

Substituent Effects on the Inclusion of 1-Alkyl-6-alkoxy-quinolinium in 4-Sulfonatocalix[*n*]arenes

Véronique Wintgens,[†] Cédric Lorthioir,[‡] Zsombor Miskolczy,[§] Catherine Amiel,[†] and László Biczók^{*,§}

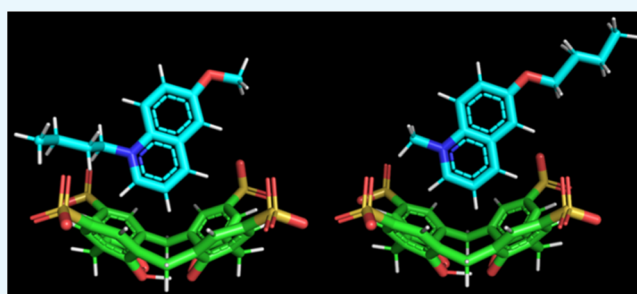
[†]Université Paris Est, ICMPE (UMR 7182), CNRS, UPEC, 2 rue Henri Dunant, F 94320 Thiais, France

[‡]Sorbonne Université, CNRS, Collège de France, Laboratoire de Chimie de la Matière Condensée de Paris (LCMCP), 4 Place Jussieu, 75005 Paris, France

[§]Institute of Materials and Environmental Chemistry, Research Centre for Natural Sciences, Hungarian Academy of Sciences, P.O. Box 286, 1519 Budapest, Hungary

Supporting Information

ABSTRACT: The effect of the chain length of the alkyl and alkoxy substituents on the binding characteristics of 1-alkyl-6-alkoxy-quinolinium cations was studied using 4-sulfonatocalix[4]arene (SCX4) and 4-sulfonatocalix[6]arene (SCX6) in neutral aqueous solutions at 298 K. Isothermal calorimetric titrations showed enthalpy-controlled inclusion with 1:1 stoichiometry. The equilibrium constants of complexation were always larger for the confinement in SCX4 than in its SCX6 homologue because the better matching between the host and guest sizes allowed more exothermic interaction. The binding affinity diminished with the lengthening of the aliphatic chain of the guests in the case of the association with SCX4, but insignificant change was found for SCX6 complexes. The most substantial change in the enthalpic and entropic contributions to the driving force of complex production occurred when the alkyl chain was linked to the heterocyclic nitrogen and the number of its carbon atoms varied between 1 and 4. ¹H NMR spectra evidenced that in SCX6, the 1-alkyl-6-alkoxy-quinolinium cations could be included within the macrocycle cavity. In the case of SCX4, the quinolinium ring is always inside the host, but the alkyl chain is included within SCX4 only for a short chain length (*n* up to 4). In contrast, the alkoxy chain displays a very weak interaction with the cavity irrespective of the length. Because of the outward orientation from the host, the lengthening of the alkoxy substituent of the quinolinium moiety barely influenced the thermodynamics of inclusion in SCX4. Distinct linear enthalpy–entropy correlations were found for the encapsulation in SCX4 and SCX6.



1. INTRODUCTION

The biocompatible, highly water-soluble 4-sulfonatocalix[*n*]arene (SCX_{*n*}) macrocycles represent a particularly important and intensely studied family of cavitands because of their capability to produce various supramolecular architectures,^{1–3} fluorescent sensing systems,^{4–6} and host–guest complexes.^{7,8} Their π -electron-rich, flexible cavity confines various biologically important compounds,^{9–11} drugs,^{12–14} and photochromic substances,¹⁵ thereby enhancing the solubility and stability of the guests. The binding between surfactants and SCX_{*n*} can induce association into nanoparticles,¹⁶ vesicles,^{17,18} or supramolecular micelles.^{19,20} Such self-assemblies have great potential in the design of stimuli-responsive aggregates and drug delivery vehicles.^{21,22}

6-Methoxyquinolinium derivatives are important fluorophores because they can be used in aqueous solution in a wide pH range and emit substantially Stokes-shifted strong fluorescence. Members of this class of compounds were applied as probes for the detection of the intracellular Cl[–] levels^{23–25} and the monitoring of Cl[–] transport across cell membranes.^{26,27} The photophysical behavior and inclusion

complex formation of 6-methoxy-1-methylquinolinium (C₁C₁OQ⁺) were investigated,²⁸ and the substantial change of its fluorescence was used to detect the embedment of a herbicide in 4-sulfonatocalix[4]arene (SCX4).²⁹

For the rational design of tailor-made self-assembled systems incorporating SCX_{*n*} hosts, the deeper understanding of the relationship between the molecular structure of the components and the driving force of encapsulation is of crucial importance. In the present study, we reveal how the systematic variation of the length and the location of the alkyl chain in quinolinium derivatives influence the thermodynamics of binding and the structure of the produced inclusion complex. We focus on two types of guests, 1-methyl-6-alkoxy-quinolinium (C₁C_{*n*}OQ⁺, *n* = 1, 2, 4, 6) and 1-alkyl-6-methoxy-quinolinium (C_{*n*}C₁OQ⁺, *n* = 1, 2, 4, 6, 8) cations, and the number of sulfonatophenol units in the SCX_{*n*} cavitands is varied from *n* = 4 to 6. We intend to deal only

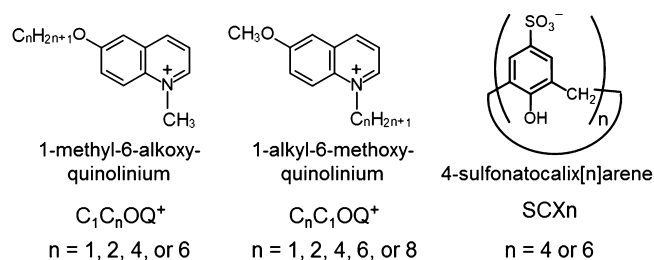
Received: April 17, 2018

Accepted: July 9, 2018

Published: August 2, 2018

with 1:1 inclusion complexes. Therefore, the larger SCX8 homologue, producing associates of various stoichiometries, were not encompassed in the present work. On the basis of the reported deprotonation constants,³⁰ we infer that one phenolic OH of SCX4 dissociated, whereas two phenolic OH substituents lost protons in 4-sulfonatocalix[6]arene (SCX6) at neutral pH. Therefore, SCX4 and SCX6 had 5 and 8 negative charges under our experimental conditions, respectively. The formulas of the employed compounds are presented in Scheme 1.

Scheme 1. Chemical Formulas of the Guest and Host Compounds



2. RESULTS AND DISCUSSION

2.1. Thermodynamics of Inclusion in SCXn. To get insight into the driving force of the confinement of 1-alkyl-6-alkoxyquinolinium ($C_nC_mOQ^+$) in SCX4 or SCX6 cavitands, isothermal titration calorimetry (ITC) measurements were performed. Figure 1 shows representative results correspond-

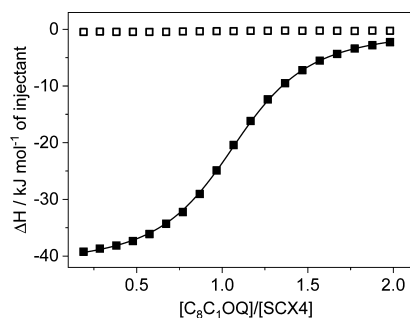


Figure 1. Variation of the integrated heat evolved per injection during the titration of 0.1 mM SCX4 solution by 2 mM $C_8C_1OQ^+$ solution (■). The line represents the fit with a 1:1 complexation model, whereas (□) displays the heat released per injection of the titrant into water.

ing to the titration of SCX4 by $C_8C_1OQ^+$ and the determination of the dilution enthalpy of the titrant. The released heat decreased upon successive additions of the ligand until the value of the dilution heat was reached because of the gradual reduction of the amount of the free binding sites. The inflexion point appeared at the equimolar component concentration, indicating the 1:1 stoichiometry of the resulting complex. The nonlinear least-squares fit of the enthalpograms provided the binding constant (K) and the enthalpy change upon encapsulation (ΔH). The standard free enthalpy (ΔG) and entropy changes (ΔS) were deduced on the basis of the following equation

$$\Delta G = -RT \ln K = \Delta H - T\Delta S \quad (1)$$

where R is the gas constant and T stands for the temperature.

Similar experiments were carried out with various $C_nC_mOQ^+$ -SCXn pairs, and the results are summarized in Table 1. All quinolinium derivatives display larger binding affinity to SCX4 than to SCX6 because the tighter fit into the smaller and more rigid homologue results in more exothermic (6–7 kJ mol⁻¹ difference) host–guest interactions. The better match of the size of the guests with the cavity of SCX4 ensures the more efficient limitation of the degrees of freedom of the components resulting in more substantial entropy diminution (2–4 kJ mol⁻¹ $T\Delta S$ difference) for the encapsulation in SCX4.

Figure 2 depicts the correlations between the binding constants and the number of carbon atoms in the aliphatic

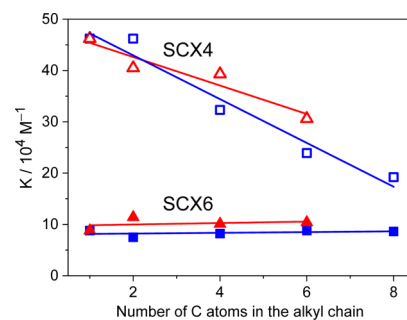


Figure 2. Equilibrium constants for the complexation of $C_nC_1OQ^+$ (squares) or $C_1C_nOQ^+$ (triangles) with SCX4 (empty symbols) and SCX6 (filled symbols) as a function of the number of carbon atoms in the aliphatic chain of the guests at pH 7 and 298 K.

chain of the guest molecules. Decreasing trends were observed when SCX4 served as a host, and the extent of this variation was more pronounced for the $C_nC_1OQ^+$ class of compounds

Table 1. Binding Constants and Thermodynamic Parameters for Inclusion Complex Formation in Neutral Aqueous Solution at 298 K

Guest	SCX4				SCX6			
	$K, 10^4 M^{-1}$	$\Delta G, kJ mol^{-1}$	$\Delta H, kJ mol^{-1}$	$T\Delta S, kJ mol^{-1}$	$K, 10^4 M^{-1}$	$\Delta G, kJ mol^{-1}$	$\Delta H, kJ mol^{-1}$	$T\Delta S, kJ mol^{-1}$
$C_1C_1OQ^+I^-$	46.2 ± 1.8	-32.4	-38.7 ± 0.4	-6.3	8.8 ± 0.4	-28.3	-32.1 ± 0.3	-3.8
$C_2C_1OQ^+Br^-$	46.2 ± 1.8	-32.4	-40.7 ± 0.4	-8.3	7.5 ± 0.3	-27.8	-33.4 ± 0.3	-5.6
$C_4C_1OQ^+I^-$	32.3 ± 1.3	-31.5	-42.6 ± 0.4	-11.1	8.2 ± 0.4	-28.1	-35.6 ± 0.4	-7.5
$C_6C_1OQ^+Br^-$	23.9 ± 1.0	-30.7	-41.6 ± 0.4	-10.9	8.8 ± 0.4	-28.3	-35.8 ± 0.4	-7.5
$C_8C_1OQ^+Br^-$	19.2 ± 0.8	-30.4	-41.7 ± 0.4	-11.3	8.6 ± 0.3	-28.2	-35.6 ± 0.4	-7.4
$C_1C_1OQ^+I^-$	46.2 ± 1.8	-32.4	-38.7 ± 0.4	-6.3	8.8 ± 0.4	-28.3	-32.1 ± 0.3	-3.8
$C_1C_2OQ^+I^-$	40.5 ± 1.6	-32.0	-37.8 ± 0.4	-5.8	11.4 ± 0.5	-28.9	-33.3 ± 0.3	-4.5
$C_1C_4OQ^+I^-$	39.3 ± 1.6	-32.0	-37.8 ± 0.4	-5.8	10.1 ± 0.4	-28.6	-30.1 ± 0.3	-1.5
$C_1C_6OQ^+I^-$	30.6 ± 1.2	-31.3	-37.8 ± 0.4	-6.5	10.4 ± 0.4	-28.6	-28.8 ± 0.3	-0.2

than for the $C_1C_nOQ^+$ family. In contrast, the K values of SCX6 complexes were practically independent of the length of the alkyl group because the entropy contribution counterbalanced the ΔH variation in the series of compounds (Table 1). The binding constants of the $C_nC_mOQ^+$ embedment in SCX4 were significantly larger than the corresponding values reported for 1-methyl-3-alkylimidazolium³¹ ($C_n\text{mim}^+$) and alkylammonium³² ($C_n\text{NH}_3^+$) guests. For these guests, the increase of the hydrophobicity and aromatic character follows this order: $C_n\text{NH}_3^+ < C_n\text{mim}^+ < C_nC_mOQ^+$. In particular, the quinolinium derivatives participate in the strongest hydrophobic host–guest interactions and additionally, are able to interact via π – π binding between the aromatic ring of the guest and the π -electron-rich core of SCX4. These two interactions will both contribute to the ΔH diminution. Indeed, the removal of a weaker-bound hydration shell from the cationic moiety in the course of host–guest complexation requires less desolvation energy for $C_nC_mOQ^+$. The more substantial binding affinity of $C_nC_mOQ^+$ cations to SCX4 compared with that of $C_n\text{mim}^+$ and $C_n\text{NH}_3^+$ arises from the larger enthalpy gain. For example, the ΔH values of SCX4 complex formation at 298 K are -17.9 , -33.9 , and -37.9 kJ mol^{-1} for $C_4\text{NH}_3^+$, $C_4\text{mim}^+$, and $C_1C_4OQ^+$, respectively, whereas the ΔS values of these processes are 16.6, -19.0 and -19.5 $\text{J mol}^{-1} \text{K}^{-1}$, respectively.^{31,32} In the case of the confinement in SCX6, $C_nC_mOQ^+$ cations had about 1 order of magnitude larger binding constant than $C_n\text{mim}^+$ guests³¹ because of the less negative ΔS for the encapsulation of the former class of compounds.

The enthalpy-driven exothermic formation of all $C_nC_mOQ^+$ complexes were accompanied by unfavorable entropy change, implying that the release of water from the hydrate shell of the constituents upon association does not counterbalance the loss of entropy due to the inclusion of the guest in the host cavity. The variation of the enthalpy and entropy terms with the alkyl substituent significantly differed for $C_nC_1OQ^+$ –SCX4 and $C_1C_nOQ^+$ –SCX4 (Figure S1). In the case of the former complexes, the gradual lengthening of the carbon chain from methyl to butyl led to ΔH and $T\Delta S$ reduction, but the change levelled off upon further extension of the aliphatic group. In contrast, ΔH and $T\Delta S$ were practically independent of the molecular structure of $C_1C_nOQ^+$ –SCX4, but both quantities increased with the alkyl length for the encapsulation of $C_1C_nOQ^+$ in SCX6. The thermodynamic parameters of $C_nC_1OQ^+$ embedment in SCX4 followed a similar trend to that found for the inclusion in SCX6. The latter cavitant may adopt inverted double partial cone conformation.³³ The manner of $C_nC_1OQ^+$ inclusion in the partial cone of SCX6 probably resembles the location in SCX4.

We found two distinct linear correlations when the entropy variation of the SCX4 and SCX6 complexes of quinolinium derivatives ($T\Delta S$) was plotted as a function of the binding enthalpy (ΔH) (Figure 3). The experimental data were fitted by the following relationship³⁴

$$T\Delta S = \alpha\Delta H + T\Delta S_0 \quad (2)$$

where the slope (α) indicates the extent to which the enthalpy change is counterbalanced by the concomitant entropy loss upon complex formation.

Enthalpy–entropy compensation has been observed for host–guest binding^{35–38} because the exothermicity of association usually restricts the movement of the constituents, thereby causing growing entropy loss. In the case of the

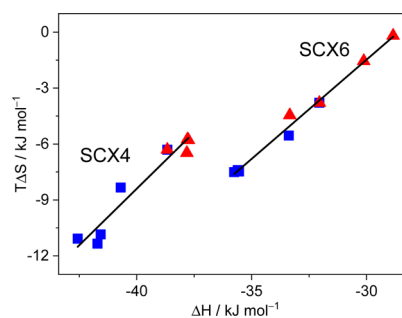


Figure 3. Enthalpy–entropy correlation for the 1:1 inclusion of $C_nC_1OQ^+$ (blue square) or $C_1C_nOQ^+$ (red triangle) in SCX4 or SCX6 in neutral aqueous solution at 298 K.

confinement in SCX4, $\alpha = 1.21$ and $T\Delta S_0 = 40.0$ kJ mol^{-1} were calculated, whereas $\alpha = 1.06$ and $T\Delta S_0 = 30.4$ kJ mol^{-1} were derived from the thermodynamic data of SCX6 complexes. The larger slope (α) and intercept ($T\Delta S_0$) of the linear ΔH versus $T\Delta S$ relationship for the confinement in the smaller SCX4 cavitant is due to the tighter binding and the release of more water molecules from the solvation shell of the reactants than in the case of the association with SCX6. The parameters found for $C_nC_mOQ^+$ –SCX4 complexes in the present study resemble those reported for the embedment of $C_n\text{mim}^+$ cations in SCX4 ($\alpha = 1.38$ and $T\Delta S_0 = 40.3$ kJ mol^{-1}).³¹ In the case of SCX6 host, α and $T\Delta S_0$ for the inclusion of $C_nC_mOQ^+$ guests were larger than those found for $C_n\text{mim}^+$ ($\alpha = 0.80$ and $T\Delta S_0 = 16.6$ kJ mol^{-1}),³¹ and approached the values previously found for 1:1 $C_n\text{mim}^+$ –SCX8 associates ($\alpha = 0.96$ and $T\Delta S_0 = 27.9$ kJ mol^{-1}).³⁹ The much larger $T\Delta S_0$ of $C_nC_mOQ^+$ –SCX6 complexes relative to that of $C_n\text{mim}^+$ –SCX6 implies that more water molecules are expelled from the host interior upon the entry of the bulkier, more hydrophobic quinolinium headgroup. The looser binding of the smaller methylimidazolium moiety causes less extensive desolvation and as a consequence, smaller entropy gain contribution. The close to unity α of $C_nC_mOQ^+$ –SCX6 complexes indicates that the change of ΔH , arising from the variation of the substitution pattern of the guest, is almost completely offset by the accompanying entropy modification. When the smaller, more hydrophilic headgroup of $C_n\text{mim}^+$ is encapsulated in SCX6, fewer degrees of freedom are limited and less water is released from the hydrate shell of the constituents into the bulk solution. As a consequence, ΔS changes in a smaller extent in the series of $C_n\text{mim}^+$ –SCX6 complexes than upon the inclusion of $C_nC_mOQ^+$ in SCX6.

2.2. Molecular Structure of the SCX n Complexes. To gain insight into the structure of the inclusion complexes, ^1H NMR experiments were performed. As a representative example, Figure 4 displays the spectra of $C_4C_1OQ^+$ –SCX4 and $C_1C_4OQ^+$ –SCX4. The broadening observed for the peaks related to the SCX4 protons results from conformational changes of the macrocycle that could correspond to inversions of the aromatic rings⁴⁰ and/or distortions of the cone conformation.⁴¹ These dynamical processes stand in the intermediate regime, as can be seen in Figure 4. In contrast, the same spectra suggest that both quinolinium derivatives undergo a fast exchange between the free and the complexed states over the characteristic NMR time scale. When the SCX4/quinolinium molar ratio is set to 2:1 and the total concentration of 10 mM is used, more than 95% of the quinolinium derivatives is complexed to SCX4 on the basis of

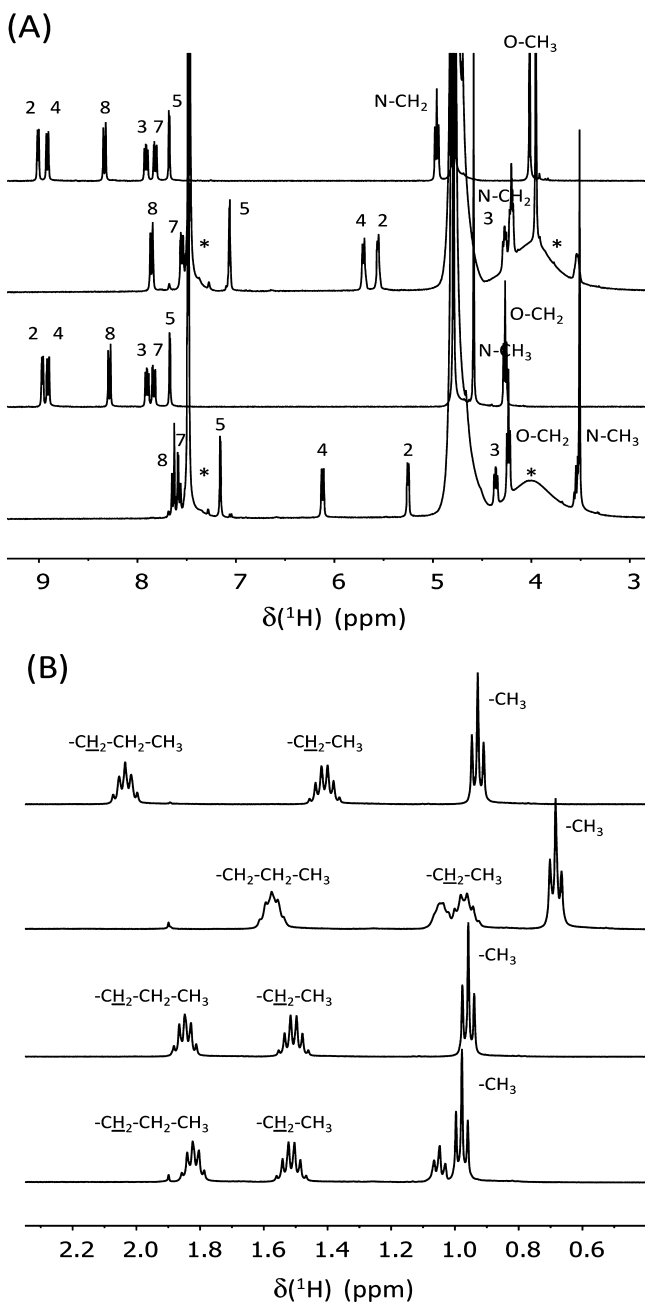


Figure 4. From top to bottom: ^1H NMR spectra of 10 mM $\text{C}_4\text{C}_1\text{OQ}^+$, 3.3 mM $\text{C}_4\text{C}_1\text{OQ}^+$ + 6.7 mM SCX4 mixture, 10 mM $\text{C}_1\text{C}_4\text{OQ}^+$, and 3.3 mM $\text{C}_1\text{C}_4\text{OQ}^+$ + 6.7 mM SCX4 mixture in D_2O at 298 K. The asterisks denote the peaks related to the SCX4 protons; (A,B) correspond to two distinct spectral ranges. The small triplet at 1.05 ppm results from an impurity present in the SCX4 compound. (The spectrum of neat SCX4 solution displays this small contribution.)

the K value (Table 1). Therefore, the ^1H chemical shifts measured under such conditions allow the probing of the quinolinium location within both $\text{C}_4\text{C}_1\text{OQ}^+-\text{SCX4}$ and $\text{C}_1\text{C}_4\text{OQ}^+-\text{SCX4}$ complexes. In the case of the former complex, the resonances related to the quinolinium and the butyl protons shift upfield in comparison to the signals of the free $\text{C}_4\text{C}_1\text{OQ}^+$ molecules. Figure 5A and Table S1 show that the most significant encapsulation-induced chemical shift variations ($\Delta\delta$) are observed for the peaks assigned to H2, H3, and H4 for all studied guests. This indicates that the

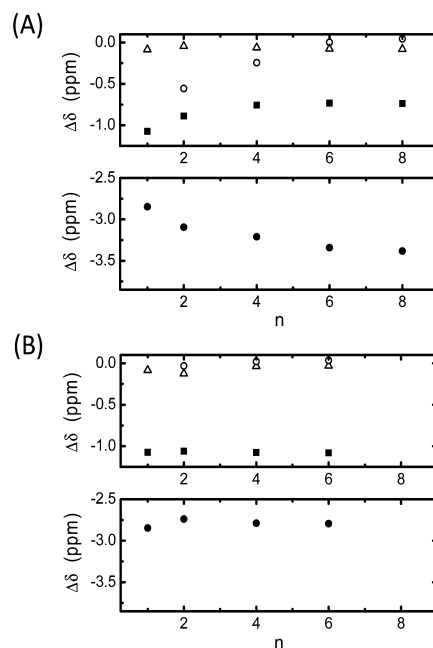


Figure 5. Alkyl chain length (n) dependence of the ^1H chemical shift variation upon complex formation for (A) $\text{C}_n\text{C}_1\text{OQ}^+-\text{SCX4}$: NCH_2 (■), H4 (●), CH_3 (○), OCH_3 (△) and (B) $\text{C}_1\text{C}_n\text{OQ}^+-\text{SCX4}$: NCH_3 (■), H4 (●), CH_3 (○), OCH_2 (△). Both complexes are obtained using 1:2 quinolinium/SCX4 molar ratio and a total concentration of 10 mM.

corresponding part of the quinolinium has the deepest immersion within the SCX4 cavity. Among the aromatic proton resonances, the H7 signal undergoes the weakest upfield shift, implying that this terminal of the quinolinium is oriented toward the aqueous phase. The $\Delta\delta$ value determined for the methoxy protons is the smallest in $\text{C}_n\text{C}_1\text{OQ}^+-\text{SCX4}$ complexes, indicating that the methoxy substituent is located far from the core of the macrocycle.

Among the alkyl protons of $\text{C}_n\text{C}_1\text{OQ}^+$, the increase of the alkyl chain length resulted in the largest chemical shift variation for the $-\text{CH}_3$ protons. As shown in Figure 5A, the $\Delta\delta$ value of this signal is reduced upon the gradual increase of the number of aliphatic carbon atoms from methyl to hexyl, while the change tends to level off for $\text{C}_n\text{C}_1\text{OQ}^+-\text{SCX4}$ with $n \geq 6$. Such a behavior suggests that the short alkyl chains ($n = 1, 2,$ and 4) were included in SCX4. Hexyl and octyl substituents are too long to be fully embedded in SCX4. Therefore, their end is located outside the cavity, exhibiting vanishing interactions with the host. This is in accord with the results of the calorimetric experiments (Table 1), which demonstrated that the exothermicity of inclusion in SCX4 initially grew upon the gradual increase of the number of aliphatic carbon atoms in the substituent linked to the heterocyclic nitrogen and then levelled off. The incorporation of the short alkyl chains ($n < 6$) within the macrocycle may rationalize the line broadening observed for their ^1H NMR peaks in the spectrum of $\text{C}_n\text{C}_1\text{OQ}^+-\text{SCX4}$, as exemplified for $n = 4$ in Figure 4B. The confinement of the aliphatic chains should indeed induce a restriction in their reorientational motions and then, a reduction of their $T_2(^1\text{H})$ relaxation times compared to free $\text{C}_n\text{C}_1\text{OQ}^+$.

As the number of carbon atoms in the alkyl chain is raised from $n = 1$ to 4, the $\Delta\delta$ values for the aromatic protons vary. In particular, a continuous increase is observed for H2 and H8

(and also NCH_2), whereas the opposite trend is observed for H3 and H4. These simultaneous variations suggest an alteration of the orientation of the quinolinium ring within the SCX4 cavity resulting from the increase of the volume occupied by the alkyl chain.

In the case of the $\text{C}_1\text{C}_n\text{OQ}^+-\text{SCX4}$ complexes, the chemical shift of the ^1H NMR peaks related to the alkoxy moiety is nearly unchanged with respect to the value observed for the solution of the corresponding uncomplexed quinolinium (Figure 5B and Table S2). Figure 4 shows that the chemical shift of each proton of the quinolinium ring of $\text{C}_1\text{C}_n\text{OQ}^+$ still displays a significant change upon the addition of SCX4. In particular, the peaks attributed to H2, H3, H4, and $\text{N}-\text{CH}_3$ protons undergo the most significant upfield shift, implying that the quinolinium ring of $\text{C}_1\text{C}_n\text{OQ}^+$ is inside the SCX4 cavity, whereas the alkoxy chain protrudes from it. The line width of the ^1H NMR peaks assigned to the alkoxy group of $\text{C}_1\text{C}_n\text{OQ}^+-\text{SCX4}$ is similar to the one observed in the spectrum of free $\text{C}_1\text{C}_n\text{OQ}^+$ (Figure 4B). In contrast to $\text{C}_n\text{C}_1\text{OQ}^+-\text{SCX4}$ ($n < 6$), the aliphatic chain is not confined within the macrocycle and as a result, no significant line broadening is found.

The $\Delta\delta$ values for the protons of the quinolinium moiety of $\text{C}_1\text{C}_n\text{OQ}^+$ and $\text{C}_n\text{C}_1\text{OQ}^+$ differ for $n \geq 2$ (Figure 5), suggesting distinct orientations of the heterocyclic ring within SCX4. Another main difference between the $\text{C}_1\text{C}_n\text{OQ}^+-\text{SCX4}$ and the $\text{C}_n\text{C}_1\text{OQ}^+-\text{SCX4}$ complexes concerns the position of the alkoxy and alkyl chains. In the case of the $\text{C}_n\text{C}_1\text{OQ}^+-\text{SCX4}$ complexes, the alkyl chains may be included within the SCX4 host for low n values ($n < 6$). As n is raised further, the alkyl chain end extends out of the cavity. This could result from a contribution of steric hindrance. In contrast, for the $\text{C}_1\text{C}_n\text{OQ}^+-\text{SCX4}$ complexes, the shortest alkoxy group ($n = 1$) is already far from the interior of the host. Therefore, the increase of this chain length does not cause any $\Delta\delta$ change (Figure 5B) and the orientation of the quinolinium ring of $\text{C}_1\text{C}_n\text{OQ}^+-\text{SCX4}$ remains unaltered. The outward orientation from the host cavity rationalizes that the lengthening of the alkoxy substituent brings about minor change in ΔH and ΔS of $\text{C}_1\text{C}_n\text{OQ}^+-\text{SCX4}$ complex formation (Table 1).

Because of the lower binding affinity to SCX6, ^1H NMR spectra were recorded for a 1:3 quinolinium/SCX6 molar ratio to ensure that more than 95% of the guest was confined to the host. The variations of the proton chemical shifts ($\Delta\delta$) upon complexation of $\text{C}_1\text{C}_1\text{OQ}^+$ with SCX6 and SCX4 are compared in Table S3 of the Supporting Information. The upfield shift of all ^1H NMR peaks upon confinement in SCX6 indicates that the entire $\text{C}_1\text{C}_1\text{OQ}^+$ cation is included in the spacious cavity of the host. In the case of the $\text{C}_1\text{C}_1\text{OQ}^+-\text{SCX6}$ complex, the upfield shift related to the H2, H3, and H4 signals were significantly smaller because of the looser binding. For the other aromatic protons of the guest, the upfield shift was found to be higher than that of the corresponding peaks in the spectrum of $\text{C}_1\text{C}_1\text{OQ}^+-\text{SCX4}$. In the larger SCX6 macrocycle, various orientations of the guest could be allowed, leading to an additional averaging of the corresponding ^1H chemical shifts.

When a methyl substituent of $\text{C}_1\text{C}_1\text{OQ}^+$ was replaced by a butyl group, the chemical shift changes induced by the incorporation into SCX6 remained similar to those of $\text{C}_1\text{C}_1\text{OQ}^+-\text{SCX6}$ and did not depend significantly on the location of the butyl moiety (Table S4, Supporting Information). All butyl proton signals of $\text{C}_4\text{C}_1\text{OQ}^+$ and

$\text{C}_1\text{C}_4\text{OQ}^+$ shifted upfield, indicating that the whole butyl chain interacts with SCX6. The larger and more flexible SCX6 macrocycle adapts better to the geometrical features of the guests than SCX4. The guest binding induced calixarene conformation change, the so-called “template effect” which has also been noticed for other associates,^{40,42,43} may explain the slightly larger alteration of ΔH and ΔS with the substituent in the series of $\text{C}_1\text{C}_n\text{OQ}^+-\text{SCX6}$ complexes (Table 1).

3. CONCLUSIONS

The confinement of $\text{C}_n\text{C}_m\text{OQ}^+$ in $\text{SCX}n$ cavities significantly depends not only on the location and the length of the hydrocarbon chain of the guest but also on the size of the macrocycle. Both the thermodynamic parameters of the embedment in $\text{SCX}n$ and the structure of the resulting $\text{C}_n\text{C}_m\text{OQ}^+-\text{SCX}n$ complexes can be tuned by the variation of the number of aliphatic carbon atoms attached to the nitrogen of the quinolinium moiety, but alteration of the alkoxy chain length causes a minor effect. ^1H NMR experiments demonstrated that in the case of SCX4, the short alkyl chains ($n \leq 4$) may be incorporated, together with the quinolinium ring, within the host cavity, while a part of longer alkyl chains are located out of the cavity. In contrast, even the short alkoxy groups were found to display very weak interactions with SCX4, while the quinolinium ring is still found within the cavity. The larger SCX6 macrocycle allows an easier inclusion of both kinds of quinolinium ($\text{C}_n\text{C}_1\text{OQ}^+$ and $\text{C}_1\text{C}_m\text{OQ}^+$), with an incorporation of either the alkyl or the alkoxy chains within the cavity when n or $m \leq 4$. From a thermodynamic point of view, the tighter binding in the smaller, more rigid SCX4 brings about larger enthalpy gain accompanied by more substantial entropy diminution, which lead to considerably larger equilibrium constant than the association with SCX6. The stability of the $\text{C}_n\text{C}_m\text{OQ}^+-\text{SCX6}$ complexes cannot be modified by the alteration of the alkyl substituent of the guest because the enthalpy variation is compensated by the concomitant entropy change. The knowledge on the relationship between the molecular structure of constituents and the thermodynamics of host–guest binding gained in the present study may be applied in the design of tailor-made supramolecular systems and self-assembled architectures.

4. EXPERIMENTAL SECTION

4.1. Materials. SCX4 and SCX6 were purchased from Acros Organics and used after drying under vacuum at 343 K overnight. Double-distilled water was used as a solvent, and $\text{SCX}n$ ($n = 4$ or 6) solutions were neutralized by the minimum volume of concentrated NaOH. The various 1-alkyl-6-alkoxy-quinolinium salts were synthesized and characterized as described in the Supporting Information. $\text{C}_1\text{C}_n\text{OQ}^+$ and $\text{C}_4\text{C}_1\text{OQ}^+$ had I^- counterions, whereas $\text{C}_2\text{C}_1\text{OQ}^+$, $\text{C}_6\text{C}_1\text{OQ}^+$, and $\text{C}_8\text{C}_1\text{OQ}^+$ were used as Br^- salts.

4.2. Sample Preparation. Stock solutions of 1-alkyl-6-alkoxy-quinolinium salts (2 mM) and $\text{SCX}n$ (0.1 mM, adjusted to pH 7 by NaOH addition) were prepared for ITC experiments. NMR experiments were performed by mixing appropriate amounts of 1-alkyl-6-alkoxy-quinolinium (6 or 10 mM) and $\text{SCX}n$ (6 or 10 mM, neutralized by NaOD) solutions in D_2O . All the experiments were carried out at pH 7.

4.3. Instrumentation. ^1H NMR spectra were recorded in deuterated dimethyl sulfoxide or in D_2O on a Bruker Avance II 400 MHz NMR spectrometer. ITC measurements were carried

out with a MicroCal VP-ITC microcalorimeter. Quinolinium solutions were injected from the computer-controlled microsyringe at an interval of 180 s into the cell (volume = 1.4569 mL) containing 0.1 mM SCX n solution at pH 7, while stirring at 450 rpm.

■ ASSOCIATED CONTENT

📄 Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/acsomega.8b00736.

Detailed descriptions of synthesis and ^1H NMR spectral characteristics of the studied quinolinium derivatives, molecular structure dependence of the thermodynamic parameters of host–guest complexes, and variation of the ^1H NMR chemical shifts upon inclusion complex formation (PDF)

■ AUTHOR INFORMATION

Corresponding Author

*E-mail: biczok.laszlo@ttk.mta.hu (L.B.).

ORCID

László Biczók: 0000-0003-2568-5942

Notes

The authors declare no competing financial interest.

■ ACKNOWLEDGMENTS

This work was supported by the BIONANO GINOP-2.3.2-15-2016-00017 project (to Z.M. and L.B.), the National Research, Development and Innovation Office (NKFIH, grant K123995 to Z.M. and L.B.), and the János Bolyai Research Scholarship Program of the Hungarian Academy of Sciences (to Z.M.).

■ REFERENCES

- (1) Atwood, J. L.; Barbour, L. J.; Hardie, M. J.; Raston, C. L. Metal sulfonatocalix[4,5]arene complexes: Bi-layers, capsules, spheres, tubular arrays and beyond. *Coord. Chem. Rev.* **2001**, *222*, 3–32.
- (2) Liu, Y.; Wang, Y.-X. Selective molecular binding and nano-supramolecular assembly of *p*-sulfonatocalix[n]arenes. *Non-Covalent Interactions in the Synthesis and Design of New Compounds*; John Wiley & Sons, Inc., 2016; pp 261–281.
- (3) Ling, I.; Alias, Y.; Raston, C. L. Structural diversity of multi-component self-assembled systems incorporating *p*-sulfonatocalix[4]-arene. *New J. Chem.* **2010**, *34*, 1802–1811.
- (4) Guo, D.-S.; Uzunova, V. D.; Su, X.; Liu, Y.; Nau, W. M. Operational calixarene-based fluorescent sensing systems for choline and acetylcholine and their application to enzymatic reactions. *Chem. Sci.* **2011**, *2*, 1722–1734.
- (5) Ghale, G.; Nau, W. M. Dynamically analyte-responsive macrocyclic host-fluorophore systems. *Acc. Chem. Res.* **2014**, *47*, 2150–2159.
- (6) Florea, M.; Kudithipudi, S.; Rei, A.; González-Álvarez, M. J.; Jeltsch, A.; Nau, W. M. A fluorescence-based supramolecular tandem assay for monitoring lysine methyltransferase activity in homogeneous solution. *Chem.—Eur. J.* **2012**, *18*, 3521–3528.
- (7) Guo, D.-S.; Wang, K.; Liu, Y. Selective binding behaviors of *p*-sulfonatocalixarenes in aqueous solution. *J. Inclusion Phenom. Macrocyclic Chem.* **2008**, *62*, 1–21.
- (8) Dsouza, R. N.; Pischel, U.; Nau, W. M. Fluorescent dyes and their supramolecular host/guest complexes with macrocycles in aqueous solution. *Chem. Rev.* **2011**, *111*, 7941–7980.
- (9) Perret, F.; Lazar, A. N.; Coleman, A. W. Biochemistry of the *para*-sulfonato-calix[n]arenes. *Chem. Commun.* **2006**, 2425–2438.

(10) Perret, F.; Coleman, A. W. Biochemistry of anionic calix[n]arenes. *Chem. Commun.* **2011**, *47*, 7303–7319.

(11) Megyesi, M.; Biczók, L. Considerable fluorescence enhancement upon supramolecular complex formation between berberine and *p*-sulfonatocalixarenes. *Chem. Phys. Lett.* **2006**, *424*, 71–76.

(12) Mokhtari, B.; Pourabdollah, K. Applications of calixarene nanobaskets in pharmacology. *J. Inclusion Phenom. Macrocyclic Chem.* **2012**, *73*, 1–15.

(13) Chao, J.; Song, K.; Wang, H.; Guo, Z.; Zhang, B.; Zhang, T. Study on the inclusion interaction of *p*-sulfonatocalix[n]arenes with norfloxacin. *Phys. Chem. Liq.* **2017**, *55*, 579–588.

(14) Li, Q.; Guo, D.-S.; Qian, H.; Liu, Y. Complexation of *p*-sulfonatocalixarenes with local anaesthetics guests: Binding structures, stabilities, and thermodynamic origins. *Eur. J. Org. Chem.* **2012**, 3962–3971.

(15) Miskolczy, Z.; Biczók, L. Photochromism of a merocyanine dye bound to sulfonatocalixarenes: Effect of pH and the size of macrocycle on the kinetics. *J. Phys. Chem. B* **2013**, *117*, 648–653.

(16) Harangozó, J. G.; Wintgens, V.; Miskolczy, Z.; Guigner, J.-M.; Amiel, C.; Biczók, L. Effect of macrocycle size on the self-assembly of methylimidazolium surfactant with sulfonatocalix[n]arenes. *Langmuir* **2016**, *32*, 10651–10658.

(17) Wang, K.; Guo, D.-S.; Zhao, M.-Y.; Liu, Y. A supramolecular vesicle based on the complexation of *p*-sulfonatocalixarene with protamine and its trypsin-triggered controllable-release properties. *Chem.—Eur. J.* **2016**, *22*, 1475–1483.

(18) Francisco, V.; Basilio, N.; Garcia-Rio, L.; Leis, J. R.; Maques, E. F.; Vázquez-Vázquez, C. Novel catanionic vesicles from calixarene and single-chain surfactant. *Chem. Commun.* **2010**, *46*, 6551–6553.

(19) Basilio, N.; Spudeit, D. A.; Bastos, J.; Scorsin, L.; Fiedler, H. D.; Nome, F.; García-Río, L. Exploring the charged nature of supramolecular micelles based on *p*-sulfonatocalix[6]arene and dodecyltrimethylammonium bromide. *Phys. Chem. Chem. Phys.* **2015**, *17*, 26378–26385.

(20) Wintgens, V.; Miskolczy, Z.; Guigner, J.-M.; Amiel, C.; Harangozó, J. G.; Biczók, L. Reversible nanoparticle-micelle transformation of ionic liquid-sulfonatocalix[6]arene aggregates. *Langmuir* **2015**, *31*, 6655–6662.

(21) Guo, D.-S.; Liu, Y. Supramolecular chemistry of *p*-sulfonatocalix[n]arenes and its biological applications. *Acc. Chem. Res.* **2014**, *47*, 1925–1934.

(22) Wang, K.; Guo, D.-S.; Wang, X.; Liu, Y. Multistimuli responsive supramolecular vesicles based on the recognition of *p*-sulfonatocalixarene and its controllable release of doxorubicin. *ACS Nano* **2011**, *5*, 2880–2894.

(23) Baù, L.; Selvestrel, F.; Arduini, M.; Zamparo, I.; Lodovichi, C.; Mancin, F. A cell-penetrating ratiometric nanoprobe for intracellular chloride. *Org. Lett.* **2012**, *14*, 2984–2987.

(24) Inglefield, J. R.; Schwartz-Bloom, R. D. Fluorescence imaging of changes in intracellular chloride in living brain slices. *Methods* **1999**, *18*, 197–203.

(25) Schwartz, R. D.; Yu, X. Optical imaging of intracellular chloride in living brain slices. *J. Neurosci. Methods* **1995**, *62*, 185–192.

(26) Illsley, N. P.; Verkman, A. S. Membrane chloride transport measured using a chloride-sensitive fluorescent probe. *Biochemistry* **1987**, *26*, 1215–1219.

(27) Hennig, B.; Schultheiss, G.; Kunzelmann, K.; Diener, M. Ca^{2+} -induced Cl^- efflux at rat distal colonic epithelium. *J. Membr. Biol.* **2008**, *221*, 61–72.

(28) Miskolczy, Z.; Harangozó, J. G.; Biczók, L.; Wintgens, V.; Lorthioir, C.; Amiel, C. Effect of torsional isomerization and inclusion complex formation with cucurbit[7]uril on the fluorescence of 6-methoxy-1-methylquinolinium. *Photochem. Photobiol. Sci.* **2014**, *13*, 499–508.

(29) Harangozó, J. G.; Miskolczy, Z.; Biczók, L.; Wintgens, V.; Lorthioir, C. Effect of host-guest complex formation on the fluorescence of 6-methoxy-1-methyl-quinolinium cation with 4-sulfonatocalix[4]arene: utilization as a fluorescent probe for the

study of difenzoquat binding. *J. Inclusion Phenom. Macrocyclic Chem.* **2015**, *81*, 377–384.

(30) Suga, K.; Ohzono, T.; Negishi, M.; Deuchi, K.; Morita, Y. Effect of various cations on the acidity of *p*-sulfonatocalixarenes. *Supramol. Sci.* **1998**, *5*, 9–14.

(31) Wintgens, V.; Biczók, L.; Miskolczy, Z. Thermodynamics of host-guest complexation between *p*-sulfonatocalixarenes and 1-alkyl-3-methylimidazolium type ionic liquids. *Thermochim. Acta* **2011**, *523*, 227–231.

(32) Stödeman, M.; Dhar, N. Microcalorimetric titration of a tetra-*p*-sulfonated calix[4]arene with alkylammonium ions in aqueous solution. *J. Chem. Soc., Faraday Trans.* **1998**, *94*, 899–903.

(33) Atwood, J. L.; Clark, D. L.; Juneja, R. K.; Orr, G. W.; Robinson, K. D.; Vincent, R. L. Double partial cone conformation for Na₈{calix[6]arene sulfonate}·20.5 H₂O and its parent acid. *J. Am. Chem. Soc.* **1992**, *114*, 7558–7559.

(34) Inoue, Y.; Hakushi, T.; Liu, Y.; Tong, L.; Shen, B.; Jin, D. Thermodynamics of molecular recognition by cyclodextrins. 1. Calorimetric titration of inclusion complexation of naphthalenesulfonates with α -, β -, and γ -cyclodextrins: enthalpy-entropy compensation. *J. Am. Chem. Soc.* **1993**, *115*, 475–481.

(35) Inoue, Y.; Liu, Y.; Tong, L. H.; Shen, B. J.; Jin, D. S. Calorimetric titration of inclusion complexation with modified β -cyclodextrins. Enthalpy-entropy compensation in host-guest complexation: from ionophore to cyclodextrin and cyclophane. *J. Am. Chem. Soc.* **1993**, *115*, 10637–10644.

(36) Rekharsky, M. V.; Inoue, Y. Complexation thermodynamics of cyclodextrins. *Chem. Rev.* **1998**, *98*, 1875–1918.

(37) Masson, E.; Ling, X.; Joseph, R.; Kyeremeh-Mensah, L.; Lu, X. Cucurbituril chemistry: A tale of supramolecular success. *RSC Adv.* **2012**, *2*, 1213–1247.

(38) Houk, K. N.; Leach, A. G.; Kim, S. P.; Zhang, X. Binding Affinities of Host-Guest, Protein-Ligand, and Protein-Transition-State Complexes. *Angew. Chem., Int. Ed.* **2003**, *42*, 4872–4897.

(39) Wintgens, V.; Amiel, C.; Biczók, L.; Miskolczy, Z.; Megyesi, M. Host-guest interactions between 4-sulfonatocalix[8]arene and 1-alkyl-3-methylimidazolium type ionic liquids. *Thermochim. Acta* **2012**, *548*, 76–80.

(40) Shinkai, S.; Araki, K.; Matsuda, T.; Manabe, O. NMR determination of association constants for aqueous calixarene complexes and guest template effects on the conformational freedom. *Bull. Chem. Soc. Jpn.* **1989**, *62*, 3856–3862.

(41) Shinkai, S.; Mori, S.; Koreishi, H.; Tsubaki, T.; Manabe, O. Hexasulfonated calix[6]arene derivatives: A new class of catalysts, surfactants, and host molecules. *J. Am. Chem. Soc.* **1986**, *108*, 2409–2416.

(42) Arimura, T.; Kubota, M.; Araki, K.; Shinkai, S.; Matsuda, T. “Template effects” on calixarene conformations through host-guest-type interactions. *Tetrahedron Lett.* **1989**, *30*, 2563–2566.

(43) Shinkai, S.; Araki, K.; Kubota, M.; Arimura, T.; Matsuda, T. Ion template effects on the conformation of water-soluble calixarenes. *J. Org. Chem.* **1991**, *56*, 295–300.