INVESTIGATION OF SORPTION OF CALIX[4]ARENE AND CALIX[4]RESORCINARENE TETRAALKYL DERIVATIVES WITH THE LiChrosorb RP 18 SURFACE BY RP HPLC AND MOLECULAR MODELLING METHODS

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Tetrahydroxycalix[4]arene, tetraalkoxyca1x[4]arenes and tetraalkylcalix[4]resorcinarenes are capable to reversible adsorption on the surface of the hydrophobic support LiChrosorb RP 18 under conditions of the reversed-phase high performance liquid chromatography. The main chromatographic characteristics of tetrahydroxycalix[4]arene, a series of tetraalkoxyca1x[4]arenes and tetraalkylcalix[4]resorcinarenes have been determined. It has been found that in the conditions selected (the mobile phase is acetonitrile/water, 86/14, v/v) the retention times of the compounds studied are in the wide range from 4.67 min to 88.0 min depending on the nature of the substituents in the macrocyclic skeleton of the molecule. Thus, the retention times increase with increase of the length of the alkyl substituents at the lower rim of the calixarene macrocycle. According to the molecular modelling data the most effective sorption is explained by the inclusion of LiChrosorb RP 18 octadecyl groups into the lipophilic cavity formed by the benzene ring of the molecular cavity formed by the alkyl substituents at the lower rim of the calixarene/calixresorcinarene. Thus, the molecular cavity formed by the benzene rings remains opened for the Host-Guest complexation with the analyte molecules. This sorption of tetraalkoxycalix[4]arenes and tetraalkylcalix[4]resorcinarenes on the LiChrosorb RP 18 surface can significantly improve its resolution ability in RP HPLC analysis. The influence of replacement of tetrahydroxycalix[4]arene hydroxyl protons with the alkyl groups on the conformation of the macrocycle has been studied.
характеристики тетрагидроксикаликс[4]арена, серии тетраалкоксикаликс[4]аренов и тетраалкилкаликс[4]резорцинаренов. Установлено, что в выбранных условиях (подвижная фаза – ацетонитрил/вода в соотношении 86/14 по объему) времена удерживания исследуемых соединений находятся в широком диапазоне 4.67-88.0 мин в зависимости от природы заместителей в макроциклическом скелете молекулы. При этом время удерживания возрастает по мере увеличения длины алкильных заместителей на нижнем ободе каликсаренового макроцикла. Согласно данным молекулярного моделирования наиболее эффективная сорбция происходит за счет включения октадецильных групп насадки в липофильную полость, образованную алкильными заместителями нижнего обода макроцикла каликсарена/каликсрезорцинаrena. При этом молекулярная полость, сформированная бензольными кольцами макроцикла, остается открытым для комплексообразования с молекулами анализов. Такая сорбция тетраалкоксикаликс[4]аренов и тетраалкилкаликс[4]резорцинаренов на поверхности хроматографической насадки LiChrosorb RP 18 может существенно улучшить ее селективность по отношению к анализам в условиях обращенно-фазной высокоэффективной жидкостной хроматографии. Исследовано влияние замены гидроксильных протонов тетрагидроксикаликс[4]арена алкильными группами на конформацию макроциклического скелета молекулы.

Calixarenes – macrocyclic compounds available through cyclocondensation of para-substituted phenols with formaldehyde – are widely used as molecular platforms for design of specific receptors that are capable of highly selective recognition between fairly similar substrates: different cations, anions and neutral molecules [1, 2, 3, 4, 5, 6, 7]. The ability to recognize different substrates and form the Host-Guest supramolecular complexes with them is the base for wide application of calixarenes functionalized at the upper/lower rim by different groups in chemistry, physics, biology, nanotechnologies.

Calixarenes are widely used in chromatography [8, 9]. Different substrate-receptor interactions of calixarenes with organic molecules and biomolecules are studied by the HPLC method [9]. Such investigations are important in the design of mobile or stationary chromatographic phases based on calixarenes with improved separating properties. The analysis of polycyclic compounds, isomeric nitroanilines, sulfonamides, nucleic bases, nucleosides, active pharmaceutical ingredients of Doxepin, Chloroprotixene, Flupentixol, Promethazine, Chromazine, Perazine, Levomepromazine, Chlorpromazin, Paracetamol on the calixarene based stationary phases CALTREX® has been reported [10, 11, 12, 13, 14, 15, 16, 17].

The aim of the work was to study sorption of tetrahydroxycalix[4]arene (CA-OH), tetraalkoxycalix[4]arenes (CA-OC) and tetraalkylkalix[4]resorcinarenes (RA-C) (Scheme) on the LiChrosorb RP 18 surface in acetonitrile-water by RP HPLC and molecular modelling methods.

Experimental Part

Calix[4]arenes CA and calix[4]resorcinarenes RA were synthesized by the methods [18]. Acetonitrile was obtained from the Acros Organics company.

RP HPLC analysis (a liquid chromatograph Hitachi, Ltd., Tokyo, Japan) was performed in isocratic conditions using the chromatographic support LiChrosorb RP 18 and MeCN/H2O (86/14, v/v) as a mobile phase. The research was carried out at the temperature of 26°C. The UV detector was operated at 254 nm and the flow rate was 0.6 ml/min.

Lipophilicity of Log P of CA and RA was calculated by the RP HPLC method from the equation: Log P = 6.195 x (log k'). The coefficient 6.195 is the ratio of Log P value of p-dichlorobenzene (3.37) [19] to its log k' determined by the RP HPLC method in this work.

The initial molecular modelling of CA, RA and their complexes was carried out by the molecular mecha-
Retention times \( t_R \), capacity factors \( k' \) and lipophilicity \( \text{Log P} \) of calix[4]arenes CA and calix[4]resorcinarenes RA

| CA, RA | \( t_R \), min | \( k' \) | \( \text{Log P} \) |
|--------|--------------|--------|--------------|
| CA-OH  | 6.26         | 1.09   | 0.08         |
| CA-OC3 | 28.33        | 8.44   | 5.74         |
| CA-OC6 | 45.60        | 14.20  | 7.14         |
| CA-OC8 | 61.00        | 19.33  | 7.97         |
| RA-C1  | 4.67         | 0.56   | -1.56        |
| RA-C3  | 6.17         | 1.06   | 0.16         |
| RA-C5  | 9.11         | 2.04   | 1.92         |
| RA-C7  | 35.58        | 10.86  | 6.42         |
| RA-C13 | 76.54        | 24.51  | 8.60         |
| RA-C15 | 88.00        | 28.33  | 8.99         |

Calixarenes CA and calixresorcinarenes RA under conditions of the given analysis were registered on the chromatograms as sharp peaks. CA and RA retention times \( t_R \) (in the range from 4.67 min to 88.0 min), capacity factors \( k' \) and lipophilicity \( \text{Log P} \) are presented in Table.

The chromatograms of some calix[4]arenes and calix[4]resorcinarenes are presented in Fig. 1.

A linear character of adsorption isotherms of CA and RA reflects their reversible sorption on the LiChrosorb-RP18 support (Fig. 2, 3). Capacity factors \( k' \) and retention times \( t_R \) of CA, RA are linearly dependent on the number of carbon atoms in the alkyl substituents at the lower rim of the macrocycle (Fig. 4, 5). The increase of the number of carbon atoms improves sorption on the hydrophobic LiChrosorb RP 18 surface and increases \( k' \) and \( t_R \) values.

The values of \( k' \) and \( t_R \) are correlated with the CA, RA molecular masses and \( \text{Log P} \) values which, in turn, are linearly dependent on the number of carbon atoms in the alkyl substituents.

Such correlations confirm the hydrophobic interactions of CA, RA with octadecyl groups on the LiChrosorb RP 18 surface. Probably, intensification of hydrophobicity of the CA, RA molecules, as well as polarity of the mobile phase can lead to its irreversible sorption on the hydrophobic LiChrosorb RP 18 surface as it was shown in [20]. This approach can be used for physical modification of chromatographic stationary phases by the calixarene-Hosts.

The process of the analyte (A) sorption in the LiChrosorb RP18 (S) – calixarene (L) chromatographic system can be described as follow:

\[
[L][A] \xleftrightarrow{K_1} [LS] + A \xleftrightarrow{K_2} L + S
\]

If the calixarene (L) binds with the support surface (S) stronger than with the analyte (A), \((K_1 > K_2)\), the analyte absorbs on the calixarene modified support surface. When the analyte (A) binds stronger with the calixarene (L), then with the support surface \((K_2 > K_1)\) the supramolecular complex \((LA)\) will absorb on the surface.

To understand the nature of the interaction of calixarenes and calixresorcinarenes with the chromatographic support the molecular modelling of the structure of their Host-Guest complexes with siloxyoctadecane \(H_3\text{SiO(CH}_2)_7\text{CH}_3(SOD)\) as a model site of LiChrosorb RP 18 has been performed.
Fig. 2. Adsorption isotherm of calixarenes CA ($R^2=0.98-0.99$).

Fig. 3. Adsorption isotherm of calixresorcinarenes RA ($R^2=0.98-0.99$).

Fig. 4. Dependence of $k'$ on the number of carbon atoms in the CA aliphatic chain ($R^2 = 0.99$).
In accordance with the modelling the tetrahydroxy-calix[4]arene \( \text{CA-OH} \) forms with \( \text{SOD} \) the complex \( \text{A} \) where the methyl group inserts in the molecular cavity formed by benzene rings of the macrocycle. The molecular cavity has the regular cone conformation stabilized by CH-π interactions of the \( \text{SOD} \) methyl group with the calixarene benzene rings, as well as by the circular system of the intramolecular hydrogen bonds between hydroxyl groups at the lower rim (Fig. 6A). In accordance with the calculations the complex \( \text{B} \) of \( \text{SOD} \) with the \( \text{CA-OH} \) lower rim (Fig. 5B) is significantly weaker compared to complex \( \text{A} \).

The relative energies \( \Delta E \) of the complexes \( \text{A}, \text{B} \) are -8.303 kcal/mol and -2.160 kcal/mol, respectively.

Replacement of the hydrogen atoms in the hydroxyl groups of \( \text{CA-OH} \) to the alkyl groups changes the macrocycle conformation from the regular cone to the flattened cone. This flattening decreases the CH-π interactions in the complex \( \text{C} \) of \( \text{SOD} \) with \( \text{CA-C8} \) (Fig. 6C) compared to the analogous complex \( \text{A} \).

The relative energies \( \Delta E \) of the complexes \( \text{A}, \text{C} \) are -8.303 kcal/mol and -6.073 kcal/mol, respectively.

The complex \( \text{D} \) of \( \text{SOD} \) with \( \text{CA-C8} \) is more stable \( (\Delta E = -9.387 \text{ kcal/mol}) \). In the complex \( \text{SOD} \) deeply penetrates in the lipophilic cavity formed by the calixarene octyl groups and holds there by the van-der-Waals forces (Fig. 6D). In the RP HPLC conditions the complex \( \text{D} \) can be additionally stabilized by hydrophobic interactions.

The similar complex \( \text{E} \) is formed between \( \text{SOD} \) and the \( \text{RA} \). The most stable is the complex \( \text{E} \) (-11.891 kcal/mol) with \( \text{RA-C15} \) where \( \text{SOD} \) penetrates into the lipophilic cavity formed by four pentadecyl groups (Fig. 6E).

Calculation of the complexes \( \text{A-E} \) with 6000 water molecules shows only small changes in their structures compared to those calculated in vacuum.

**Conclusions**

Tetrahydroxy-calix[4]arene, tetraalkoxy-calix[4]arenes and tetraalkyl-calix[4]resorcinarenes are capable to the reversible sorption on the LiChrosorb RP 18 support surface in RP HPLC conditions (the mobile phase is MeCN/H\(_2\)O, 86/14, v/v). The retention times of the compounds studied are in the wide range of 4.67-88.0 min depending on the nature of the substituents in the calixarene macrocyclic skeleton. The retention time increases with the lengthening of the alkyl substituent at the lower rim of the macrocycle.
As it has been shown by the molecular modelling, the most effective sorption is observed for the complexes where the LiChrosorb’s octadecyl groups are included into the lipophilic cavity formed by the alkyl substituents at the calixarene/calix[4]resorcinarene lower rim. Therefore, the molecular cavity formed by the benzene rings of the macrocycle remains opened for complexation with the analyte molecules. Such sorption of tetraalkoxycalixarenes and tetraalkylcalix[4]resorcinarenes on the LiChrosorb RP 18 support surface can significantly improve the resolution of analytes under the reversed-phase high-performance liquid chromatography conditions.

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