tetrapropoxycalixarene **1** immobilized in the *cone* conformation and the subsequent reaction with aq. HCl provided **2a** in 67% yield. The subsequent transformation (isopentyl nitrite/HCl) of this organomercurial intermediate into nitroso derivative **2b** and the final reduction with RANEY® nickel and hydrazine afforded amine **2c** in a good overall yield¹⁴ (Scheme 1). Target receptors were obtained by the reaction of **2c** with appropriate commercially available aromatic isocyanates (*p*-X-C₆H₄-NCS, X = NO₂, *n*-Bu, OMe, CF₃) in dichloromethane at room temperature. Purification by preparative thin layer chromatography on silica gel provided **3a–d** in 33–60% isolated yields.

The structures of receptors **3a–d** were confirmed by the combination of HRMS and NMR techniques. Thus, the HRMS ESI⁺ of **3a** showed a peak at *m/z* = 973.4101, which is in good agreement with the [M + Na]⁺ (973.4106) ion predicted for the product. The multiplicity and splitting pattern of signals in



Scheme 1 Synthesis of *meta* substituted ureidocalix[4]arenes: (i) (1) Hg(TFA)₂/CHCl₃, (2) HCl; (ii) *i*-amyl nitrite/HCl; (iii) NH₂NH₂·H₂O/Ni(R); and (iv) CH₂Cl₂, RT.

the ¹H NMR spectrum (400 MHz, DMSO-*d*₆, 298 K) are fully consistent with the expected C₁ symmetry. Thus, the presence of four different signals for the terminal methyl groups (0.89, 0.90, 1.08 and 1.09 ppm) refers to the inherently chiral structure of the entire system. Also, the presence of two sets of doublets (2 × 4) for the axial and equatorial protons (with typical geminal coupling constants ~13–15 Hz) of the CH₂ bridges is consistent with the predicted structure. The singlets at 9.56 and 8.44 ppm (ureido N–H protons) and two doublets in the aromatic region at 8.19 and 7.70 ppm reflect the presence of the *p*-nitrophenylureido moiety within the molecule.

The structure of receptor **3a** was further confirmed by single crystal X-ray studies. Compound **3a** crystallized in the triclinic system with space group *P*₁; the unit cell contained two molecules of MeOH (crystallization solvent) that formed the 3a·2MeOH complex. As can be seen in Fig. 2a, the molecule adopts a typical *pinched cone* conformation, where two aromatic subunits are slightly tilted towards the interior of the cavity, while the other two opposing units are directed outwards. If we define the main plane of the molecule using the carbon atoms of the CH₂ bridging groups, the individual interplanar angles Φ are 125.75, 77.52, 125.63 and 82.54° starting clockwise from the subunit bearing the ureido group (see Fig. 2a). The urea function is located on the phenolic subunit pointing out of the cavity (Φ = 125.75), and the planar urea group itself is sharply twisted from the plane of the phenol at an angle of 67.47°.

An interesting binding motif was found within the crystal packing of substance **3a** (Fig. 2b). Two methanol molecules form a dimer by short hydrogen bonding (O–H...O distance = 2.101 Å) outside the calixarene cavity. The urea unit is connected by HB interactions of NH bonds to the first methanol;

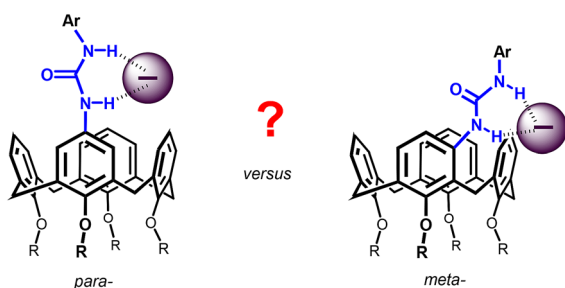


Fig. 1 Ureido receptors based on calix[4]arene: *para*- vs. *meta*-regioisomers.



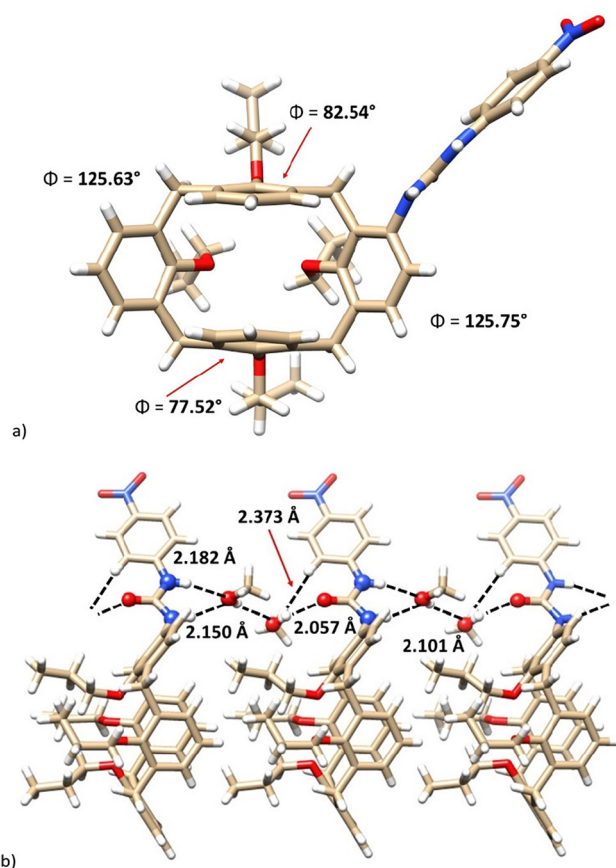


Fig. 2 X-ray structure of **3a**: (a) top view into the cavity with the corresponding interplanar angles and (b) hydrogen bonding motif of ureido functions.

the N–H...O distances are 2.182 and 2.150 Å. At the same time, the second methanol molecule is attached to another urea through HB interaction with the carbonyl (O–H...O=C distance is 2.057 Å) and by the interaction between the *ortho* CH bond of the adjacent nitro-substituted aromatic unit and the CH bond from the methanol (C_{ar}–H...H–C = 2.373 Å). The result is an endless belt of interconnected ureidocalixarenes, which have the same stereochemistry (identical enantiomers). Since the inherently chiral compound **3a** crystallizes as a racemic mixture, the same arrangement of the opposite enantiomer is found within the crystal (50 : 50 ratio). Finally, the opposite homochiral bands point towards each other through the ureido functions and are interconnected by the π – π interactions between the nitrophenyl units.

After evaluating interactions in the solid state, the properties of the systems in solution were subsequently monitored. Therefore, the ureido derivatives were studied by dilution experiments in order to confirm or disprove the possible self-aggregation behaviour of receptors.¹⁵ Here, the insufficient solubility did not allow us to reach concentrations higher than 10 mM even in DMSO (*e.g.* compound **3a** did not dissolve in concentrations exceeding 5 mM). This observation was general for all mono-*meta*-substituted receptors. Therefore, the

complexation abilities of receptors **3a–d** towards selected anions were studied using standard ¹H NMR and/or UV/vis titration experiments in a HB competitive environment. The anion solution was gradually added to the solution of **3a** in DMSO-*d*₆ to obtain various receptor : anion ratios (specific for each measurement, see the ESI†). In all cases, upon the addition of the anion, a significant down-field shift of ureido NH signals was observed, indicating complexation under fast exchange conditions (for example, see Fig. 3). When the formation of the host–guest complex was monitored by UV/vis, a redshift of receptor absorbance with a distinctive isosbestic point was observed.

The calculation of the corresponding complexation constants was based on the analysis of binding isotherms obtained from the urea NH protons, and/or the parts of the absorption curves where the changes in absorbance were the most significant. The non-linear curve fitting of experimental data was performed using Bindfit software.¹⁶ The stoichiometry of the complexes was determined by Job plot analysis¹⁷ or it was based on the Bindfit output, where the 1 : 1 model provided the best fit among all the stoichiometries (1 : 1, 1 : 2, and 2 : 1) tested.¹⁸

The complexation properties of receptor **3a** are summarized in Table 1, where they are also compared with the corresponding *para* isomer **4** (obtained as previously described¹⁹). The receptors showed no changes in the ¹H NMR spectra upon dilution of their DMSO-*d*₆ solutions (see Fig. S37 and S38†), which excludes receptor self-aggregation¹⁵ under the measurement conditions. As can be seen, the complexation constants *K* for benzoate, chloride and hydrogen sulfate anions are almost identical for both receptors, indicating no influence of the regioisomers (*meta* vs. *para*). The only distinct difference can be found for acetate and especially for dihydrogen phosphate, where the constant for the *para* isomer **4** is twice that of the *meta* analogue **3a** (compare *K*_{H₂PO₄}(**4**) = 2020 vs. *K*_{H₂PO₄}(**3a**) = 1060).

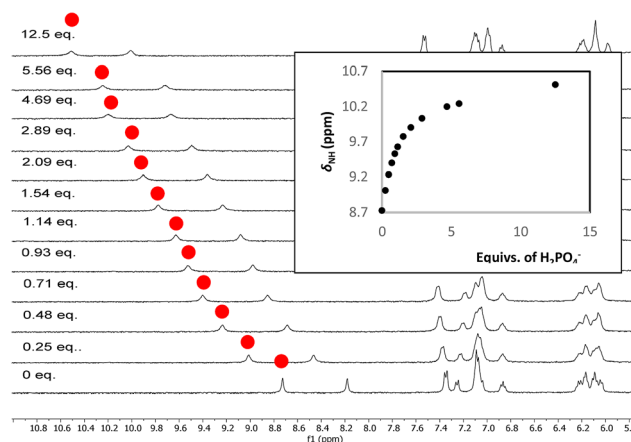


Fig. 3 ¹H NMR titration of receptor **3c** with TBA⁺ H₂PO₄[–] (DMSO-*d*₆, 400 MHz, 298 K), inset: the titration curve for the same system.



Table 1 Binding constants of receptors **3a** and **4** towards selected anions determined by ^1H NMR titration (400 MHz, $\text{DMSO}-d_6$, 298 K)

Anion ^a	K^b 3a	K 4
H_2PO_4^-	1060 ^c	2020 ^c
AcO^-	950 ^c	1320 ^c
BzO^-	830	845
Cl^-	50	35
HSO_4^-	4	6

^a All anions used as TBA (tetra-*n*-butylammonium) salts. ^b Error, when estimated, was <10%. ^c Determined by UV/vis titration in DMSO (298 K).

As several tested species can be considered as basic anions (H_2PO_4^- , benzoate, acetate), the interaction of **3a** with tetra-butylammonium hydroxide was also studied. As shown in Fig. S42,† deprotonation of the NH bond of receptor **3a** leads to a dramatic shift of the absorbance maximum, corresponding to a wavelength around 480 nm, which we never observed during the common titration experiments with any of the above-discussed anions (compare the record for acetate in Fig. S42†).

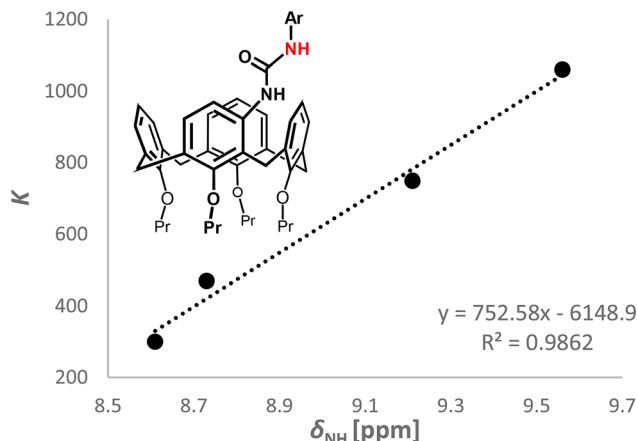
In order to demonstrate the effect of the substitution of the aryl residue on the complexation of the anions, a series of *meta* substituted receptors were further expanded to include **3b–d** and their binding abilities towards the dihydrogen phosphate anion were examined. As shown in Table 2, the strength of complexation is proportional to the electronic effect of the chosen substituent. Thus, the nitro group (the strongest EWG) exhibits the highest complexation constant $K_{\text{H}_2\text{PO}_4}(\mathbf{3a}) = 1060$, while the weakest binding was found for the methoxy derivative (EDG) $K_{\text{H}_2\text{PO}_4}(\mathbf{3d}) = 300$.

It is interesting that the chemical shift of the urea NH signal of the receptor (Table 2) can be correlated with the strength of the complex – the higher the shift, the higher the complexation constant. By plotting the complexation constants K against the chemical shift δ of the attached hydrogen, we even observed a linear dependence (see Fig. 4). A simple prediction of the binding constant for receptors of the same structural type, based on the chemical shift of urea NH signals, can be advantageous, e.g. in determining the correct choice of titration conditions for a new receptor.

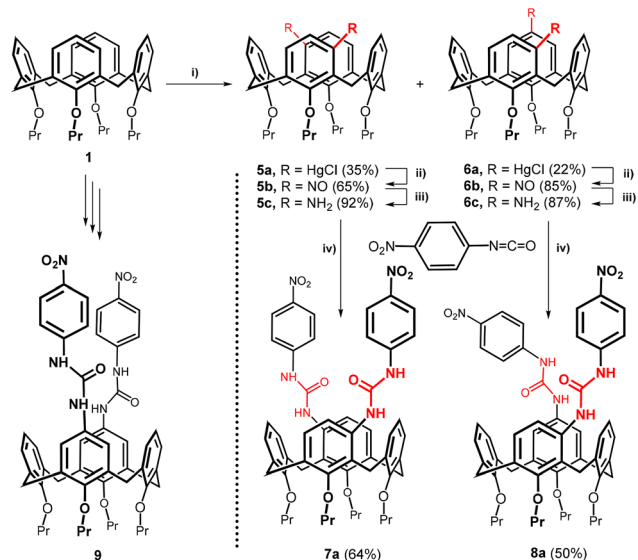
Table 2 Binding constants of receptors **3a–d** towards the H_2PO_4^- anion (^1H NMR titration, 400 MHz, $\text{DMSO}-d_6$, 298 K) and the NMR shifts of the corresponding NH signals

Receptor	K^a	δ_{NH} [ppm]
3a	1060 ^b	9.56
3b	750	9.21
3c	470	8.73
3d	300	8.61

^a Error, when estimated, was <10%. ^b Determined by UV/vis titration in DMSO (298 K).

**Fig. 4** Dependence of the K values on the chemical shifts of NH (compounds **3a–d**).

In order to study the effect of system preorganization on the complexation behaviour, we attempted to introduce two urea functionalities into the upper rim of calix[4]arene using an organomercury protocol described by our group.^{12b} Briefly, the starting tetrapropoxycalix[4]arene **1** was reacted with two equivalents of $\text{Hg}(\text{TFA})_2$ in chloroform to provide a mixture of two distally dimercurated calixarenes: *meta,meta* **5a** and *meta,para* **6a** isomers in an approx. 3 : 2 ratio (Scheme 2). Without any isolation, the crude reaction product was reacted with isoamyl nitrite/aq. HCl to yield a mixture of nitroso derivatives **5b** and **6b**, which was smoothly separated by column chromatography on silica gel.²⁰ The resulting amines **5c** and **6c** were obtained by reduction with hydrazine hydrate in the presence of RANEY®-nickel. Since the corresponding nitro derivative **3a**

**Scheme 2** Synthesis of *meta* substituted diureidocalix[4]arenes **7a** and **8a**: (i) (1) $\text{Hg}(\text{TFA})_2/\text{CDCl}_3$, (2) HCl; (ii) isoamyl nitrite/HCl; (iii) $\text{NH}_2\text{NH}_2 \cdot \text{H}_2\text{O}/\text{Ni}(\text{R})$; and (iv) CH_2Cl_2 , RT.

showed the highest complexation constants, the reaction with *p*-nitrophenyl isocyanate was used in both cases. The respective distally substituted diureido isomers **7a** and **8a** were isolated in 64 and 50% yields, respectively.

The ^1H NMR spectra of compounds **7a** and **8a** (400 MHz, $\text{DMSO}-d_6$, 298 K) reflect the expected symmetry of the molecules well. Thus, receptor **7a** shows 2×2 doublets for the CH_2 bridging groups ($J \sim 13$ Hz) together with two NH signals (9.59 and 8.54 ppm) of urea groups, suggesting the C_2 symmetry of **7a**. On the other hand, the splitting pattern and number of signals in the ^1H NMR spectrum of **8a** (e.g. 2×4 doublets for equatorial and axial CH bonds of the bridging CH_2 units, four different signals for NH bonds (9.55, 9.36, 8.65 and 8.42 ppm)) are consistent with the predicted C_1 symmetry.

Once again, solubility issues play an important role, limiting the range of values of potential binding constants under correct conditions (for the corresponding dilution experiments in $\text{DMSO}-d_6$, see Fig. S39–S41†). For receptor **7a**, the clear solution was not obtained until the concentration decreased below 2 mM; therefore only strongly interacting anions were studied. The ^1H NMR titrations of compound **7a** (*meta,meta*) and the Job plot analysis revealed that 2 : 1 complexes (anion : calix) are formed for all anions investigated (H_2PO_4^- , BzO^- and AcO^-). It is therefore obvious that the individual ureido groups function independently and there is no mutual cooperation between them. Thus, each urea grabs its own anionic species, leading to the aforementioned 2 : 1 stoichiometry. As shown in Table 3, the corresponding binding constants K (1 : 1) and the overall binding constants β (2 : 1) were calculated using a non-cooperative model as the most appropriate choice.

The corresponding *para,para* isomer **9**¹⁹ was used for direct comparison of the complexation properties of the receptors. As shown in Table 3, this isomer behaves very similarly to the preceding one. Although dihydrogen phosphate is captured somewhat more strongly, even in this case, a 2 : 1 complex is formed with independent complexation of both urea moieties.

Finally, the complexation study of receptor **8a** (*meta,para*) yielded unexpected results. With basic anions, this isomer has been shown to form complexes with a 1 : 1 stoichiometry (Table 4). This suggests that in this case there is a cooperative binding of one anion by both urea units. This leads to the fact that the complexation constant towards dihydrogen phosphate $K_{\text{H}_2\text{PO}_4}(\mathbf{8a}) = 13\,600$ is by far the highest among all measured

Table 4 Binding constants K (1 : 1) and the overall binding constants β (2 : 1) of receptor **8a** towards selected anions^a (^1H NMR titration, 400 MHz, $\text{DMSO}-d_6$, 298 K)

Anion ^a	K [1 : 1]	β [2 : 1] ^b
H_2PO_4^-	13 600 ^c	—
BzO^-	2450 ^c	—
AcO^-	4170 ^c	—
Cl^-	60	900
HSO_4^-	15	56

^a Used as TBA (tetra-*n*-butylammonium) salts. ^b 2 : 1 stoichiometry (anion : calix), fitted by a non-cooperative model. ^c Determined by UV/vis titration in DMSO (298 K).

receptors – essentially an order of magnitude higher than that of the corresponding monoureido receptor **3a** ($K_{\text{H}_2\text{PO}_4}(\mathbf{3a}) = 1060$). The same trend can be seen for the acetate anion although the effect was not so pronounced. The results for BzO^- were somewhat in the middle (between cooperating and independent action of urea's sites). However, as no mathematical tool was able to combine these actions in a single fitting process, we decided to fit the raw data by a simple 1 : 1 stoichiometry. On the other hand, anions that cannot be captured synchronously from both functional groups (chloride and hydrogen sulphate, Table 4) again clearly form complexes with a 2 : 1 stoichiometry (Fig. 5).

All *meta*-substituted receptors represent inherently chiral systems that could be useful for chiral anion recognition. To test this hypothesis, we attempted to resolve racemic mixtures of compounds **3a**, **7a**, and **8a** using chiral phase chromatography. Unfortunately, all our efforts did not lead to the goal and it was not possible to separate the urea derivatives into pure enantiomers.

Therefore, we focused on other properties of these receptors. Since our compounds carry the $-\text{NO}_2$ group in their structure, we tried to evaluate their electrochemical reduction. The combination of steady state (RDE) and dynamic methods (CV) provided sufficient information about the reduction transformation. Receptors **3a** and **4** were shown to be reduced in three well-separated steps (Fig. 6a and S53†), regardless of the substitution (*meta* vs. *para*).

This behaviour has been already reported for various compounds bearing acidic hydrogens together with a nitro group

Table 3 Binding constants K (1 : 1) and the overall binding constants β (2 : 1) of receptor **7a** towards selected anions^a (^1H NMR titration, 400 MHz, $\text{DMSO}-d_6$, 298 K)

Anion ^a	K [1 : 1]	β [2 : 1] ^b
H_2PO_4^-	1660	6.89×10^5
BzO^-	1440	5.18×10^5
AcO^-	1160	3.36×10^5
Comp 9 / H_2PO_4^-	3350	2.81×10^6

^a Used as TBA (tetra-*n*-butylammonium) salts. ^b 2 : 1 stoichiometry (anion : calix), fitted by a non-cooperative model.

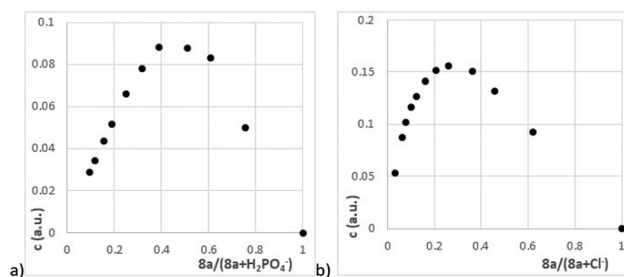


Fig. 5 Job plot (^1H NMR, 400 MHz, $\text{DMSO}-d_6$, 298 K) recorded for receptor **8a** (a) with H_2PO_4^- and (b) with Cl^- anions.



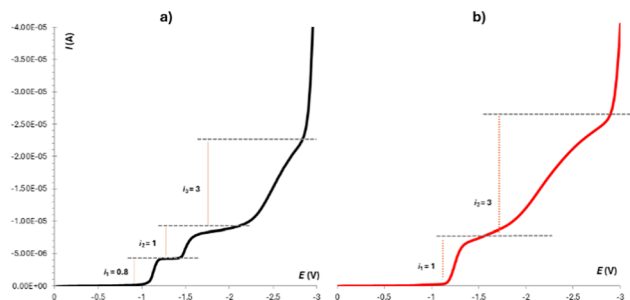


Fig. 6 Linear sweep voltammetry of **3a** (9×10^{-4} M) in DMSO (0.1 M TBAPF₆) (W = RDE = glassy carbon – \varnothing 1 mm, Ref = SCE, Aux = Pt) recorded with a scan rate of 10 mV s^{-1} using a rotating rate of 100 s^{-1} : (a) in the absence of any anion and (b) in the presence of H_2PO_4^- .

within the same structure. The mechanism describing this transformation is known in the literature as the self-protonation mechanism.²¹ Interestingly, we found out that the urea's hydrogens might be “protected” upon the addition of an anion (H_2PO_4^-), when a strong complex is formed by means of hydrogen bonds from the urea unit (NH). This suppresses the possibility of self-protonation, which results in the common two-step reduction (Fig. 6b).²² This property could therefore be used for electrochemical sensing of phosphates.

Conclusions

In conclusion, calix[4]arenes bearing ureido units at the *meta* position(s) of the upper rim of the calixarene skeleton were prepared. The ^1H NMR and UV/vis titrations proved that these systems are capable of effectively complexing selected anions even in a highly competitive environment (such as $\text{DMSO}-d_6$). The monoureido derivatives showed approximately the same complexation ability irrespective of the substitution (*para* vs. *meta* isomers). On the other hand, the bisureas at the upper rim demonstrated interesting differences in complexation. While the *meta,meta* and *para,para* isomers preferred 2 : 1 complexes (e.g. with the dihydrogen phosphate anion), a similar *meta,para* isomer formed a 1 : 1 complex based on the synchronous complexation of the anion by both ureido groups. This suggests that the regioselective introduction of a urea moiety into the upper rim of calix[4]arene brings with it the possibility of “tuning” the complexation properties depending on the mutual arrangement of the functional groups (substitution pattern).

Experimental

General experimental procedures

All chemicals were obtained from commercial sources and used as received without further purification. Solvents were dried and distilled using conventional methods. TLC was performed on Merck foil sheets with silica gel 60 F254 (Merck). Column chromatography was performed on silica gel 60 with particle size 0.063–0.200 mm (Merck). Preparative thin-layer

chromatography was performed on self-prepared glass plates ($25 \times 25 \text{ cm}$) covered by silica gel 60 GF₂₅₄ containing CaSO_4 (Merck). ^1H and ^{13}C NMR spectra were obtained using Agilent 400-MR DDR2, JEOL-ECZL400G (^1H : 400 MHz, ^{13}C : 100 MHz) and Bruker Avance DRX 500 (^1H : 500 MHz, ^{13}C : 125 MHz) spectrometers at 298 K. Deuterated solvents used are indicated in each case. Chemical shifts are reported as δ values in parts per million (ppm) and were referenced to the residual peak of the solvent or TMS as an internal standard; coupling constants (J) are expressed in Hz. NMR data were processed and displayed using MestReNova and TopSpin software. Melting points were measured on a Heitzsch Mikroskop – Polytherm A (Wagner & Munz, Germany) and are not corrected. The mass spectra analyses were performed on a Q-TOF (Micromass) spectrometer, using ESI ionisation in positive mode. Infrared spectra were measured on an FT-IR spectrometer Nicolet iS50 (Thermo-Nicolet, USA) connected with a heatable Golden Gate Diamante ATR-Unit GladiATR (PIKE, USA) in KBr. 64 Scans for one spectrum were co-added at a spectral resolution of 4 cm^{-1} . The spectra were processed using Omnic 9 (Thermo-Nicolet Instruments Co., USA) with baseline correction.

Synthetic procedures

General procedure for the synthesis of mono- and diureido-calix[4]arenes. The corresponding calixarene was dissolved in 7–12 mL of DCM under an argon atmosphere. The isocyanate (2.1 eq. for disubstituted derivatives, 1.1 eq. for monosubstituted derivatives) was added to the solution. The mixture was stirred overnight at 25°C and then 5 mL of methanol was added. The reaction mixture was stirred another 20 minutes at 25°C . The solvent was removed under reduced pressure from the reaction mixture. The crude product was isolated by preparative thin layer chromatography on silica gel.

4-{N'-(4-Nitrophenyl)ureido}-25,26,27,28-tetrapropoxycalix[4]arene (cone) (3a). Calixarene **3a** was prepared according to the general procedure for ureido derivatives by reacting calixarene **2c** (0.141 g, 0.230 mmol) and 4-nitrophenylisocyanate (0.041 g, 0.253 mmol) in 10 mL dry DCM. The product was purified by preparative TLC (eluent acetone : cyclohexane 10 : 90) to give the title compound **3a** as a yellow solid (0.102 g, 58%), m.p. $143\text{--}146^\circ\text{C}$. ^1H -NMR (500 MHz, $\text{DMSO}-d_6$): δ 9.54 (s, 1H, $-\text{NH}-$), 8.43 (s, 1H, $-\text{NH}-$), 8.18 (d, $J = 9.10 \text{ Hz}$, 2H, Ar-H), 7.68 (d, $J = 9.10 \text{ Hz}$, 2H, Ar-H), 7.20 (d, $J = 8.2 \text{ Hz}$, 1H, Ar-H), 7.07 (td, $J = 7.0 \text{ Hz}$, $J = 1.5 \text{ Hz}$, 2H, Ar-H), 7.06 (d, $J = 8.0 \text{ Hz}$, 1H, Ar-H), 6.85 (t, $J = 7.5 \text{ Hz}$, 1H, Ar-H), 6.23 (t, $J = 7.5 \text{ Hz}$, 1H, Ar-H), 6.18 (t, $J = 7.5 \text{ Hz}$, 1H, Ar-H), 6.16 (d, $J = 7.5 \text{ Hz}$, 1H, Ar-H), 6.11 (dd, $J = 7.5 \text{ Hz}$, $J = 1 \text{ Hz}$, 1H, Ar-H), 6.08 (dd, $J = 7.5 \text{ Hz}$, $J = 1 \text{ Hz}$, 1H, Ar-H), 6.04 (d, $J = 7.5 \text{ Hz}$, 1H, Ar-H), 4.34 (d, $J = 13.5 \text{ Hz}$, 1H, Ar- CH_2 -Ar), 4.33 (d, $J = 13.0 \text{ Hz}$, 1H, Ar- CH_2 -Ar), 4.31 (d, $J = 13.5 \text{ Hz}$, 1H, Ar- CH_2 -Ar), 4.15 (d, $J = 13.5 \text{ Hz}$, 1H, Ar- CH_2 -Ar), 4.00–3.87 (m, 4H, $-\text{O}-\text{CH}_2-$), 3.65–3.59 (m, overlap, 4H, $-\text{O}-\text{CH}_2-$), 3.49 (d, $J = 13.5 \text{ Hz}$, 1H, Ar- CH_2 -Ar), 3.16 (d, 1H + 2H, Ar- CH_2 -Ar), 2.02–1.88 (m, 4H, $-\text{O}-\text{CH}_2-\text{CH}_2-$), 1.83 (sextet, $J = 7.2 \text{ Hz}$, 4H, $-\text{O}-\text{CH}_2-\text{CH}_2-$), 1.08 (t, $J = 7.3 \text{ Hz}$, 3H, $-\text{O}-(\text{CH}_2)_2-\text{CH}_3$), 1.07 (t, $J = 7.3 \text{ Hz}$, 3H, $-\text{O}-(\text{CH}_2)_2-\text{CH}_3$), 0.90 (t, $J = 7.5 \text{ Hz}$, 3H, $-\text{O}-(\text{CH}_2)_2-\text{CH}_3$), 0.88 (t, $J = 7.5 \text{ Hz}$, 3H, $-\text{O}-$



(CH₂)₂-CH₃). ¹³C{¹H} NMR (125 MHz, DMSO-*d*₆): δ 157.7, 157.3, 154.9, 154.8, 152.7, 146.7, 136.3, 136.2, 134.7, 132.9, 132.83, 132.80, 132.6, 132.5, 130.3, 128.7, 128.6, 127.9, 127.3, 127.2 (2×), 126.7, 125.2 (2×), 121.9, 121.64, 121.59, 118.3, 117.2 (2×), 76.6, 76.4, 76.1, 75.9, 30.2, 30.1, 29.9, 24.0, 23.1, 23.0, 22.6, 22.5, 10.69, 10.67, 9.77, 9.76. HRMS ESI⁺: (C₄₇H₅₃N₃O₇) *m/z* calcd 794.3776 [M + Na]⁺, found 794.3783 [M + Na]⁺. IR (KBr) ν 2961, 2931, 2874, 1717, 1667, 1594, 1504, 1455, 1329, 1300, 1245, 1207, 1175, 1111, 1087, 1037, 1005, 964, 850, 753, 691 cm⁻¹.

4-{N'-(4-Trifluoromethylphenyl)ureido}-25,26,27,28-tetrapropoxycalix[4]arene (cone) (**3b**). Calixarene **3b** was prepared according to the general procedure for ureido derivatives by reacting calixarene **2c** (0.116 g, 0.190 mmol) and 4-trifluoromethylphenylisocyanate (0.03 mL, 0.21 mmol) in 7 mL dry DCM. The product was purified by preparative TLC (eluent DCM:cyclohexane 90:10) and then once again (eluent acetone:cyclohexane 15:85) to give the title compound **3b** as a white solid (0.09 g, 60%), m.p. 170–173 °C. ¹H-NMR (500 MHz, DMSO-*d*₆): δ 9.22 (s, 1H, -NH-), 8.32 (s, 1H, -NH-), 7.65 (d, *J* = 7.8 Hz, 2H, Ar-H), 7.61 (d, *J* = 7.8 Hz, 2H, Ar-H), 7.21 (d, *J* = 8.2 Hz, 1H, Ar-H), 7.07 (td, *J* = 7.7 Hz, *J* = 1.1 Hz, 2H, Ar-H), 7.06 (d, *J* = 8.2 Hz, 1H, Ar-H), 6.85 (t, *J* = 7.5 Hz, 1H, Ar-H), 6.22 (t, *J* = 7.5 Hz, 1H, Ar-H), 6.17 (t, *J* = 8.5 Hz, 1H, Ar-H), 6.16 (d, *J* = 7.6 Hz, 1H, Ar-H), 6.10 (d, *J* = 7.6 Hz, 1H, Ar-H), 6.08 (d, *J* = 7.6 Hz, 1H, Ar-H), 6.03 (d, *J* = 7.6 Hz, 1H, Ar-H), 4.34 (d, *J* = 13.5 Hz, 1H, Ar-CH₂-Ar), 4.33 (d, *J* = 13.5 Hz, 1H, Ar-CH₂-Ar), 4.31 (d, *J* = 13.5 Hz, 1H, Ar-CH₂-Ar), 4.14 (d, *J* = 13.5 Hz, 1H, Ar-CH₂-Ar), 4.03–3.86 (m, 4H, -O-CH₂-), 3.63 (t, *J* = 6.3 Hz, 2H, -O-CH₂-), 3.62 (t, *J* = 6.3 Hz, 2H, -O-CH₂-), 3.50 (d, *J* = 13.5 Hz, 1H, Ar-CH₂-Ar), 3.16 (~d, overlap, 1H + 2H, Ar-CH₂-Ar), 2.01–1.88 (m, 4H, -O-CH₂-CH₂-), 1.82 (sextet, *J* = 7.2 Hz, 4H, -O-CH₂-CH₂-), 1.08 (t, *J* = 7.2 Hz, 3H, -O-(CH₂)₂-CH₃), 1.07 (t, *J* = 7.2 Hz, 3H, -O-(CH₂)₂-CH₃), 0.89 (t, *J* = 7.6 Hz, 3H, -O-(CH₂)₂-CH₃), 0.88 (t, *J* = 7.6 Hz, 3H, -O-(CH₂)₂-CH₃). ¹³C{¹H} NMR (125 MHz, DMSO-*d*₆): δ 157.7, 157.3, 154.9, 154.8, 153.0, 143.8, 136.3, 136.2, 135.0, 132.9, 132.83, 132.79, 132.2, 130.1, 128.7, 128.6, 127.8, 127.2, 127.1 (*J* = 4 Hz), 126.7, 126.09, 126.07, 124.6 (*J* = 272 Hz), 121.9, 121.64, 121.59, 121.4 (*J* = 32 Hz), 118.1, 117.5, 76.6, 76.4, 76.1, 75.9, 30.2, 30.1, 29.9, 23.9, 23.1, 23.0, 22.6, 22.5, 10.69, 10.67, 9.76. HRMS-ESI⁺: (C₄₈H₅₃F₃N₂O₅) *m/z* calcd 817.3799 [M + Na]⁺, found 817.3803 [M + Na]⁺. IR (KBr) ν 2963, 2933, 2875, 1659, 1606, 1550, 1457, 1324, 1208, 1118, 1069, 1008, 965, 840, 760 cm⁻¹.

4-{N'-(4-Butylphenyl)ureido}-25,26,27,28-tetrapropoxycalix[4]arene (cone) (**3c**). Calixarene **3c** was prepared according to the general procedure for ureido derivatives by reacting calixarene **2c** (0.112 g, 0.185 mmol) and 4-butylphenylisocyanate (0.036 g, 0.203 mmol) in 6 mL dry DCM. The product was purified by preparative TLC (eluent acetone:cyclohexane 10:90) to give the title compound **3c** as a yellow solid (0.048 g, 33%), m.p. 230–232 °C. ¹H-NMR (500 MHz, DMSO-*d*₆): δ 8.69 (s, 1H, -NH-), 8.15 (s, 1H, -NH-), 7.34 (d, *J* = 8.0 Hz, 2H, Ar-H), 7.25 (d, *J* = 8.2 Hz, 1H, Ar-H), 7.12–7.00 (m, overlap, 5H, Ar-H), 6.86 (t, *J* = 7.5 Hz, 1H, Ar-H), 6.26–6.12 (m, 3H, Ar-H), 6.09 (d, *J* = 7.5 Hz, 1H, Ar-H), 6.07 (d, *J* = 7.5 Hz, 1H, Ar-H), 6.03 (d, *J* =

7.5 Hz, 1H, Ar-H), 4.34 (d, *J* = 13.3 Hz, 2H, Ar-CH₂-Ar), 4.30 (d, *J* = 13.3 Hz, 1H, Ar-CH₂-Ar), 4.03–3.85 (m, 4H, -O-CH₂-), 3.66–3.58 (m, 4H, -O-CH₂-), 3.50 (d, *J* = 13.3 Hz, 1H, Ar-CH₂-Ar), 3.15 (d, *J* = 13.3 Hz, 2H, Ar-CH₂-Ar), 3.14 (d, *J* = 13.3 Hz, 2H, Ar-CH₂-Ar), 2.01–1.88 (m, 4H, -O-CH₂-CH₂-), 1.87–1.77 (m, 4H, -CH₂-), 1.55–1.46 (m, 2H, -CH₂-), 1.34–1.24 (m, 2H, -CH₂-), 1.08 (t, *J* = 7.50 Hz, 6H, -CH₃), 1.08 (overlap, 1H, -CH₂-), 0.92–0.83 (m, 9H, -CH₃ + 1H, -CH₂-). ¹³C{¹H} NMR (125 MHz, DMSO-*d*₆): δ 157.7, 157.3, 154.8, 154.7, 153.2, 137.7, 136.34, 136.26, 135.6, 135.4, 132.9, 132.8, 132.7 (2C), 131.6, 129.5, 128.71, 128.65, 128.5 (2C), 127.8, 127.2, 127.13, 127.09, 126.7, 121.9, 121.64, 121.58, 118.0 (2C), 117.7, 76.6, 76.5, 76.1, 75.9, 34.1, 33.3, 30.2, 30.1, 29.9, 23.7, 23.1, 23.0, 22.6, 22.5, 21.7, 13.8, 10.7 (2C), 9.8 (2C). HRMS-ESI⁺: (C₅₁H₆₂N₂O₅) *m/z* calcd 805.4551 [M + Na]⁺, found 805.4539 [M + Na]⁺. IR (KBr) 2960, 2927, 2873, 1655, 1605, 1550, 1456, 1206, 1088, 967, 759 cm⁻¹.

4-{N'-(4-Methoxyphenyl)ureido}-25,26,27,28-tetrapropoxycalix[4]arene (cone) (**3d**). Calixarene **3d** was prepared according to the general procedure for ureido derivatives by reacting calixarene **2c** (0.097 g, 0.159 mmol) and 4-methoxyphenylisocyanate (0.02 mL, 0.176 mmol) in 5 mL dry DCM. The product was purified by preparative TLC (eluent acetone:cyclohexane 20:80) to give the title compound **3d** as a white solid (0.051 g, 42%), m.p. 198–200 °C. ¹H-NMR (500 MHz, DMSO-*d*₆): δ 8.64 (s, 1H, -NH-), 8.14 (s, 1H, -NH-), 7.34 (d, *J* = 9.0 Hz, 2H, Ar-H), 7.23 (d, *J* = 8.2 Hz, 1H, Ar-H), 7.11–7.00 (m, overlap, 3H, Ar-H), 6.86 (t, *J* = 7.5 Hz, 1H, Ar-H), 6.85 (d, *J* = 9.0 Hz, 2H, Ar-H), 6.24–6.11 (m, 3H, Ar-H), 6.08 (td, *J* = 7.8 Hz, *J* = 1.2 Hz, 2H, Ar-H), 6.02 (dd, *J* = 7.7 Hz, *J* = 1.2 Hz, 1H, Ar-H), 4.33 (d, *J* = 13.3 Hz, 2H, Ar-CH₂-Ar), 4.30 (d, *J* = 13.3 Hz, 1H, Ar-CH₂-Ar), 4.12 (d, *J* = 13.3 Hz, 1H, Ar-CH₂-Ar), 4.04–3.84 (m, 4H, -O-CH₂-), 3.70 (s, 3H, -Ar-CH₃), 3.66–3.57 (m, 4H, -O-CH₂-), 3.50 (d, *J* = 13.3 Hz, 1H, Ar-CH₂-Ar), 3.16 (d, *J* = 13.0 Hz, 2H, Ar-CH₂-Ar), 3.13 (d, *J* = 13.0 Hz, 2H, Ar-CH₂-Ar), 2.04–1.87 (m, 4H, -CH₂-), 1.87–1.76 (m, 4H, -CH₂-), 1.08 (t, *J* = 7.5 Hz, 6H, -CH₃), 0.88 (t, *J* = 7.5 Hz, 6H, -CH₃), 0.87 (t, *J* = 7.5 Hz, 6H, -CH₃). ¹³C NMR (101 MHz, DMSO-*d*₆) δ 157.7, 157.3, 154.8, 154.8, 154.2, 153.4, 136.3, 136.2, 135.7, 133.1, 132.9, 132.8, 132.74, 132.70, 131.5, 129.5, 128.7, 128.6, 127.7, 127.2, 127.1, 126.7, 121.8, 121.6, 121.6, 119.6, 117.6, 114.0, 76.6, 76.4, 76.1, 75.9, 55.1, 30.2, 30.1, 29.9, 23.7, 23.1, 23.0, 22.6, 22.5, 10.7, 9.8. HRMS-ESI⁺: (C₄₈H₅₆N₂O₆) *m/z* calcd 779.4030 [M + Na]⁺, found 779.4026 [M + Na]⁺. IR (KBr) 2961, 2933, 2874, 1647, 1510, 1481, 1455, 1229, 1207, 1089, 1036, 1006, 966, 826, 759 cm⁻¹.

4,16-Bis{N'-(4-nitrophenyl)ureido}-25,26,27,28-tetrapropoxycalix[4]arene (cone) (**7a**). Calixarene **7a** was prepared according to the general procedure for ureido derivatives by reacting calixarene **5c** (0.0535 g, 0.0851 mmol) and 4-nitrophenylisocyanate (2.1 eq. 0.029 g, 0.178 mmol) in 7 mL dry DCM. The product was precipitated from acetonitrile to give the title compound **7a** as a yellow solid (0.052 g, 64%), m.p. 255–260 °C. ¹H-NMR (500 MHz, DMSO-*d*₆): δ 9.57 (s, 1H, -NH-), 8.53 (s, 1H, -NH-), 8.19 (d, *J* = 9.20 Hz, 4H, Ar-H), 7.69 (d, *J* = 9.20 Hz, 4H, Ar-H), 7.26 (d, *J* = 7.75 Hz, 2H, Ar-H), 7.13 (d, *J* = 7.75 Hz, 2H, Ar-H), 6.17 (t, *J* = 7.66 Hz, 2H, Ar-H), 6.06 (d, *J* = 7.45 Hz, 2H, Ar-H),



5.94 (d, $J = 7.45$ Hz, 2H, Ar-H), 4.30 (d, $J = 13.3$ Hz, 2H, Ar-CH₂-Ar), 4.13 (d, $J = 13.3$ Hz, 2H, Ar-CH₂-Ar), 4.05–3.98 (m, 2H, -O-CH₂), 3.96–3.88 (m, 2H, -O-CH₂), 3.61 (t, $J = 6.50$ Hz, 4H, -O-CH₂), 3.50 (d, $J = 13.3$ Hz, 2H, Ar-CH₂-Ar), 3.17 (d, $J = 13.3$ Hz, 2H, Ar-CH₂-Ar), 1.97 (sextet, $J = 7.61$ Hz, 4H, -O-CH₂-CH₂-), 1.83 (sextet, $J = 7.61$ Hz, 2H, -O-CH₂-CH₂-), 1.09 (t, $J = 7.3$ Hz, 6H, -O-(CH₂)₂-CH₃), 0.88 (t, $J = 7.3$ Hz, 6H, -O-(CH₂)₂-CH₃). ¹³C{¹H} NMR (125 MHz, DMSO-*d*₆, 333 K): δ 157.7, 154.5, 152.5, 146.4, 140.8, 134.7, 132.5, 132.3, 132.3, 130.2, 127.6, 126.9, 126.3, 124.8, 121.5, 117.8, 117.1, 76.2, 75.7, 29.8, 23.6, 22.8, 22.2, 10.4, 9.4. HRMS-ESI⁺: (C₅₄H₅₈N₆O₁₀) m/z calcd 973.4106 [M + Na]⁺; found 973.4101 [M + Na]⁺. IR (KBr) 2965, 2930, 2874, 1664, 1555, 1510, 1482, 1454, 1331, 1218, 1176, 1111, 1085, 965, 851, 752 cm⁻¹.

4,17-Bis[N'-(4-nitrophenyl)ureido]-25,26,27,28-tetrapropoxy-calix[4]arene 8a. Calixarene **8a** was prepared according to the general procedure for ureido derivatives by reacting calixarene **6c** (0.122 g, 0.193 mmol) and 4-nitrophenylisocyanate (2.1 eq. 0.066 g, 0.405 mmol) in 12 mL dry DCM. The product was purified by preparative TLC (eluent acetone : cyclohexane 30 : 70) to give the title compound **8a** as a yellow solid (0.09 g, 50%), m.p. 220–225 °C. ¹H-NMR (500 MHz, DMSO-*d*₆): δ 9.55 (s, 1H, -NH-), 9.36 (s, 1H, -NH-), 8.65 (s, 1H, -NH-), 8.41 (s, 1H, -NH-), 8.18 (d, $J = 9.10$ Hz, 2H, Ar-H), 8.16 (d, $J = 9.10$ Hz, 2H, Ar-H), 7.70 (d, $J = 9.10$ Hz, 2H, Ar-H), 7.68 (d, $J = 9.10$ Hz, 2H, Ar-H), 7.19 (d, $J = 8.2$ Hz, 1H, Ar-H), 7.13 (m, 2H, Ar-H), 7.03 (d, $J = 8.2$ Hz, 1H, Ar-H), 6.28 (t, $J = 7.4$ Hz, 1H, Ar-H), 6.23–6.18 (m, overlap, 4H, Ar-H), 6.10 (dd, $J = 7.0$ Hz, $J = 1$ Hz, 1H, Ar-H), 4.33 (d, $J = 13.0$ Hz, 1H, Ar-CH₂-Ar), 4.32 (d, $J = 13.0$ Hz, 1H, Ar-CH₂-Ar), 4.31 (d, $J = 13.0$ Hz, 1H, Ar-CH₂-Ar), 4.15 (d, $J = 13.0$ Hz, 1H, Ar-CH₂-Ar), 4.02–3.94 (m, 1H, -O-CH₂-), 3.94–3.96 (m, 3H, -O-CH₂-), 3.68–3.59 (m, 4H, -O-CH₂-), 3.49 (d, $J = 13.0$ Hz, 1H, Ar-CH₂-Ar), 3.16 (d, $J = 13.0$ Hz, 1H, Ar-CH₂-Ar), 3.13 (d, $J = 13.0$ Hz, 2H, Ar-CH₂-Ar), 2.01–1.88 (m, 4H, -O-CH₂-CH₂-), 1.83 (sextet, $J = 7.0$ Hz, 2H, -O-CH₂-CH₂-), 1.82 (sextet, $J = 7.0$ Hz, 2H, -O-CH₂-CH₂-), 1.07 (t, $J = 7.4$ Hz, 3H, -O-(CH₂)₂-CH₃), 1.06 (t, $J = 7.4$ Hz, 3H, -O-(CH₂)₂-CH₃), 0.90 (t, $J = 7.4$ Hz, 3H, -O-(CH₂)₂-CH₃), 0.89 (t, $J = 7.4$ Hz, 3H, -O-(CH₂)₂-CH₃). ¹³C{¹H} NMR (125 MHz, DMSO-*d*₆): δ 157.7, 155.0, 154.9, 152.8, 152.7, 152.0, 146.71, 146.66, 140.8, 136.3, 136.2, 134.7, 133.0, 132.8, 132.74, 132.69 (2 \times), 132.5, 130.2, 127.8, 127.3, 127.2, 127.1, 126.9, 125.2 (2 \times), 121.7, 119.63, 119.55, 118.3, 117.3, 117.2, 76.7, 76.5, 76.1, 76.0, 30.4, 30.3, 29.9, 24.1, 23.1, 23.0, 22.6, 22.5, 10.7, 10.6, 9.84, 9.79. HRMS-ESI⁺: (C₅₄H₅₈N₆O₁₀) m/z calcd for 973.4106 [M + Na]⁺, found 973.4110 [M + Na]⁺. IR (KBr) 2962, 2935, 2874, 1658, 1596, 1552, 1508, 1461, 1328, 1298, 1211, 1178, 1110, 1002, 846, 748, 687 cm⁻¹.

Titration experiments

Dilution experiments preceded titrations to confirm or exclude the self-aggregation processes in the solution. Depending on the solubility of the corresponding urea-based receptors, the studies were performed in DMSO-*d*₆ in the concentration range from 26 mM to 0.6 mM. The appropriate data are shown in the ESI.†

All titrations were performed at a constant receptor concentration, specific for individual measurements. Anions were gradually added to the solution of receptors in the form of their TBA salt. The ¹H NMR titrations were performed in DMSO-*d*₆ using a Bruker Avance 400 spectrometer (Bruker Biospin, Rheinstetten, Germany). The UV/vis titrations were performed in DMSO (HPLC grade 99.9%, Merck) using a double beam UV-1800 spectrophotometer (Shimadzu). All UV/vis spectra were recorded in the wavelength region from 270 to 800 nm, with steps of 1 nm, in cuvettes with pathlengths of 2 or 1 mm (depending on the absorbance of receptors). The association constants of the resulting complexes were evaluated using the freeware program Bindfit¹⁶ using the most pronounced shifts in spectra (¹H NMR) or the whole parts of the absorption curves, where the changes in absorbance were the most significant (UV/vis).

The complexation ability of receptors was expressed by association constant K and/or overall association constant β . In the cases of **7a**, **8a** and **9**, the binding ability was evaluated using the overall association constant β by testing four different models for data fitting. The right choice of the model (non-cooperative model) was justified by the accuracy of the fit of experimental points and the fulfilment of the additional fitting conditions, as reported by Thordarson.¹⁸

Electrochemistry

A combination of steady-state (RDE) and dynamic methods (CV) was applied to study the electrochemical properties of some prepared molecules. The measurements were performed in DMSO solution (for DNA and peptide synthesis, containing max 0.025% H₂O, Merck) using 0.1 M TBAPF₆ (from TCI > 98%) as the supporting electrolyte, with the concentration of the corresponding urea derivative as specified in the ESI.† Due to low conductivity, the three-electrode systems were applied in all cases. As the reference electrode, a saturated calomel electrode (SCE) separated from the investigated sample by a bridge filled by the blank (DMSO-electrolyte solution) was applied. As the auxiliary electrode, Pt sheet was chosen. The material of the working electrode was dependent on the particular experiment. For linear sweep voltammetry on RDE (glassy carbon – diameter 1 mm) at a scan rate of 10 mV s⁻¹, several rotation rates (100, 250, 500 and 1000 s⁻¹) were used. For cyclic voltammetry (scan rates 100, 200, and 500 mV s⁻¹), HMDE, glassy carbon electrode (diameter 1 mm) or Pt disk electrode (diameter 1 mm) were used. All these experiments were carried out in an undivided 20 mL cell (using 10 mL of the appropriate solution), and before analyses, the solutions were deaerated using argon (99.998%, Messer). The experiments were carried out using the computer-driven digital potentiostat PGSTAT101 (Autolab-Metrohm) controlled by the software NOVA 2.1.3.

X-ray measurements

Crystallographic data for 3a. $M = 836.04$ g mol⁻¹, triclinic system, space group $P\bar{1}$, $a = 9.3111(4)$ Å, $b = 15.9194(7)$ Å, $c = 16.2178(7)$ Å, $\alpha = 98.430(2)^\circ$, $\beta = 106.068(2)^\circ$, $\gamma = 95.002(2)^\circ$, $Z = 2$,



$V = 2264.31(17) \text{ \AA}^3$, $D_c = 1.226 \text{ g cm}^{-3}$, $\mu(\text{Cu-K}\alpha) = 0.68 \text{ mm}^{-1}$, crystal dimensions of $0.202 \text{ mm} \times 0.209 \text{ mm} \times 0.295 \text{ mm}$. Data were collected at $180(2) \text{ K}$ on a D8 Venture diffractometer equipped with a Photon II detector and an Incoatec microfocus sealed tube with Cu-K α radiation. The structure was solved by direct methods²³ and anisotropically refined by full matrix least squares on F^2 using the CRYSTALS suite of programs²⁴ to a final value of $R = 0.0552$ and $wR = 0.1465$, using 8282 independent reflections ($\theta_{\text{max}} = 68.550^\circ$), 560 parameters and 0 restraints. The hydrogen atoms bonded to carbon atoms were placed in calculated positions, and all hydrogen atoms were refined with riding constraints. The disordered solvent positions were found in difference electron density maps and refined with no restrictions. MCE²⁵ was used for visualization of electron density maps. The occupancies of disordered functional groups were constrained to full. The structure was deposited in the Cambridge Structural Database under number CCDC 2381330.†

Data availability

The data supporting this article have been included as part of the ESI.†

Crystallographic data for compound **3a** have been deposited at the CCDC under CCDC 2381330 and can be obtained from <https://www.ccdc.cam.ac.uk>.†

Conflicts of interest

There are no conflicts of interest to declare.

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