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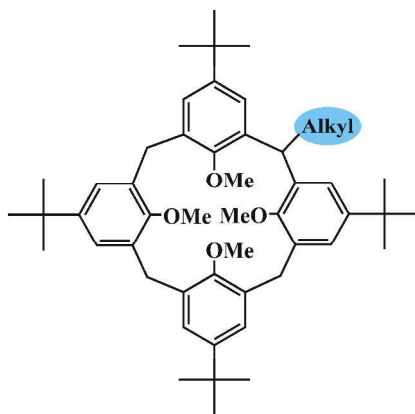
Influence of laterally attached alkyl groups on the conformational behaviour of a basic calix[4]arene: Combined NMR, Molecular Mechanics and X-ray study

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Horizontal broadening:

Calix[4]arenes featuring the structural mode of lateral alkyl substituents were prepared. As proven by special ROESY- and EXSY-2D-NMR experiments, the lateral alkyl group substituents are not exercising a decisive influence on the conformational behaviour of the parent calixarene, suggesting a promising approach for horizontal expansion of the calixarene platform.

**Calixarenes**

*M. Gruner, C. Fischer, T. Gruber,
E. Weber**.....

Influence of laterally attached alkyl groups on the conformational behaviour of a basic calix[4]arene: Combined NMR, Molecular Mechanics and X-ray study

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8 **Influence of laterally attached alkyl groups on the**
9 **conformational behaviour of a basic calix[4]arene: Combined**
10 **NMR, Molecular Mechanics and X-ray study**
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5 Three new calixarenes **3-5** featuring an alkyl residue of different chain length attached
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7 to one of the central ring methylene groups of the basic calix[4]arene **1** have been
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9 prepared. A systematic study that includes also the lower homologous compound **2**,
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11 showing the effect of the alkyl substitution on the conformational behaviour of the
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13 calixarene framework in comparison with the unsubstituted parent compound **1** is
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15 reported. The application of special 2D NMR techniques, 2D-EXSY and ROESY
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17 method, at various temperatures establishes that the calixarenes **2-5** adopt the *partial*
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19 *cone* conformation of lower symmetry and far less the symmetric *cone* and *1,2-alternate*
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21 conformations. In solution, they undergo a fast interconversion with relatively low
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23 activation energies of about 15 kcal/mol at room temperature. The conformer
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25 distribution is reproduced well by molecular mechanistic calculations (MMFF94)
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27 indicating the present conformers to assume the lowest steric energies. A single-crystal
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29 X-ray structure of the lateral ethyl derivative **2** corroborates these results, showing the
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31 molecule in a sterically favourable *partial cone* conformation.
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Keywords: calixarenes; lateral substitution; synthesis; conformational study; solution
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43 2D NMR measurements; X-ray crystal structure
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50 Introduction

51 Calixarenes and their derivatives are an important class of compounds that have been
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53 extensively investigated regarding their supramolecular recognition properties during
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55 the past two decades (1). Calixarenes have also emerged as very attractive building
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57 blocks for the development of highly ingenious functional systems and devices (2). In
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59 both of these respects, much work has been dedicated to the synthesis of calixarenes
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modified on the upper and lower rims, giving rise to an enormous variety of compound

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3 structures (3). Homocalixarenes (4), thiocalixarenes (5) and others (1) where the
4 bridging methylene units of the conventional calixarene framework are completely
5 replaced by modifying organic groups or hetero atoms, represent further currently
6 studied yet rather well known types of structural modifications of the calixarene family
7 (6). By way of contrast, calixarenes featuring a substitution of only one of the
8 methylene bridges are presently rare (7), although these laterally substituted calixarenes
9 may open interesting aspects of calixarene chemistry including linked constructions of
10 oligotopic receptors (8) as well as systems distinguished by particular anchoring of the
11 calixarene to a specific target or support (2, 9). As a result, the conformational
12 behaviour of this particular compound type is also studied only fragmentary in a few
13 cases of modified tetrahydroxycalix[4]- and -calix[5]arenes (7*c-d*). This warrants to
14 deal with the conformational property of laterally monosubstituted calixarenes more
15 broadly. Hence, a series of calixarenes that feature a lateral substitution of alkyl
16 groups of different chain length and includes a well-known parent compound was
17 considered for this purpose. The concrete system of compounds which was chosen for
18 this study, as specified in Fig. 1, covers the tetra-*tert*-butyltetramethoxycalix[4]arene (1)
19 as the reference and 2-5 as the test compounds. This shows an advantage since the
20 conformational behaviour of 1 is known rather detailed from previous work (10).
21 Moreover, for the synthesis of 2-5 one can follow a literature method elaborated for the
22 calixarene 2 by Scully (7*a*). The conformational properties of 2-5, which are described
23 here, were determined from a detailed solution NMR study. This involves
24 measurements at various temperatures and uses special 2D NMR techniques to yield a
25 full description of the conformer distribution and the free activation energies of
26 conformational interconversion. We also report on the crystal structure of the
27 corresponding laterally ethyl monosubstituted calixarene 2, being the first crystal
28 structure of a laterally monosubstituted tetramethoxycalix[4]arene.

Results and discussion

Synthesis

The calixarene **1**, which is the basic compound for the synthesis of **2-5**, was obtained from two-stage methylation reaction of 5,11,17,23-tetra-*tert*-butyl-25,26,27,28-tetrahydroxycalix[4]arene (*11*) using in a starting phase-transfer process sodium hydroxide, dimethyl sulphate and tetrabutylammonium bromide to yield the intermediate compound with two distal methyl groups, followed by treatment with sodium hydride and iodomethane in a final methylation step (*12*). A corresponding one-step methylation method of the parent tetrahydroxycalixarene with sodium hydride and iodomethane, described in the literature (*10a*, *13*), did not prove competitive in its result. The target calixarenes **2-5** featuring different laterally substituted alkyl groups were prepared by deprotonation of **1** with *n*-butyllithium and reaction with respective alkyl halides following the procedure described by Scully (*7a*).

Conformational study

Conformational predominance and conformational flexibility are important structural parameters of a calixarene molecule, being decisive of its supramolecular behaviour both in solution and solid state (*1-3*). While the so-called *cone* conformation, providing a spaceous cavity structure, is perhaps the most frequent conformation of the existing calixarenes (*1*, *14*), the other types of molecular conformations, including *partial cone*, *1,2-* and *1,3-alternate* conformation, are somewhat more rarely found though not less important for a designed purpose (*15*). As a rule, these conformations are mainly determined by the ring size or the nature and the space required of the substituents being present at the upper and lower rims of the calixarene molecule. Conformational property

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3 will also depend on the physical condition, including type of solvent as well as the
4 presence of a proper guest molecule. However, in no manner, *O*-alkylated calixarenes
5 that feature an alkyl residue laterally attached to one of the methylene groups of the
6 central ring have been studied conformationally. Only analogous
7 tetrahydroxycalixarenes were taken for a corresponding study (7c-d).

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10 A well-known molecular structure is that of the tetra-*tert*-
11 butyltetramethoxycalix[4]arene (**1**), which was chosen as the parent compound for the
12 test. According to the NMR data (10), **1** in a solution of CDCl₃ exists in an equilibrium
13 of all four conventional conformers of which the *partial cone* conformation is the
14 preferred one. A *partial cone* conformer is also reported for the solvent free calixarene **1**
15 (16) as well as for the furan (17) and dichloromethane (18) inclusion compounds in the
16 solid state, while a corresponding sodium complex shows a *cone* conformation both in
17 the crystal structure and in CDCl₃/CD₃CN solution due to coordination of the sodium
18 ion (10b). Thus, it is clear that a potential change of the conformational behaviour in the
19 present cases of calixarenes, in comparison with **1**, is attributed to the laterally
20 substituted alkyl groups.

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22 A complex pattern of signals in the NMR spectra of the alkyl substituted
23 calixarenes **2-5** in both CDCl₃ and tetrachloroethane-d₂ (TCl-d₂) at various temperatures
24 indicates that the compounds exist in different conformers, as demonstrated exemplary
25 for the lateral propyl derivative **3** in Fig. 2. To minimize the signal broadening and
26 separate the overlapping region of methine and methylene signals, we carried out the
27 NMR experiments at an optimized temperature of 269 K, yielding three different signals
28 for the methine protons between 4.47 and 4.79 ppm in CDCl₃ for compounds **2-5**.
29 Detailed 2D NMR measurements using COSY-, ¹H/¹³C-correlated HSQC (19) and
30 HMBC (20) as well as NOESY and ROESY experiments gave a full determination of
31 conformers (specified in Fig. 3; ¹H and ¹³C chemical shifts are summarized in Tables
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1S-4S of the Supplementary Information). A special denotation of the calixarene scaffold in the three observed conformations is illustrated in Fig. 3, indicating similarities in the chemical shift for different pairs of atoms.

Depending on the alignment of the four arene units with reference to the lateral substituent, there are 16 different conformational isomers possible (Fig. 4). Six of them are mirror images of each other resulting in identical NMR spectra. The 2D experiments show that the laterally substituted *n*-alkyl calixarenes, studied here, exist mainly in one *partial cone* conformation (*pacoD* in Fig.4), with an equatorial substituent between two *syn* orientated arene units, and also as *1,2-alternate* (*alt2C* in Fig. 4) and *cone* (*coneB* in Fig. 4) conformers (Scheme 1), revealed in the region of the OCH₃ groups of the HSQC spectrum (Fig. 5). In the ROESY spectrum (Fig. 6), all conformers possess stronger positive cross peaks between the axial methine H-1 protons in the bridge and the near H-8 protons of the methoxy groups than between these protons and the H-3 protons of the same aromatic rings, indicating the vicinity of one of the methoxy groups on the lower rim to H-1. Further characteristic specific interactions between aromatic protons of the calixarene and protons of OCH₃ or *tert*-butyl groups neighbouring the methylene bridge are marked in Fig. 6. These correlations suggest the orientation of arene units of the calixarene ring for each individual isomer. For example, the *paco* isomer is characterized by the cross peaks 8▲'/11▲', 16▲ /13▲, 16▲'/5▲' and 16▲'/13▲, and the *1,2-alternate* species by the 8■/11■ and 8■ /13■ relation.

Furthermore, in Fig. 6, negative cross-peaks between different H-1 proton signals can be observed due to chemical exchange processes in the form of a dynamic interconversion of conformers. To extract the rate constants for the conformational chemical exchange (k_{chem}), the 2D-EXSY method (21) was used, and thus the free activation energies $\Delta G^{\#}$ of the interconversion has been calculated (Table 1). Interestingly, compounds 2-5 display a fast *partial cone/cone* interconversion involving

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3 values of free activation energy rather comparable to the reported ~13 kcal/mol for the
4 laterally unsubstituted prototype calixarene **1** in the same solvent (*10a*). The observed
5 influence of solvent and temperature on the conformational behaviour is in agreement
6 with the common expectation. In this connection, the conformational reorientation was
7 found to be faster at higher temperature and retarded in solution of the larger and less
8 polar solvent molecules of tetrachloroethane-*d*₂ (TCI-*d*₂) compared to CDCl₃.
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17 The conformer distribution of **1** and **2-5**, summarized in Table 2, shows that the
18 *partial cone* conformation predominates the present series of tetra-*tert*-
19 butyltetramethoxycalix[4]arenes in CDCl₃, varying only little by introduction of the
20 lateral alkyl substituents, i.e. the predominance of the *partial cone* conformation is only
21 slightly affected. Nevertheless, its frequency decreases to some extent for the benefit of
22 the rarely occurring *cone*- and *1,2-alternate* structures in the following manner. With
23 increasing chain length, the ratio of *partial cone* conformation remains nearly
24 unchanged, while the *cone* conformation decreases and the *1,2-alternate* conformation
25 increases. However, as contrasted with the prototype calixarene **1**, for none of the
26 studied compounds a *1,3-alternate* conformation was detected. This points to a
27 predominance of thermodynamically stable conformers and suggests the avoidance of
28 an isoclinical arrangement of the lateral substituent, presumably for energetic reasons.
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48 **MM Calculation**

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52 To evaluate the steric influence of the lateral alkyl substituent, relative energies of the
53 different conformers of **2** as a model compound, summarized in Fig. 4, were determined
54 using the MMFF94 force field (22) as implemented in the SYBYL software. As proven
55 by previous mechanistic studies (7*d*), all conformations with an axial arrangement of the
56 lateral substituent are energetically discriminated due to their higher angle bending
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energy. The predominance of the *partial cone* conformation resulting from the NMR experiments, bearing the lateral substituent in an equatorial position between *syn* orientated arene units in solution (*pacoD*) may thus be explained with the avoidance of repulsion between the α -C-atom of the lateral alkyl moiety and the neighbouring methoxy groups, indicated by their low sterical energies (Table 3). Analogously, the frequency and stability of the *coneB* conformer detected by NMR can be assumed due to its similar low ground state energy. It is also shown that the energy gap between the *1,2-alternate* (*alt2C*) and the *1,3-alternate* conformer is small, giving evidence of their potential presence in solution. The fact that only the *1,2-alternate* conformer is detected in the NMR could result from its higher energy gap to the lowest energy *partial cone* conformer (~ 3.8 kcal/mol for *alt2C*) leading to separate signals, whereas the interconversion to the *1,3-alternate* conformer is too fast on the NMR time scale.

Crystal structure of compound 2·CHCl₃

In order to compare the conformational results from the solution NMR study with the conformational behaviour in the solid state, we tried to obtain suitable crystals of the compounds **2-5** for X-ray study. Unfortunately, we were unsuccessful in getting high-quality crystals in the case of **3-5**, presumably due to the high flexibility of the rather extended alkyl chains. However, the corresponding laterally ethyl substituted compound **2** yielded colourless crystals (space group *Pca2₁*) from a solvent mixture of methanol and chloroform (2:1) suitable for X-determination (Table 4). The crystals were analyzed to be a stoichiometric 1:1 solvate with chloroform. Not surprisingly, also in the solid state, one low energy *partial cone* conformer bearing the lateral substituent in an equatorial position between *syn* orientated arene units is found (Fig. 7). This result is in agreement both with the predominance of the *partial cone* conformation in solution and the lowest energy conformer (*pacoD*) from MM calculation.

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3 Selected conformational parameters of the calixarene molecule **2** are summarized
4 in Tab. S5. The opposite phenyl rings A and C differ only little from coplanarity [8.9°
5 (1)], while the aromatic rings B and D are orientated in an angle of 42.8° (1). Similar to
6 a comparable 2:1 solvate of the lateral unsubstituted calixarene **1** with dichloromethane
7 (18), the methoxy group of ring D points outwards and maintains no interactions with
8 the other rings. In consequence of the compact structure caused by the *partial cone*
9 conformer, the calixarene does not allow a cavitate inclusion of solvent molecules,
10 which therefore are located interstitially in the crystal lattice. With reference to the
11 crystal packing, the calixarene molecules are orientated translatory staggered along the
12 crystallographic *b* axis (Fig. 8). Corresponding to the nonpolar character of the
13 calixarene, the interactions are limited to weak C-H \cdots π -contacts (23) involving guest
14 and calixarene molecules [$d(\text{C1GA}\cdots\text{centroid C}) = 3.569$ (1) Å] or the calixarene
15 molecules among each other [$d(\text{C33}\cdots\text{centroid B}) = 3.486$ (1) Å] as specified in Tab. S6.
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39 Conclusions

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41 According to the literature (1, 7), calixarenes showing a substitution of a single
42 methylene bridge, especially in the case of an alkyl substitution, are a rather low
43 developed class of compounds. As derivatives, of the archetype
44 tetramethoxycalix[4]arene **1**, only the first two members of the potential compound
45 series, i.e. the methyl- and ethyl (**2**) derivatives have previously been synthesized and
46 preliminarily described regarding their conformational behaviour. With the present
47 calixarenes **3-5** featuring the corresponding mode of lateral substitution but using higher
48 alkyl groups of different chain length, new examples of this particular compound type
49 have now been made available. They enable to perform a first systematic study of the
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3 influence of the given alkyl groups on the conformational property of the
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6 tetramethoxycalix[4]arene system.

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8 The following main conclusions can be obtained from this study: (1) The lateral
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10 attachment of an alkyl residue to one of the central ring methylene groups of the basic
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12 calixarene **1** has no substantial consequence on the conformational behaviour. That is,
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14 independent of the chain length, a respective substitution of an alkyl group does not
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16 hamper the predominance of the *partial cone* conformation of **1** in chloroform solution,
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18 as proven by NMR spectroscopy. This particular conformation was also found to be low
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20 energetic in the calculated sterical energies. Moreover, the crystal structure of
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22 compound **2** in its solvated form with chloroform is in structural agreement, showing
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24 the *partial cone* conformation in the crystalline state. (2) As against the unsubstituted
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26 calixarene **1**, the free activation energy for the conformational interconversion of **2-5** in
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28 solution is found only moderately higher. Hence, it is shown that (3) an alkyl
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30 substitution of one of the hydrogen atoms of a methylene group between the flanking
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32 aryl residues of the central ring offers a very promising possibility for coupling of a
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34 basic tetramethoxy-calix[4]arene structural element to spacer or linker units in order to
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36 realize future developments of new complex framework structures without substantial
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38 impairment of the initial calixarene conformation, while corresponding modifications at
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40 the upper and lower rim sites normally result in distinct changes of the conformational
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42 behaviour.
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50 Future work is required to explore the effect of more than one laterally attached
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52 alkyl groups or of a substituent containing polar or functional units in the side chain.
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54 Investigations into this direction are aiming work of our group.
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Experimental

Materials

1-Iodopropane (Fluka), 1-iodopentane (Acros) and 1-bromodecane (Fluka) were used as received. The *n*-BuLi (1.6 M in hexane) was purchased from Acros. Analytical TLC was performed on precoated silica gel plates (Merck, 60 F₂₅₄). Organic solvents were purified by standard procedures (24).

Techniques

Melting points were determined with a hot-stage microscope (VEB Dresden Analytik) and are uncorrected. ¹H and ¹³C NMR spectra were recorded using a Bruker DRX 500 spectrometer. The chemical shifts (δ) are reported in ppm relative to TMS = 0 ppm. Mass spectra were recorded with an EI-MS (Finnigan MAT 8200). The elemental analyses were performed with a Heraeus CHN rapid analyzer.

Synthesis of 5,11,17,23-tetra-*tert*-butyl-25,26,27,28-tetramethoxycalix[4]arene (1)

A mixture of 5,11,17,23-tetra-*tert*-butyl-25,26,27,28-tetrahydroxycalix[4]arene (11), NaOH (aqueous, 50 p.c.), dimethyl sulphate and tetrabutylammonium bromide in toluene was reacted following a known procedure (12). Recrystallization from MeOH/CHCl₃ (1:1) yielded 60% of intermediate 5,11,17,23-tetra-*tert*-butyl-26,28-dihydroxy-25,27-dimethoxycalix[4]arene as colorless solid, m.p. 235-238°. Following the literature protocol (13), this compound was caused to react with NaH and iodomethane to afford 67% of calixarene **1** as a colorless solid, m.p. 242-244 °C (lit.(12) 242-243 °C).

General procedure for the synthesis of the laterally monoalkylated calix[4]arenes 2-5

The synthesis of the products **2-5** follows the protocol described by Scully et al. (7a). To a stirred solution of **1** (1.0 mmol) in dry THF (30 ml), *n*-BuLi (1.6 N in *n*-hexane, 2.9 ml, 4.6 mmol) was added at room temperature under argon. After having stirred for 30 min, the respective 1-halogenalkane (8.0 mmol) was added via syringe to the solution, causing a change of its color from red to yellow. Stirring was continued for 1 h. The solvent was removed under reduced pressure and the residue partitioned into water/dichloromethane (1:1, 60 ml). The organic layer was washed with water (2 x 30 ml), dried (MgSO₄) and evaporated under reduced pressure. Recrystallization from MeOH yielded the products as colorless solids. Experimental, MS and elemental analysis data of the individual compounds are specified below. For the detailed 2D NMR experiments, all substances were dried for 24 h under vacuum at 50 °C. The respective NMR data of **2-5** are given in the Supplementary Materials (Tables S1-S4). The NMR data for **2** correspond to the literature (7a).

5,11,17,23-Tetra-tert-butyl-25,26,27,28-tetramethoxy-2-ethylcalix[4]arene (2)

1-Bromoethane was used. Yield 0.42 g (57%), mp. 183-185 °C [lit. (7a) 184-186 °C].

5,11,17,23-Tetra-tert-butyl-25,26,27,28-tetramethoxy-2-n-propylcalix[4]arene (3)

1-Iodopropane was used. Yield 0.34 g (46%), mp. 132-135 °C. Anal. Calcd for C₅₁H₇₀O₄ · 0.5 MeOH: C 81.06, H 9.51. Found: C 80.99, H 9.78 %. MS (70 eV), *m/z* (%): 747 (M⁺, 100), 748 (52), 358 (35), 177 (28).

5,11,17,23-Tetra-tert-butyl-25,26,27,28-tetramethoxy-2-n-pentylcalix[4]arene (4)

1-Iodopentane was used. Yield 0.35 g (47%), mp. 76-80 °C. Anal. Calcd for C₅₃H₇₄O₄: C 82.12, H 9.62. Found: C 82.15, H 9.80 %. MS (70 eV), *m/z* (%): 775 (M⁺, 100), 776 (64), 372 (43), 177 (14).

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5,11,17,23-Tetra-tert-butyl- -25,26,27,28-tetramethoxy-2-n-decylcalix[4]arene (5)

1-Bromodecane was used. Yield 0.37 g (44%), mp. 139-144 °C. Anal. Calcd for $C_{58}H_{84}O_4 \cdot 1.5 \text{ MeOH}$: C 80.00, H 10.15. Found: C 79.85, H 9.88 %. MS (70 eV), *m/z* (%): 845 (M^+ , 100), 846 (63), 407 (42), 177 (32).

X-ray crystallography

Crystals of **2** suitable for X-ray diffraction analysis were obtained by slow recrystallization from methanol/chloroform (2:1) solution. The intensity data were collected on a Bruker APEX II diffractometer with MoK_α radiation ($\lambda = 0.71073 \text{ \AA}$) using ω - and φ -scans. Reflections were corrected for background and Lorentz polarization effects. Preliminary structure models were derived by application of direct methods (25) and were refined by full-matrix least squares calculation based on F^2 for all reflections (26). All hydrogen atoms were included in the models in calculated positions and were refined as constrained to bonding atoms. Relevant crystal data together with the refinement details are listed in Table 4.

Crystallographic data for the structural analysis have been deposited with the Cambridge Crystallographic Data Centre, No. CCDC 748215. Copies of this information may be obtained free of charge from: The Director, CCDC, 12 Union Road, Cambridge, CB2 1EZ, UK. Fax (+44) 1223-336-033; E-mail: deposit@ccdc.cam.ac.uk or www: www.ccdc.cam.ac.uk.

Supplementary materials

¹H and ¹³C NMR data of **2-5** in CDCl₃ and tetrachloroethane-d₂, selected conformational parameters and geometric parameters of potential hydrogen bond type interactions from the crystal structure of **2** · CHCl₃ are available as Supplementary Materials, available online.

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Table 1. (a) Rate constants k_{chem} and (b) free activation energies ΔG^\ddagger of conformational interconversion according to the methine protons H-1 of compounds **2-5** and for comparison of compound **1**, calculated by the 2D-EXSY method.

(a)

Interconversion	$k_{\text{chem}} (\text{s}^{-1})$									
	2				3				4	5
	CDCl ₃ , 295 K	CDCl ₃ , 264 K	TCl-d ₂ , 295 K	TCl-d ₂ , 263K	CDCl ₃ , 295 K	CDCl ₃ , 263 K	TCl-d ₂ , 295 K	TCl-d ₂ , 260K	CDCl ₃ , 265 K	CDCl ₃ , 265 K
paco/cone	fast	18.9	fast	-	fast	18.2	fast	-	15.9	19.5
paco/1,2-alt.	52.0	0.82	14.4	0.7	14.5	0.5	7.3	0.4	0.5	1.3

(b)

Interconversion	$\Delta G^\ddagger (\text{kcal mol}^{-1})^{\text{a,b}}$									
	2				3				4	5
paco/cone	-	13.8	-	-	-	13.7	-	-	13.9	13.7
paco/1,2-alt.	14.9	15.4	15.7	15.6	15.7	15.6	16.1	15.6	15.7	15.2

^a Interconversion paco/1,2-alt. for **3** at coalescence in TCl-d₂ (360 K): 17.5 kcal mol⁻¹.

^b Interconversion paco/1,2-alt. for unsubstituted compound **1** at coalescence in TCl-d₂ (363 K): 16.4 kcal mol⁻¹.

Table 2. Mole fractions of conformers for compounds **1-5** in solution.

conformer	1				2				3				4	5
	CDCl ₃ , 295 K	CDCl ₃ , 265 K	TCI-d ₂ , 295 K	TCI-d ₂ , 263 K	CDCl ₃ , 295 K	CDCl ₃ , 261 K	TCI-d ₂ , 295 K	TCI-d ₂ , 260 K	CDCl ₃ , 295 K	CDCl ₃ , 261 K	TCI-d ₂ , 295 K	TCI-d ₂ , 260 K	CDCl ₃ , 264 K	CDCl ₃ , 263 K
paco (▲,▲')	85	84	83	84	90	74	88	79	88	72	87	80	73	71
cone (o)	7	9	6	6	-	17	-	10	-	17	-	10	16	15
1,2-alt. (M)	8	7	10	9	10	9	12	11	12	10	13	10	11	14

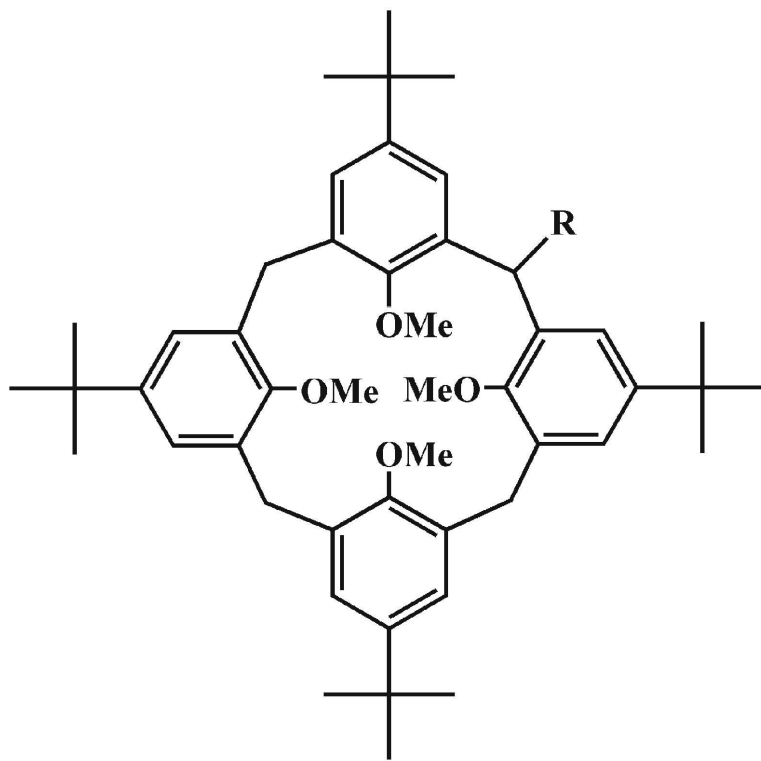
Table 3. Relative energies (in kcal mol⁻¹) for possible conformers of ethyl compound **2** obtained by the MMFF94 force field (SYBYL) and calculated populations.

Conformation	Relative energy (kcal mol ⁻¹)	Relative to <i>pacoD</i> (kcal mol ⁻¹)	Population (%)
<i>coneA</i>	225.76	0.01	47.13
<i>coneB</i>	233.15	7.40	-
<i>pacoA</i>	233.99	8.24	-
<i>pacoB</i>	227.77	2.02	1.58
<i>pacoC</i>	227.58	1.83	2.18
<i>pacoD</i>	225.75	0.00	47.93
<i>alt2A</i>	237.02	11.27	-
<i>alt2B</i>	233.00	7.25	-
<i>alt2C</i>	229.50	3.75	0.08
<i>alt3</i>	227.98	2.23	1.11

Table 4. Crystal data and selected details of the data collection and refinement calculations of compound **2** · CHCl₃.

Compound	2 · CHCl ₃
Empirical formula	C ₅₁ H ₆₉ Cl ₃ O ₄
Formula weight (g mol ⁻¹)	852.4
Crystal system	orthorhombic
Space group	Pca2 ₁
<i>a</i> (Å)	23.9269(5)
<i>b</i> (Å)	16.8437(4)
<i>c</i> (Å)	12.6634(3)
α (°)	90.00
β (°)	90.00
γ (°)	90.00
<i>V</i> (Å ³)	5103.6(2)
<i>Z</i>	4
<i>F</i> (000)	1832
<i>D</i> _c (Mg m ⁻³)	1.109
μ (mm ⁻¹)	0.219
Data collection	
Temperature (K)	93(2)
No. of collected reflections	57108
within the θ -limit (°)	1.21- 28.10
Index ranges $\pm h, \pm k, \pm l$	-31/31, -22/21, -16/10
No. of unique reflections	9944
<i>R</i> _{int}	0.0968
Refinement calculations: full-matrix least- squares on all <i>F</i> ² values	
Weighting expression <i>w</i> ^a	$[\sigma^2(F_o^2) + (0.1788P)^2 + 4.6405P]^{-1}$
No. of refined parameters	537
No. of F values used [<i>I</i> > 2σ(<i>I</i>)]	8817
Final <i>R</i> -Indices	
<i>R</i> (=Σ Δ <i>F</i> / Σ <i>F</i> _o)	0.0843
<i>wR</i> on <i>F</i> ²	0.2405
<i>S</i> (=Goodness of fit on <i>F</i> ²)	1.092
Final Δρ _{max} /Δρ _{min} (e Å ⁻³)	0.000

$$^a P = (F_o^2 + 2F_c^2)$$



1 R = H

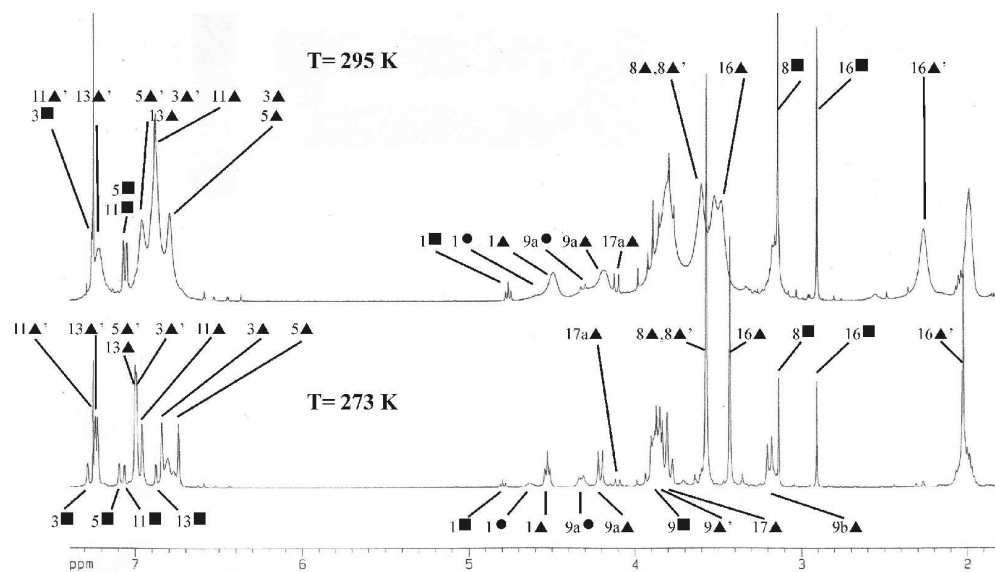
2 R = C₂H₅

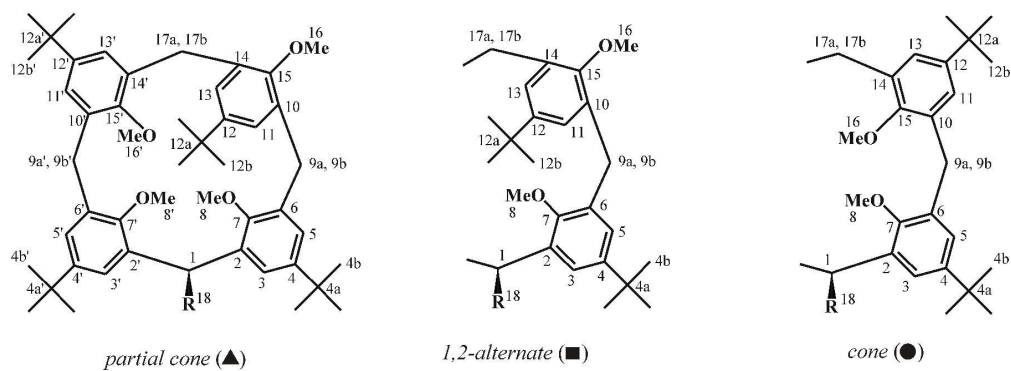
3 R = *n*-C₃H₇

4 R = *n*-C₅H₁₁

5 R = *n*-C₁₀H₂₁

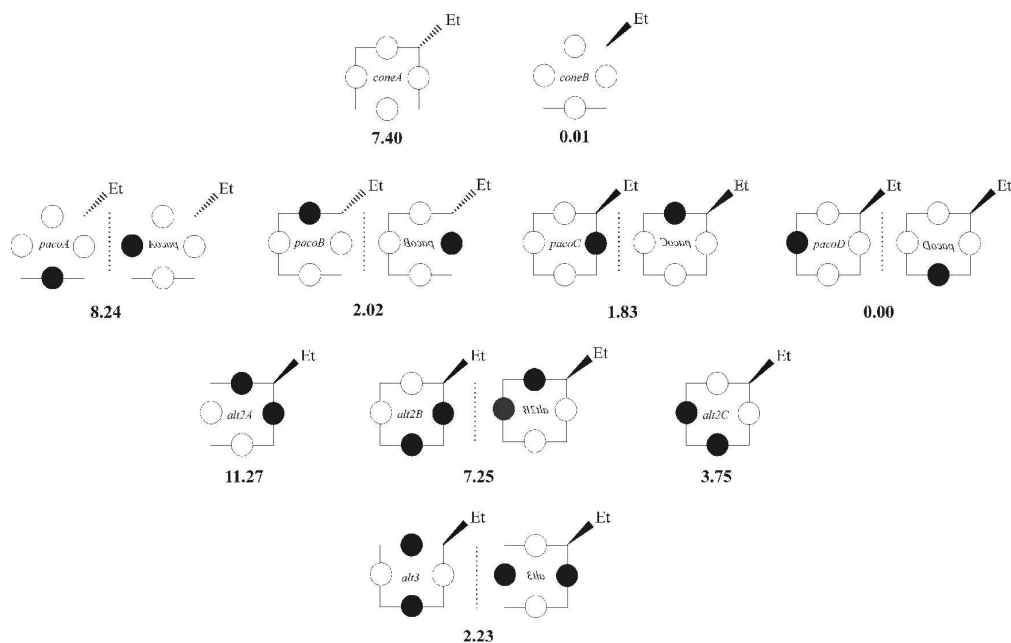
560x877mm (72 x 72 DPI)



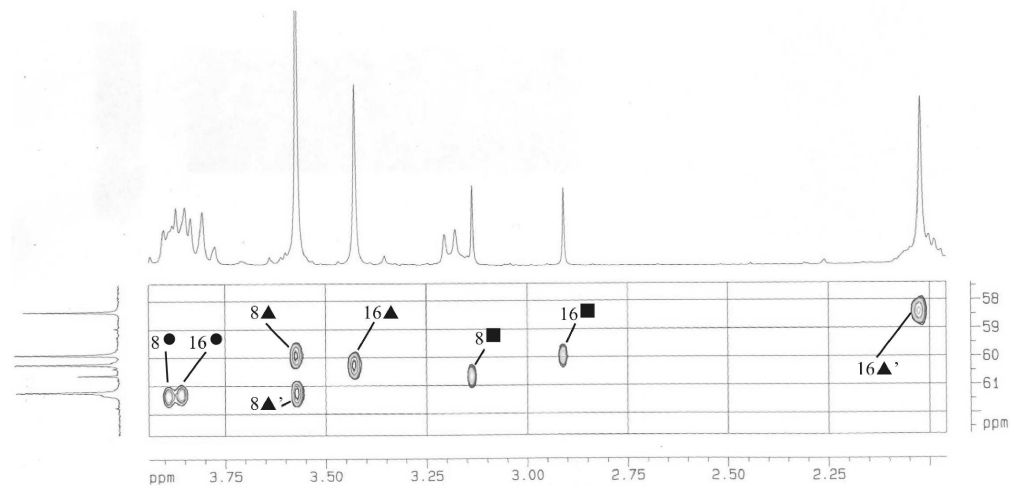


- 2: R = CH₂¹⁹CH₃²⁰
 3: R = CH₂¹⁹CH₂²⁰CH₃²¹
 4: R = CH₂¹⁹CH₂²⁰CH₂²¹CH₂²²CH₃²³
 5: R = CH₂¹⁹CH₂²⁰CH₂²¹CH₂²²CH₂²³CH₂²⁴CH₂²⁵CH₂²⁶CH₂²⁷CH₃

1432x751mm (96 x 96 DPI)

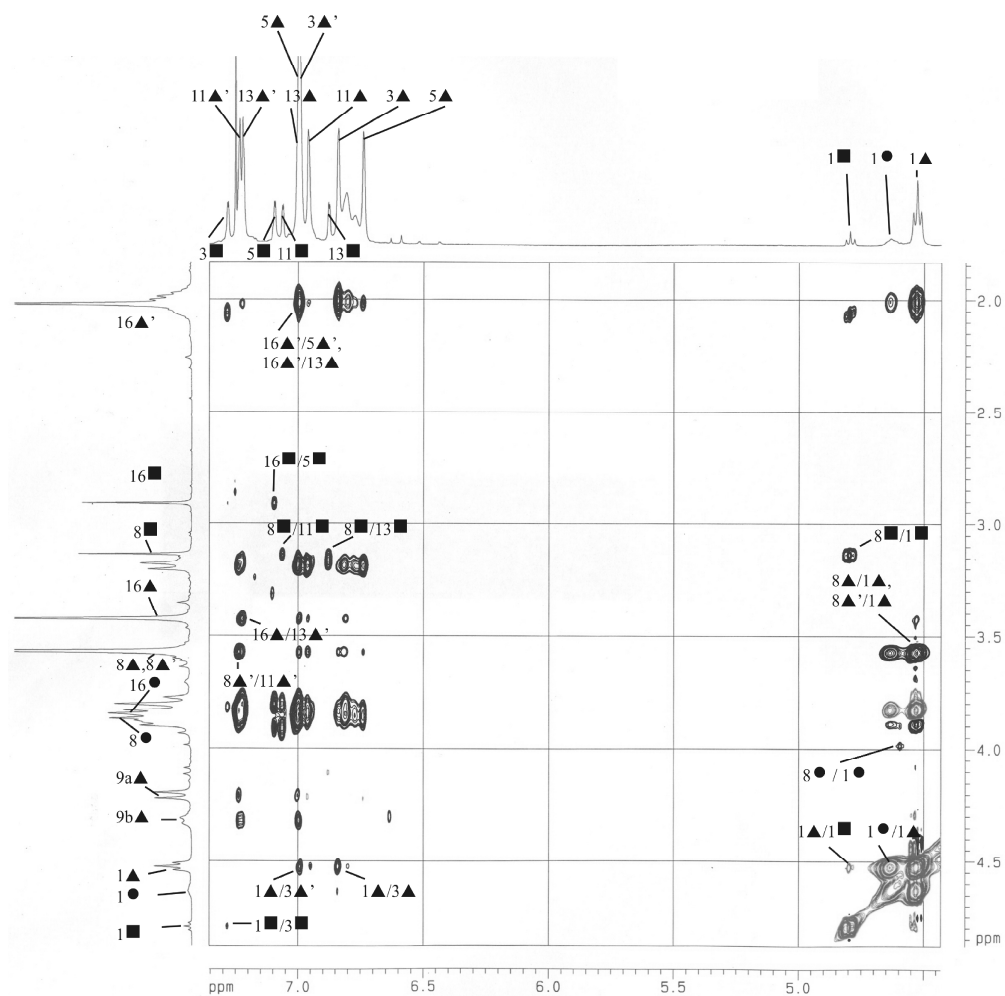


1844x1160mm (72 x 72 DPI)



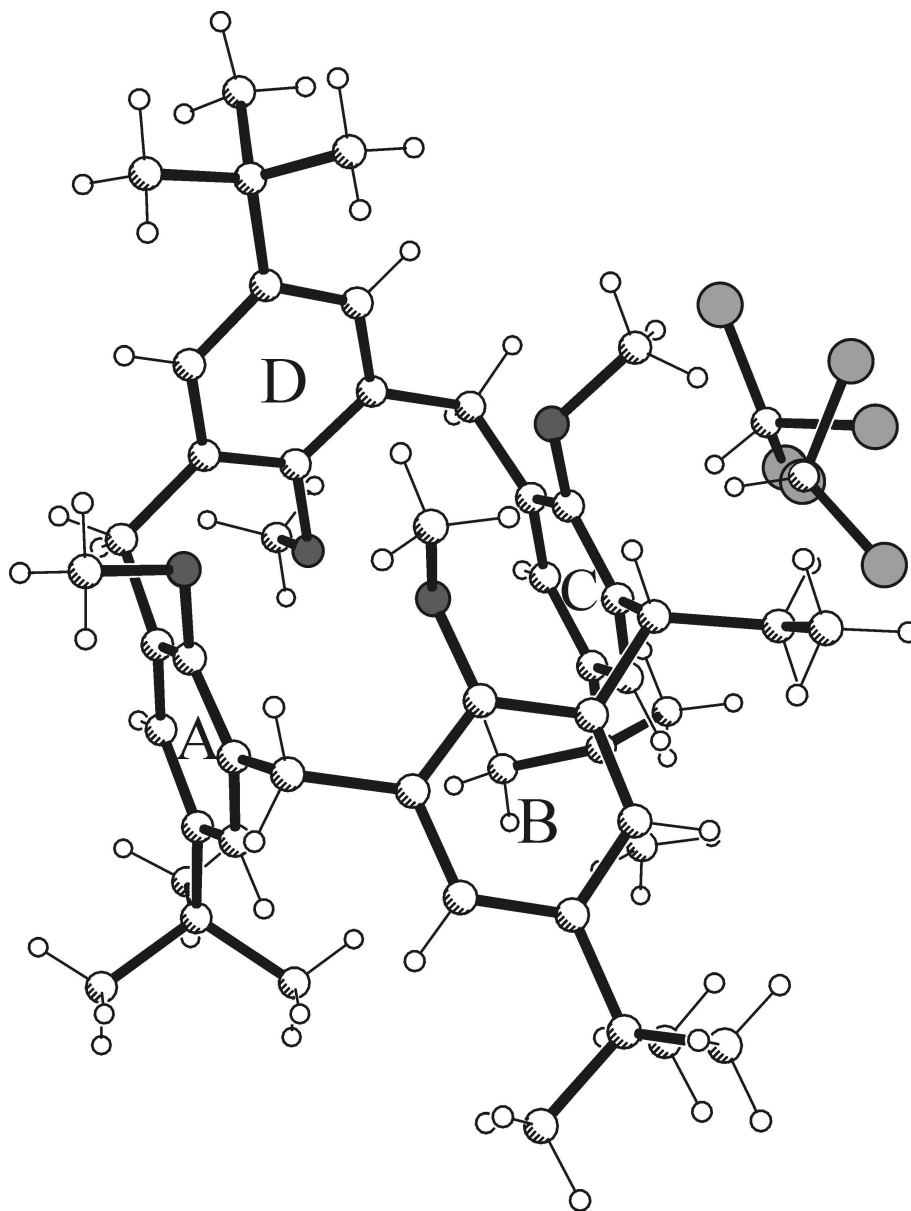
1603x786mm (72 x 72 DPI)

Review Only

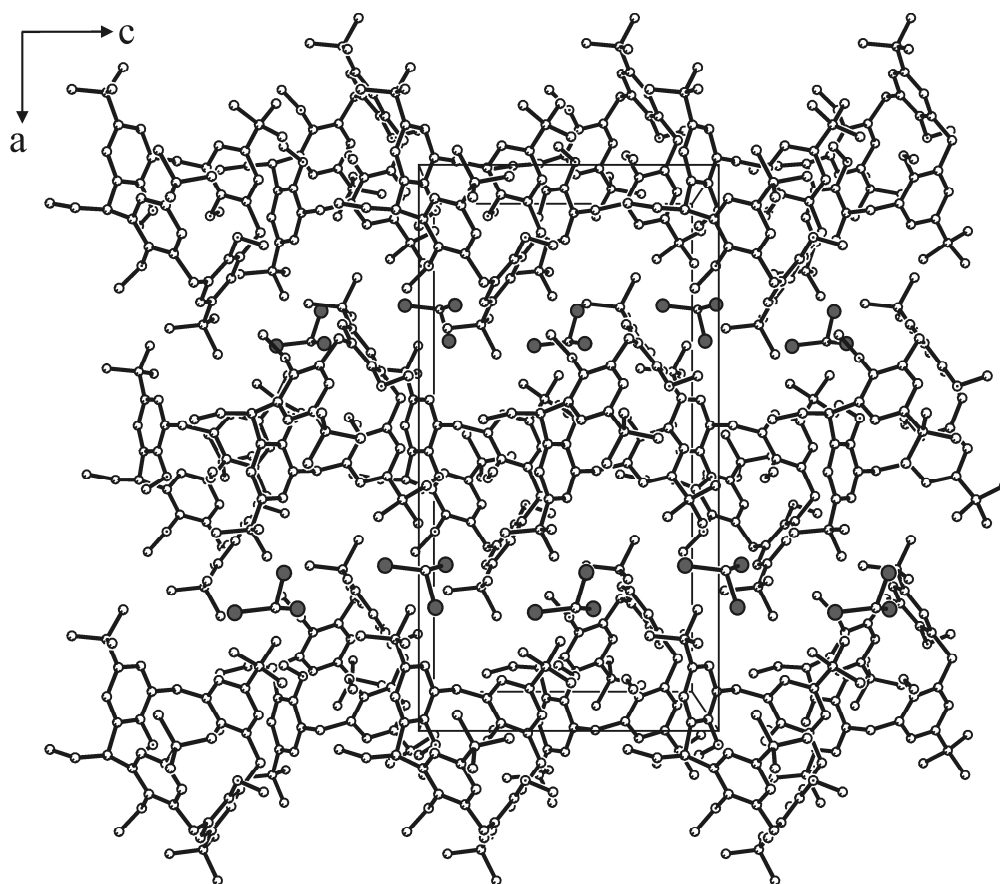


1623x1603mm (72 x 72 DPI)

MUN

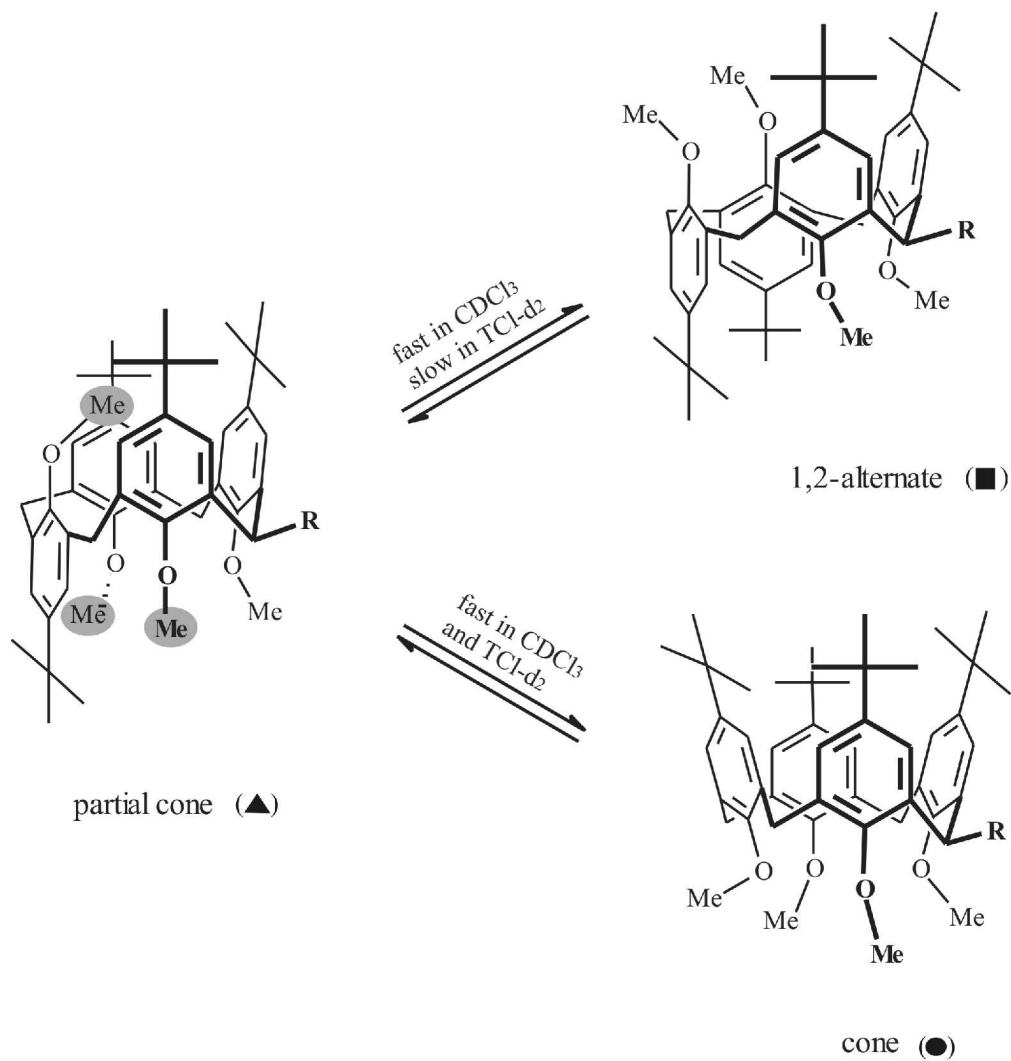


981x1292mm (72 x 72 DPI)



1497x1305mm (72 x 72 DPI)

Only



1108x1189mm (72 x 72 DPI)



Supplementary Materials

Table S1a. ¹H-NMR-spectrum of **2** in CDCl₃ at 264 K.

paco (▲,▲')	1,2-alternate(O)	cone(M)
7.24 (s, br, 1H, H-11')	7.28 (d, ⁴ J=2.0 Hz, 2H, H-3)	7.2 (s, br, H-11)
7.22 (s, br, 1H, H-13')	7.10 (d, ⁴ J=1.8 Hz, 2H, H-5)	6.81 (s, br, H-13)
7.00 (s, br, 2H, H-5')	7.06 (d, ⁴ J=1.5 Hz, 2H, H-11)	6.81 (s, br, H-3)
6.99 (s, br, 1H, H-13)	6.86 (d, ⁴ J=1.5 Hz, 2H, H-13)	6.76 (s, br, H-5)
6.98 (s, br, 1H, H-3')		
6.96 (s, br, 1H, H-11)		
6.84 (s, br, 1H, H-3)		
6.74 (s, br, 1H, H-5)		
4.39 (t, ³ J=8.0 Hz, H-1)	4.70 (t, ³ J=8.2 Hz, 1H, H-1)	4.48 (t, br, 1H, H-1)
4.21 (d, ² J=12.8 Hz, H-9a)	4.10 (d, ² J=13.4 Hz, 1H, H-17a)	4.32 (d, br, H-9a)
3.90 (d, ² J=15.4 Hz, H-9a')	3.19 (1H, H-17b)	3.18 (H-9b)
3.86 (d, ² J=15.4 Hz, H-17a)	3.93 (1H, H-9a)	4.12 (H-17a)
3.80 (d, ² J~15 Hz, H-9b')	3.78 (1H, H-9b)	3.17 (H-17b)
3.79 (d, ² J=15.3 Hz, H-17b)		
3.19 (d, ² J=13.0 Hz, H-9b)		
3.57 (s, 3H, H-8')	3.14 (s, 6H, H-8)	3.87 (H-8)
3.56 (s, 3H, H-8)	2.90 (s, 6H, H-16)	3.85 (H-16)
3.42 (s, 3H, H-16)		
2.03 (s, 3H, H-16')		
1.39 (s, 9H, H-12b')	1.29 (s, 18H, H-4b)	not discriminable
1.23 (s, 9H, H-12b)	1.25 (s, 18H, H-12b)	
1.22 (s, 9H, H-4b')		
1.09 (s, 9H, H-4b)		
2.06 (m, 2H, H-18)	2.09 (m, 2H, H-18)	2.06 (H-18)
1.05 (t, 3H, ³ J= 7.2 Hz, H-19)	0.99 (t, 3H, H-19)	1.05 (H-19)

Table S1b. ^{13}C -NMR-spectrum of **2** in CDCl_3 at 264 K.

	paco ($\blacktriangle, \blacktriangle'$)	1,2-alternate (O)	cone (M)
154.84 (C _i -15)	155.11 (C _i -7')	154.64 (C _i -15)	154.9 (br, C _i -15)
154.59 (C _i -7)	154.64 (C _i -15')	153.79 (C _i -7)	
144.91 (C _i -12)	145.01 (C _i -4')	144.45 (C _i -12)	144.3 (br, C _i -4,12)
144.53 (C _i -4)	143.22 (C _i -12')	144.00 (C _i -4)	
136.84 (C _i -2)	137.84 (C _i -2')	137.97 (C _i -2)	137.1 (br, C _i -2)
135.24 (C _i -10)	133.13 (C _i -10')	135.11 (C _i -14)	133.9 (br, C _i -10)
133.34 (C _i -6)	133.13 (C _i -14')	132.68 (C _i -10)	133.8 (br, C _i -14)
132.77 (C _i -14)	132.74 (C _i -6')	132.60 (C _i -6)	133.5 (br, C _i -6)
125.62 (CH,C-13)	126.39 (CH, C-13')	126.15 (CH, C-5)	126.3 (br, C-11)
125.10 (CH,C-11)	126.30 (CH, C-11')	125.86 (CH, C-11)	124.4 (br, C-13)
124.53 (CH,C-5)	125.41 (CH,C-5')	125.31 (CH, C-13)	124.0 (br, C-5)
121.84(CH,C-3)	121.85 (CH,C-3')	122.04 (CH,C-3)	121.4 (br,C-3)
60.24 (CH ₃ ,C-16)	59.90 (CH ₃ ,C-8')	60.64 (CH ₃ ,C-8)	61.23 (CH ₃ ,C-8)
61.21 (CH ₃ ,C-8)	58.36 (CH ₃ ,C-16')	59.91 (CH ₃ ,C-16)	61.20 (CH ₃ ,C-16)
36.89 (CH,C-1)		36.27 (CH,C-1)	37.39 (CH, C-1)
36.86 (CH ₂ ,C-17)	37.55 (CH ₂ ,C-9')	36.86 (CH ₂ ,C-9)	31.0 (CH ₂ , C-9)
31.00 (CH ₂ ,C-9)		30.11 (CH ₂ ,C-17)	30.9 (CH ₂ , C-17)
26.87 (CH ₂ ,C-18)		27.72 (CH ₂ ,C-18)	26.87 (CH ₂ , C-18)
13.48 (CH ₃ ,C-19)		13.39 (CH ₃ ,C-19)	13.50 (CH ₃ , C-19)
3.88 (C,C-4a)	34.02 (C,C-4a')	34.04 (C,C-4a)	33.8 (br, C, C-4a, 12a)
33.88 (C,C-12a)	33.88 (C,C-12a')	33.92 (C,C-12a)	
31.41 (CH ₃ ,C-12b)	31.53 (CH ₃ ,C-12b')	31.47 (CH ₃ ,C-4b)	
31.28 (CH ₃ ,C-4b)	31.36 (CH ₃ ,C-4b')	31.36 (CH ₃ ,C-12b)	

Table S1c. $^1\text{H-NMR}$ -spectrum of **2** in $\text{TCl}_4\text{-d}_2$ at 263 K.

paco (▲,▲')	1,2-alternate(O)	cone(M)
7.06 (s, br, 1H, H-11')	7.06 (s, br, H-3)	7.09 (H-11)
7.05 (s, br, 1H, H-13')	6.92 (s, br, H-5)	6.6-6.7 (H-3, 5, 13)
6.92 (s, br, 1H, H-3')	6.90 (d,br, 2H, H-11)	
6.87 (s, br, 1H, H-11)	6.74 (d,br, 2H, H-13)	
6.83 (s, br, 2H, H-13,5')		
6.59 (s, br, 1H, H-3)		
6.52 (s, br, 1H, H-5)		
4.25 (t, $^3\text{J}=7.8$ Hz, H-1)	4.43 (t, $^3\text{J}=7.8$ Hz, 1H, H-1)	4.3 (t, br, H-1)
4.06 (d, $^2\text{J}=12.5$ Hz, H-9a)	3.96 (d, 1H, H-17a)	4.14 (br, H-9a)
3.75 (d, H-17a)	3.73 (d, 2H, H-9a)	3.03 (br, H-9b)
3.69 (d, H-17b)	3.64 (d, 2H, H-9b)	4.00 (br, H-17a)
3.09 (d, $^2\text{J}=12.4$ Hz, H-9b)	3.02 (d, 1H, H-17b)	3.05 (br, H-17b)
3.73 (d, $^2\text{J}\sim 10$ Hz, H-9a')		
3.70 (d, $^2\text{J}\sim 10$ Hz, H-9b')		
3.48 (s, 3H, H-8)	2.96 (s, 6H, H-8)	3.74 (H-8)
3.39 (s, 3H, H-8')	2.76 (s, 6H, H-16)	3.71 (H-16)
3.25 (s, 3H, H-16)		
1.60 (s, 3H, H-16')		
1.25 (s, 9H, H-12b')	1.15 (s, 18H, H-4b)	1.29 (H-12b)
1.12 (s, 18H, H-4b',12b')	1.12 (s, 18H, H-12b)	1.11 (H-4b)
0.90 (s, 9H, H-4b)		
1.96 (m, 1H, H-18a)	1.93 (m, 1H, H-18)	not discriminable
1.89 (m, 1H, H-18b)	0.86 (t, 3H, H-19)	
0.90 (t, 3H, H-19)		

Table S1d. ^{13}C -NMR-spectrum of **2** in TCI-d_2 at 263 K.

	paco (▲,▲')	1,2-alternate(O)	cone(M)
154.75 (C _i -7)	155.47 (C _i -7')	154.04 (C _i -7)	
155.10 (C _i -15)	154.84 (C _i -15')	154.75 (C _i -15)	
144.60 (C _i -4)	145.32 (C _i -4')	144.27 (C _i -4)	
145.28 (C _i -12)	143.55 (C _i -12')	145.32 (C _i -12)	
137.00 (C _i -2)	138.39 (C _i -2')	138.30 (C _i -2)	
136.05 (C _i -10)	133.38 (C _i -10')	132.95 (C _i -10)	
133.10 (C _i -14)	133.38 (C _i -14')	135.33 (C _i -14)	
133.09 (C _i -6)	133.10 (C _i -6')	132.83 (C _i -6)	
125.93 (CH,C-13)	126.68 (CH, C-11')	126.43 (CH, C-5)	126.7 (CH, C-11)
125.61 (CH,C-11)	(CH, C-13')	126.28 (CH, C-11)	124.9 (CH, C-13)
124.87 (CH,C-5)	125.60 (CH,C-5')	125.74 (CH, C-13)	124.4 (CH, C-5)
122.47 (CH,C-3)	122.03 (CH,C-3')	122.47 (CH,C-3)	121.9 (CH,C-9)
61.84 (CH ₃ ,C-8)	60.29 (CH ₃ ,C-8')	60.85 (CH ₃ ,C-8)	61.5 (<i>br</i> , CH ₃
60.53 (CH ₃ ,C-16)	58.77 (CH ₃ ,C-16')	60.29 (CH ₃ ,C-16)	C-8;16)
37.89 (CH,C-1)		36.73 (CH,C-1)	
31.36 (CH ₂ ,C-9)		37.31 (CH ₂ ,C-9)	
37.31 (CH ₂ ,C-17)	38.28 (CH ₂ ,C-9')	30.25 (CH ₂ ,C-17)	
27.32 (CH ₂ ,C-18)		28.19 (CH ₂ ,C-18)	
14.01 (CH ₂ ,C-19)		14.01 (CH ₂ ,C-19)	
34.37 (C,C-4a)	34.22 (C,C-4a')	34.33 (C,C-4a)	
34.15 (C,C-12a)	34.18 (C,C-12a')	34.18 (C,C-12a)	
31.82 (CH ₃ ,C-12b)	31.91 (CH ₃ ,C-4b')	31.87 (CH ₃ ,C-4b)	
31.62 (CH ₃ ,C-4b)	31.87 (CH ₃ ,C-12b')	31.81 (CH ₃ ,C-12b)	

Table S2a. $^1\text{H-NMR}$ -spectrum of **3** in CDCl_3 at 269 K.

paco ($\blacktriangle, \blacktriangle'$)	1,2-alternate(O)	cone(M)
7.23 (s, br, 1H, H-11')	7.28 (d, $^4J=2.2$ Hz, 2H, H-3)	7.23 (s, br, H-11)
7.22 (s, br, 1H, H-13')	7.09 (d, $^4J=2.1$ Hz, 2H, H-5)	6.80 (s, br, H-13)
7.00 (s, br, 2H, H-5', 13)	7.06 (d, $^4J=2.2$ Hz, 2H, H-11)	6.80 (s, br, H-3)
6.99 (s, br, 1H, H-3')	6.87 (d, $^4J=2.2$ Hz, 2H, H-13)	6.77 (s, br, H-5)
6.96 (s, br, 1H, H-11)		
6.84 (s, br, 1H, H-3)		
6.74 (s, br, 1H, H-5)		
4.53 (t, $^3J=7.8$ Hz, H-1)	4.79 (t, $^3J=9.0$ Hz, 1H, H-1)	4.63 (t, br, 1H, H-1)
4.21 (d, $^2J=12.8$ Hz, H-9a)	4.10 (d, $^2J=13.3$ Hz, 1H, H-17a)	4.32 (d, $^2J=11.2$ Hz, H-9a)
3.91 (d, $^2J=15.4$ Hz, H-9a')	3.16 (d, 1H, H-17b)	3.19 (d, $^2J\sim 12$ Hz, H-9b)
3.87 (d, $^2J=15.4$ Hz, H-17a)	3.92 (d, $^2J\sim 16$ Hz, 2H, H-9a)	4.13 (H-17a)
3.82 (d, $^2J\sim 15$ Hz, H-9b')	3.79 (d, $^2J\sim 16$ Hz, 2H, H-9b)	3.18 (H-17b)
3.78 (d, $^2J=15.3$ Hz, H-17b)		
3.19 (d, $^2J=13.0$ Hz, H-9b)		
3.57 (s, 3H, H-8)	3.13 (s, 6H, H-8)	3.88 (H-8)
3.57 (s, 3H, H-8')	2.91 (s, 6H, H-16)	3.87 (H-16)
3.43 (s, 3H, H-16)		
2.02 (s, 3H, H-16')		
1.38 (s, 9H, H-12b')	1.29 (s, 18H, H-4b)	1.22 (H-4b)
1.23 (s, 9H, H-4b')	1.25 (s, 18H, H-12b)	1.09 (H-12b)
1.22 (s, 9H, H-12b)		
1.09 (s, 9H, H-4b)		
2.00 (m, 2H, H-18)	2.05 (m, 2H, H-18)	2.00 (H-18)
1.46 (m, 2H, H-19)	1.39 (H-19)	1.46 (H-19)
1.01 (t, 3H, $^3J=7.2$ Hz, H-20)	0.96 (t, 3H, $^3J=7.2$ Hz, H-20)	1.01 (t, $^3J=7.2$ Hz, H-20)

Table S2b. ^{13}C -NMR-spectrum of **3** in CDCl_3 at 269 K.

	paco (▲,▲')	1,2-alternate(O)	cone(M)
154.56 (C _i -7)	154.98 (C _i -7')	153.64 (C _i -7)	152.2 (br, C _i -7)
154.82 (C _i -15)	154.56 (C _i -15')	154.61 (C _i -15)	
144.49 (C _i -4)	144.91 (C _i -4')	143.96 (C _i -4)	144.35 (C _i -4)
144.91 (C _i -12)	143.22 (C _i -12')	144.43 (C _i -12)	144.35 (C _i -12)
136.91 (C _i -2)	138.05 (C _i -2')	138.09 (C _i -2)	137.22 (C _i -2)
135.30 (C _i -10)	133.11 (C _i -10')	135.10 (C _i -14)	133.84 (C _i -14)
133.27 (C _i -14)	133.11 (C _i -14')	132.68 (C _i -10)	133.83 (C _i -10)
132.78 (C _i -6)	133.72 (C _i -6')	132.60 (C _i -6)	133.52 (C _i -6)
125.61 (CH,C-13)	126.36 (CH, C-13')	126.11 (CH, C-5)	126.26 (CH, C-5)
125.10 (CH,C-11)	126.27 (CH, C-11')	125.87 (CH, C-11)	124.48 (CH, C-11)
124.49 (CH,C-5)	125.37 (CH,C-5')	125.30 (CH, C-13)	124.02 (CH, C-13)
121.93 (CH,C-3)	121.93 (CH,C-3')	122.13 (CH,C-3)	121.52 (CH,C-3)
61.22 (CH ₃ ,C-8)	59.89 (CH ₃ ,C-8')	60.63 (CH ₃ ,C-8)	61.23 (CH ₃ , C-8)
60.23 (CH ₃ ,C-16)	58.36 (CH ₃ ,C-16')	59.89 (CH ₃ ,C-16)	61.21 (CH ₃ , C-16)
35.10 (CH,C-1)		34.31 (CH,C-1)	35.56 (CH, C-1)
30.98 (CH ₂ ,C-9)	37.84 (CH ₂ ,C-9')	36.88 (CH ₂ ,C-9)	36.98 (CH ₂ ,C-9)
36.83 (CH ₂ ,C-17)		30.06 (CH ₂ ,C-17)	30.98 (CH ₂ ,C-17)
36.15 (CH ₂ ,C-18)		37.09 (CH ₂ ,C-18)	36.15 (CH ₂ ,C-18)
21.53 (CH ₂ ,C-19)		21.71 (CH ₂ ,C-19)	21.53 (CH ₂ ,C-19)
14.39 (CH ₃ ,C-20)		14.23 (CH ₃ ,C-20)	14.39 (CH ₂ ,C-20)
33.89 (C,C-4a)	33.89 (C,C-4a')	34.02 (C,C-4a)	not discriminable
33.89 (C,C-12a)	33.89 (C,C-12a')	33.89 (C,C-12a)	
31.27 (CH ₃ ,C-4b)	31.41 (CH ₃ ,C-4b')	31.48 (CH ₃ ,C-4b)	
31.36 (CH ₃ ,C-12b)	31.52 (CH ₃ ,C-12b')	31.36 (CH ₃ ,C-12b)	

Table S2c. $^1\text{H-NMR}$ -spectrum of **3** in TCI-d_2 at 265 K.

paco (▲,▲')	1,2-alternate(O)	cone(M)
7.08 (s, br, 1H, H-11')	7.08 (H-3)	7.11 (H-11)
7.06 (s, br, 1H, H-13')	6.93 (d, $^4J=2.2$ Hz, 2H, H-5)	6.69 (H-13)
6.94 (s, br, 1H, H-3')	6.92 (d, $^4J=2.0$ Hz, 2H, H-11)	6.66 (H-3)
6.88 (s, br, 1H, H-11)	6.77 (d, $^4J=1.9$ Hz, 2H, H-13)	6.61 (H-5)
6.85 (s, br, 2H, H-13,5')		
6.61 (s, br, 1H, H-3)		
6.53 (s, br, 1H, H-5)		
4.40 (t, $^3J=7.7$ Hz, H-1)	4.55 (t, $^3J=8.0$ Hz, 1H, H-1)	4.46 (H-1)
4.07 (d, $^2J=12.6$ Hz, H-9a)	3.97 (d, $^2J=13.0$ Hz, 1H, H-17a)	4.13 (H-9a)
3.74 (d, $^2J=15.7$ Hz, H-17a)	3.74 (d, $^2J=15.5$ Hz, 2H, H-9a)	3.03 (H-9b)
3.68 (d, $^2J=15$ Hz, H-17b)	3.65 (d, $^2J=15.9$ Hz, 2H, H-9b)	4.00 (H-17a)
3.07 (d, $^2J=12.8$ Hz, H-9b)	3.03 (d, $^2J=13.7$ Hz, 1H, H-17b)	3.06 (H-17b)
3.73 (d, $^2J=10$ Hz, H-9a')		
3.70 (d, $^2J=10$ Hz, H-9b')		
3.49 (s, 3H, H-8)	2.97 (s, 6H, H-8)	3.77 (H-8)
3.40 (s, 3H, H-8')	2.78 (s, 6H, H-16)	3.75 (H-16)
3.27 (s, 3H, H-16)		
1.62 (s, 3H, H-16')		
1.27 (s, 9H, H-12b')	1.16 (s, 18H, H-4b)	1.27 (H-12b)
1.13 (s, 18H, H-4b', 12b')	1.13 (s, H-12b)	1.13 (H-4b)
0.92 (s, 9H, H-4b)		
1.94 (m, 1H, H-18a)	1.89 (m, 1H, H-18)	1.94 (H-18a)
1.83 (m, 1H, H-18b)	1.27 (m, 1H, H-19)	1.83 (H-18b)
1.35 (m, 2H, H-19)	0.85 (t, 3H, $^3J=7.3$ Hz, H-20)	1.35 (H-19)
0.90 (t, 3H, $^3J=7.3$ Hz, H-20)		0.90 (H-20)

Table S2d. ^{13}C -NMR-spectrum of **3** in TCI-d_2 at 265 K.

	paco (▲,▲')	1,2-alternate(O)	cone(M)
154.74 (C _i -7)	155.34 (C _i -7')	153.94 (C _i -7)	
155.07 (C _i -15)	154.73 (C _i -15')	154.69 (C _i -15)	
144.55 (C _i -4)	145.34 (C _i -4')	144.22 (C _i -4)	
145.27 (C _i -12)	143.54 (C _i -12')	144.55 (C _i -12)	
137.07 (C _i -2)	138.61 (C _i -2')	138.45 (C _i -2)	
136.15 (C _i -10)	133.36 (C _i -10')	132.95 (C _i -10)	
133.12 (C _i -14)	133.36 (C _i -14')	135.32 (C _i -14)	
133.01 (C _i -6)	133.01 (C _i -6')	132.81 (C _i -6)	
125.91 (CH,C-13)	126.63 (CH, C-11')	126.39 (CH, C-5)	126.6 (CH, C-5)
125.62 (CH,C-11)	(CH, C-13')	126.30 (CH, C-11)	124.8 (CH, C-11)
124.83 (CH,C-5)	125.57 (CH,C-5')	125.74 (CH, C-13)	124.3 (CH, C-13)
122.60 (CH,C-3)	122.08 (CH,C-3')	122.60 (CH,C-3)	122.0 (CH,C-3)
61.87 (CH ₃ ,C-8)	60.28 (CH ₃ ,C-8')	60.82 (CH ₃ ,C-8)	61.6 (<i>br</i> , CH ₃ C-8;16)
60.52 (CH ₃ ,C-16)	58.77 (CH ₃ ,C-16')	60.33 (CH ₃ ,C-16)	
35.46 (CH,C-1)		34.60 (CH,C-1)	36.0 (<i>br</i> , CH, C-1)
31.36 (CH ₂ ,C-9)		37.35 (CH ₂ ,C-9)	
37.26 (CH ₂ ,C-17)	38.25 (CH ₂ ,C-9')	30.48 (CH ₂ ,C-17)	
36.57 (CH ₂ ,C-18)		37.35 (CH ₂ ,C-18)	
22.01 (CH ₂ ,C-19)		22.28 (CH ₂ ,C-19)	
14.90 (CH ₃ ,C-20)		14.77 (CH ₃ ,C-20)	
34.13 (C,C-4a)	34.23 (C,C-4a')	34.32 (C,C-4a)	
34.38 (C,C-12a)	34.19 (C,C-12a')	34.19 (C,C-12a)	
31.60 (CH ₃ ,C-4b)	31.88 (CH ₃ ,C-4b')	31.82 (CH ₃ ,C-4b)	
31.82 (CH ₃ ,C-12b)	31.88 (CH ₃ ,C-12b')	31.88 (CH ₃ ,C-12b)	

Table S3a. ¹H-NMR-spectrum of **4** in CDCl₃ at 265 K.

paco (▲,▲')	1,2-alternate(O)	cone(M)
7.21 ((d), 1H, H-11')	7.26 (d, H-3)	7.20 (H-11)
7.20 ((d), 1H, H-11')	7.07 (d, ⁴ J=2.4 Hz, 2H, H-5)	6.78 (H-13)
6.98 (((d), 1H, H-5')	7.04 (d, ⁴ J=2.3 Hz, 2H, H-11)	6.78 (H-3)
6.97 ((d), 2H, H-3')	6.86 (d, ⁴ J=2.4 Hz, 2H, H-13)	6.75 (H-5)
6.97 ((d), 1H, H-13)		
6.94 ((d), 1H, H-11)		
6.81 ((d), 1H, H-3)		
6.72 ((d), 1H, H-5)		
4.47 (t, ³ J=7.9 Hz, H-1)	4.73 (t, ³ J=8.2 Hz, 1H, H-1)	4.57 ((t), H-1)
4.18 (d, ² J=12.8 Hz, H-9a)	4.08 (d, ² J=13.3 Hz, 1H, H-17a)	4.30 (br, H-9a)
3.86 (d, ² J=14.0 Hz, H-17a)	3.87 (d, ² J=14.7 Hz, 2H, H-9a)	3.17 (br, H-9b)
3.84 (d, ² J=13.0 Hz, H-9a')	3.75 (d, 2H, H-9b)	not discriminable
3.83 (d, ² J=14.2, H-17b)		
3.80 (d, ² J=13.8 Hz, H-9b')		
3.17 (d, ² J=13.0 Hz, H-9b)		
3.55 (s, 3H, H-8)	3.11 (s, 6H, H-8)	3.88 (H-8)
3.55 (s, 3H, H-8')	2.89 (s, 6H, H-16)	3.83 (H-16)
3.40 (s, 3H, H-16)		
2.00 (s, 3H, H-16')		
1.36 (s, 9H, H-12b')	1.27 (s, 18H, H-4b)	not discriminable
1.20 (s, 9H, H-4b')	1.23 (s, 18H, H-12b)	
1.20 (s, 9H, H-12b)		
1.07 (s, 9H, H-4b)		
1.97 (m, H-18)	2.03 (m, H-18)	1.97 (m, H-18)
1.41 (m, H-19)	1.34 (m, H-19)	not discriminable
1.39 (m, H-20)	1.39 (m, H-20)	
1.32 (m, H-21)	1.31 (m, H-21)	
0.87 (t, ³ J=7.0 Hz, 3H, H-22)	0.82 (m, H-22)	

Table S3b. ^{13}C -NMR-spectrum of **4** in CDCl_3 at 265 K.

	paco ($\blacktriangle, \blacktriangle'$)	1,2-alternate (O)	cone (M)
154.83 (C _i -15)	154.99 (C _i -7')	154.61 (C _i -15)	not discriminable
154.55 (C _i -7)	154.57 (C _i -15')	153.66 (C _i -7)	
144.92 (C _i -12)	144.92 (C _i -4')	144.44 (C _i -12)	not discriminable
144.50 (C _i -4)	143.24 (C _i -12')	143.99 (C _i -4)	
136.93 (C _i -2)	138.05 (C _i -2')	138.14 (C _i -2)	not discriminable
135.32 (C _i -10)	133.12 (C _i -10')	135.09 (C _i -14)	
133.28 (C _i -6)	133.12 (C _i -14')	132.69 (C _i -10)	
132.78 (C _i -14)	132.75 (C _i -6')	132.61 (C _i -6)	
125.61 (CH,C-13)	126.36 (CH, C-13')	126.11 (CH, C-5)	126.3 (br, C-11)
125.11 (CH,C-11)	126.27 (CH, C-11')	125.88 (CH, C-11)	124.5 (br, C-13)
124.50 (CH,C-5)	125.37 (CH,C-5')	125.30 (CH, C-13)	124.1 (br, C-5)
121.95 (CH,C-3)	121.91 (CH,C-3')	122.11 (CH,C-3)	121.7 (br,C-3)
61.27 (CH ₃ ,C-8)	59.91 (CH ₃ ,C-8')	60.63 (CH ₃ ,C-8)	61.3 (C-8)
60.24 (CH ₃ ,C-16)	58.37 (CH ₃ ,C-16')	59.91 (CH ₃ ,C-16)	61.3 (C-16)
35.52 (CH,C-1)		34.57 (CH,C-1)	36.1 (br, C-1)
36.82 (CH ₂ ,C-17)	37.85 (CH ₂ ,C-9')	36.90 (CH ₂ ,C-9)	31.1 (br, C-9)
31.09 (CH ₂ ,C-9)		30.06 (CH ₂ ,C-17)	not discriminable
34.03 (CH ₂ ,C-18)		34.85 (CH ₂ ,C-18)	not discriminable
32.04 (CH ₂ ,C-19)		31.09 (CH ₂ ,C-19)	
28.21 (CH ₃ ,C-20)		28.30 (CH ₃ ,C-20)	
22.60 (CH ₃ ,C-21)		22.50 (CH ₃ ,C-21)	
14.32 (CH ₃ ,C-22)		14.22 (CH ₃ ,C-22)	
33.89 (C,C-4a)	33.89 (C,C-4a')	33.89 (C,C-4a)	not discriminable
33.89 (C,C-12a)	33.89 (C,C-12a')	33.89 (C,C-12a)	
31.42 (CH ₃ ,C-12b)	31.52 (CH ₃ ,C-12b')	31.48 (CH ₃ ,C-4b)	
31.27 (CH ₃ ,C-4b)	31.42 (CH ₃ ,C-4b')	31.37 (CH ₃ ,C-12b)	

Table S4a. $^1\text{H-NMR}$ -spectrum of **5** in CDCl_3 at 263 K.

paco (▲,▲')	1,2-alternate(O)	cone(M)
7.22 ((d), 1H, H-11')	7.26 (d, ⁴ J=2.4 Hz, 2H; H-3)	7.21 (H-11)
7.21 ((d), 1H, H-13')	7.08 (d, ⁴ J=2.3 Hz, 2H, H-5)	6.79 (H-13)
6.99 (1H, H-5')	7.05 (d, ⁴ J=2.3 Hz, 2H, H-11)	6.79 (H-3)
6.98 (1H, H-3')	6.86 (d, ⁴ J=2.4 Hz, 2H, H-13)	6.76 (H-5)
6.98 (1H, H-13)		
6.95 ((d), 1H, H-11)		
6.82 ((d), 1H, H-3)		
6.72 ((d), 1H, H-5)		
4.48 (t, ³ J=7.7 Hz, H-1)	4.74 (t, ³ J=8.0 Hz, 1H, H-1)	4.58 ((t), H-1)
4.20 (d, ² J=12.8 Hz, H-9a)	4.09 (d, ² J=13.3 Hz, 1H, H-17a)	4.30 (br, H-9a)
3.87 (d, ² J~15.0 Hz, H-17a)	3.89 (d, ² J=14.7 Hz, 2H, H-9a)	3.18 (br, H-9b)
3.86 (d, ² J=13.5 Hz, H-9a')	3.76 (d, 2H, H-9b+)	not discriminable
3.82 (d, ² J=14.1 Hz, H-17b)		
3.81 (d, ² J=13.7 Hz, H-9b')		
3.18 (d, ² J=12.8 Hz, H-9b)		
3.56 (s, 3H, H-8)	3.12 (s, 6H, H-8)	3.87 (s, H-8)
3.56 (s, 3H, H-8')	2.89 (s, 6H, H-16)	3.82 (s, H-16)
3.41 (s, 3H, H-16)		
2.00 (s, 3H, H-16')		
1.98 (m, H-18)	1.86 (m, H-18)	2.01 (m, H-18)
1.42 (m, H-19)	1.42 (m, H-19)	not discriminable
1.36 (m, H-20)	not discriminable	
1.26-1.24 (m, H-21a-d)		
1.35 (m, H-22)		
1.26 (m, H-23)		
0.86 (t, 3H, H-24)		
1.38 (s, 9H, H-12b')		

Table S4b. ^{13}C -NMR-spectrum of **5** in CDCl_3 at 263 K.

	paco (▲,▲')	1,2-alternate(O)	cone(M)
154.81 (C _i -15)	154.97 (C _i -7')	154.60 (C _i -15)	154.8 (C _i -7)
154.54 (C _i -7)	154.54 (C _i -15')	153.63 (C _i -7)	not discriminable
144.90 (C _i -12)	144.92 (C _i -4')	144.44 (C _i -12)	not discriminable
144.50 (C _i -4)	143.24 (C _i -12')	143.99 (C _i -4)	
136.90 (C _i -2)	138.04 (C _i -2')	138.15 (C _i -2)	not discriminable
135.34 (C _i -10)	133.10 (C _i -10')	135.10 (C _i -14)	
133.25 (C _i -6)	133.10 (C _i -14')	132.66 (C _i -10)	
132.74 (C _i -14)	132.76 (C _i -6')	132.59 (C _i -6)	
125.59 (CH,C-13)	126.33 (CH, C-13')	126.10 (CH, C-5)	126.3 (br, C-11)
125.10 (CH,C-11)	126.25 (CH, C-11')	125.87 (CH, C-11)	124.5 (br, C-13)
124.47 (CH,C-5)	125.34 (CH,C-5')	125.29 (CH, C-13)	124.0 (br, C-5)
121.95 (CH,C-3)	121.90 (CH,C-3')	122.10 (CH,C-3)	121.5 (br,C-3)
61.27 (CH ₃ ,C-8)	59.90 (CH ₃ ,C-8')	60.63 (CH ₃ ,C-8)	61.3 (C-8)
60.23 (CH ₃ ,C-16)	58.35 (CH ₃ ,C-16')	59.90 (CH ₃ ,C-16)	61.3 (C-16)
35.50 (CH,C-1)		34.56 (CH,C-1)	36.0 (br, C-1)
36.79 (CH ₂ ,C-17)	37.83 (CH ₂ ,C-9')	36.85 (CH ₂ ,C-9)	31.0 (br, C-9)
30.98 (CH ₂ ,C-9)		30.06 (CH ₂ ,C-17)	not discriminable
33.88 (CH ₂ ,C-18)		32.69 (CH ₂ ,C-18)	not discriminable
28.46 (CH ₂ ,C-19)		28.10 (CH ₂ ,C-19)	
29.81 (CH ₃ ,C-20)		not discriminable	
29.63, 29.60, 29.53, 29.34 (CH ₂ , C-21a-d)			
31.87 (CH ₂ ,C-22)			
22.67 (CH ₂ ,C-23)			
14.21 (CH ₃ ,C-24)			
33.87 (C,C-4a)	33.87 (C,C-4a')	34.02 (C,C-4a)	not discriminable
33.87 (C,C-12a)	33.87 (C,C-12a')	34.02 (C,C-12a)	
31.41 (CH ₃ ,C-12b)	31.51 (CH ₃ ,C-12b')	31.46 (CH ₃ ,C-4b)	
31.25 (CH ₃ ,C-4b)	31.35 (CH ₃ ,C-4b')	31.36 (CH ₃ ,C-12b)	

Table S5. Selected conformational parameters of the calixarene **2** · CHCl₃.

Compound	2 · CHCl ₃
Interplanar angles (°) ^a	
mpla ^b /A	83.68 (1)
mpla/B	43.73 (1)
mpla/C	87.43 (1)
mpla/D	86.46 (1)
A/C	8.90 (1)
B/D	42.77 (1)

^a Aromatic rings: ring A: C(1)...C(6); ring B: C(8)...C(13); ring C: C(15)...C(20); ring D: C(22) ...C(27);

^b Best plane through atoms C(7), C(14), C(21) and C(28).

Table S6. Distances und angles of possible hydrogen-bond type interactions of **2**.

Atoms involved	Symmetry	Distances (Å)		Angles (°)
		D...A	H...A	
C1G-H1G...centroid(C) ^a	x, y, z	3.8855 (10)	2.90 (1)	170
C33-H33A...centroid(B) ^a	-x, 1-y, ½z	3.4855 (10)	2.62 (1)	147
C1GA-H1GA...centroid(C) ^a	x, y, z	3.5692 (10)	2.63 (1)	157

^a Centroid means center of gravity of the respective aromatic ring: ring A: C(1)...C(6); ring B: C(8)...C(13); ring C: C(15)...C(20); ring D: C(22)...C(27).