Tribenzotriquinacenes bearing three peripheral or bridgehead urea groups stretched into the 3-D space
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Abstract
The syntheses of tribenzotriquinacenes (TBTQ) bearing three phenylurea groupings at either the arene periphery or at the benzhydrylic bridgeheads of the rigid, convex–concave, $C_{3v}$-symmetrical molecular framework are reported. $^1$H NMR data point to supramolecular aggregation of these TBTQ derivatives in low-polarity solvents.

Introduction
In the course of our extended research on polyfunctionalized derivatives of tribenzotriquinacene (TBTQ, 1) and some hydrocarbon congeners 2–4, which provide a great variety of convex–concave molecular building blocks [1-7], we focused our attention on the synthesis of TBTQ derivatives 5–7 with three urea units located at the molecular periphery (Figure 1). Owing to the rigidity of their molecular framework, these compounds were considered of potential interest for the formation of supramolecular aggregates consisting of two or more multiply hydrogen-bonded TBTQ units. Research on the formation and chemical properties of host–guest compounds bearing urea units as linker groups has become an exciting field of scientific interest over the past few decades [8-15]. For example, cone-like building blocks such as those based on calix[4]arenes and calix[6]arenes bearing urea groups at their lower rims have been used for the generation of multi-hydrogen bonded dimers [8,9]. Therefore, the TBTQ framework, with its close similarity to that of the cyclotribenzylenes and the cyclotrimeratrylenes [16-19], should constitute a promising substrate for such studies. In the present paper, we wish to report the facile access to TBTQs that bear three phenylurea groups directed into the 3-D space in well controlled orientations. Some observations that point to non-covalent aggregation of these novel molecules are also reported.
Results and Discussion

As shown earlier, introduction of six amino groups at the periphery of the TBTQ skeleton is achievable by nitration with nitric acid (100%) and sulfuric acid (98%) followed by reduction [20]. However, this holds true only for TBTQ derivatives with alkyl groups at the benzhydryl bridgehead positions, such as the tetramethyl derivative 3 [20] and the related tripropyl analog 4 [5,7]. The monomethyl derivative 2 decomposes under these conditions. By contrast, threefold nitration of the arene periphery, leading to single functionalization at the outer positions of each of the three benzene rings, was achieved by use of sodium nitrate in trifluoroacetic acid [21]. As shown in Scheme 1, this method allowed us to convert the tetramethyl derivate 3 into a mixture of the C3-symmetrical compound 8 and the C1-symmetrical isomer 9 in apparently quantitative yield. More recently, this method was also successfully applied to the analogous nitration of compound 2 [22].

Our attempts to separate the two constitutional isomers 8 and 9 by gravity column chromatography failed due to their almost identical elution behaviour. However, 1H NMR spectroscopy of the mixture allowed us to determine the ratio of the isomers in the crude reaction mixture. As expected, the symmetrical compound 8 was the minor component and the ratio observed, 8:9 ≈ 1:3, suggested a random attack of the electrophile at each of the benzene rings even in the second and third step of the threefold nitration. This corresponds to comparable findings with multiple electrophilic substitutions of the centropolyindanes and reflects the lack of electronic interaction between the aromatic units [22].

Because of the prototypical character of three-fold nitration of a TBTQ derivative, the ability to distinguish between the C3- and the C1-symmetrical isomers 8 and 9 and to assess the relative amounts is addressed here in some detail (Figure 2). Notwithstanding the fact that the 1H NMR spectra of compounds 8 and 9 are very similar, their isomer ratio can be determined from the mixture by assuming incremental deshielding effects caused by the nitro group placed at the benzene ring which is adjacent to that bearing the proton under observation. Thus, besides the strong deshielding effect due to the ortho-NO2 groups of the very same benzene ring, the “distal” or “proximal” nitro group at the closest neighbouring ring affects the chemical shift. For example, the resonance of the (“isolated”) ortho-proton 1-H of...
isomer 8 (H^a, δ 8.19) is also affected to a minor extent by the distal 10-NO₂ group. The equivalent protons 5-H^a and 9-H^a of 8 suffer the same extra chemical shift but, interestingly, the isolated proton 5-H^a of isomer 9 is also isochronous and contributes to the doublet (\(J = 2.0\) Hz) at δ 8.19. By contrast, the remaining isolated ortho-protons, 1-H and 12-H, of isomer 9 resonate at slightly lower field (H^b, δ 8.24) due to the stronger deshielding effect of the proximal 11-NO₂ and 2-NO₂ groups, respectively. Correspondingly, this significant difference in the chemical shifts (\(Δδ = 0.05\) ppm) of protons H^a and H^a' of isomers 8 and 9 is found to occur inversely with the resonances of the non-isolated ortho-protons H^a'' and H^b'', and appear as doublets (\(J = 8.5\) Hz) at δ 7.57 and δ 7.50, respectively. Thus, the former resonance is assigned to the equivalent protons 4-H^b', 9-H^a'' and 12-H^b'' of 8 and the isochronous proton 4-H^b'' of 9, all having one proximal nitro group at the adjacent benzene ring. The latter resonance assigned to the isochronous protons 8-H^a'' and 9-H^a'' is slightly upfield-shifted (\(Δδ = 0.07\) ppm) due to the weaker deshielding effect of the respective distal 11-NO₂ and 6-NO₂ groups. Similar arguments hold true for the meta-protons H^b and H^b' of the TBTQ skeleton, with a noticeably slight perturbation in the case of H^b due to the asymmetry of 9. In fact, the unity integral ratios observed for the six different resonances a and a', b and b', and c and c' are in excellent agreement with the assumed isomer ratio 8:9 = 25:75: With this ratio, the integrals of the groups of isochronous protons, e.g., H^a' in 8 and 9 and H^a'' in 9, become equal: [H^a'] = 0.25 × 3 + 0.75 × 1 = [H^a''] = 0.75 × 2 = 1.50. Finally, it may be noted that, accordingly, the 1:3 mixture of the trinitro compounds 8 and 9 is also reflected by three ¹³C NMR lines at δ 25.54, 25.46 and 25.40 in the intensity ratio of ca. 3:6:3.

Reduction of the mixture of the trinitro compounds 8 and 9 was in the first instance achieved by use of hydrazine with iron(III) chloride and charcoal in refluxing methanol (Scheme 1) [23], and afforded a mixture of the triamino-TBTQs 10 and 11 in good yield (81%). However, catalytic hydrogenolysis of compounds 8 and 9 at ambient temperature under medium pressure [20,24,25] proved to be even more efficient and gave the TBTQ-triaminelines 10 and 11 in an even higher yield (91%). At variance from the trinitrotrobenzotriquinacenes, these derivatives were found to display quite different elution behaviour on silica gel. By use of ethyl acetate as an eluent, the retention factors of 10 and 11 were found to be 0.44 and 0.27. After sep-
Figure 2: Partial $^1$H NMR spectrum (500 MHz, CDCl$_3$) of the mixture of the trinitro-TBTQ isomers 8 and 9.

After column chromatography, the first-eluting, C$_3$-symmetrical isomer 10 was obtained in 20% and the second-eluting, C$_1$-symmetrical isomer 11 in 70% yields, in line with the isomer ratio deduced for the trinitro precursors.

Similar to the trinitro compounds 8 and 9, the triamines 10 and 11 were found to give apparently identical EI mass spectra. In all of these cases, loss of a methyl radical is, by far, the dominating primary fragmentation process. However, $^1$H and $^{13}$C NMR spectroscopy allowed us to unequivocally assign the isomers. The C$_3$-symmetry of compound 10 is reflected by the relatively well resolved doublets at δ 6.47 and 7.06 and the relatively sharp singlet at δ 6.59, all of which having unity integral ratios. As expected, the chemical shifts in the spectrum of the C$_1$-symmetrical isomer 11 are very similar to those of 10 but very slight differences of the chemical shifts are evident from the significantly broadened and more complex signals. Notably, these differences are much less pronounced as was the case with the trinitro precursors; thus, the splitting of the resonances of the isolated protons 1-H, 5-H and 12-H of isomer 11 into two singlets at δ 6.59 (1H) and δ 6.61 (2H) is minute. Moreover, and quite specifically, the proton-decoupled $^{13}$C NMR spectrum of the C$_3$-symmetrical isomer 10 exhibits a single line at δ 61.7 due to the three benzhydryllic bridgeheads and another line...
at $\delta$ 25.9 due to the three methyl groups attached to these positions, whereas the spectrum of 11 shows two sets of three lines at $\delta$ 61.1, 61.7 and 62.2 and at $\delta$ 25.7, 25.9 and 26.1, respectively.

The triamino-substituted TBTQs 10 and 11 readily added three equivalents of phenylisocyanate to give the corresponding $C_3$- and $C_1$-symmetrical TBTQ-based tris-ureas 5 and 6 in very good yields (Scheme 2). These conversions were as efficient as the corresponding reactions of the tetraaminocalix[4]arenes [10]. As expected, the colorless amorphous solids had good solubility in polar solvents but not in non-polar ones. Whereas EI and DEI mass spectrometry failed, ionization by FAB(+) furnished almost identical mass spectra that clearly showed the expected [M + H]$^+$ peaks at m/z 739 along with very minor signals for the twofold condensation product. However, the $^1$H NMR spectra of compounds 5 and 6 gave no hint to such impurities.

In analogy to the triamino precursors 10 and 11, the TBTQ-tris-ureas 5 and 6 gave very similar $^1$H NMR spectra even at 600 MHz but, again, the spectrum of the non-symmetrical 6 isomer revealed significant splitting of the peaks. The spectrum of the $C_3$-symmetrical 5 isomer exhibited two characteristic singlets at $\delta$ 8.54 and $\delta$ 8.60 for the two sets of amido protons. $^1$H,$^1$H-COSY measurements allowed us to assign these resonances to the “outer” (Ph–NHCO) and the “inner” (TBTQ–NHCO) amido protons, respectively, of the three equivalent urea groups. In fact, the singlet for the adjacent protons, 1-H, 5-H and 9-H ($\delta$ 7.55), at the TBTQ nucleus, as well as the doublet for the six ortho-protons of the phenyl groups ($\delta$ 7.43), were found to be useful for this assignment. The three equivalent bridgehead methyl groups of 5 resonate at $\delta$ 1.57 and those of the central methyl group appear at $\delta$ 1.30.

Although the bridgehead chemistry of the tribenzotriquinacenes has been extensively investigated [1-4,7], the attachment of multifunctional groupings at the convex surface of the TBTQ skeleton has been only scarcely examined [4]. Therefore, we subjected the previously described bridgehead triaminotri-benzotriquinacene 12 [4] to the reaction with phenylisocyanate in analogy to the conversion of the peripheral triamines 10 and 11, and isolated the corresponding $C_{3v}$-symmetrical TBTQ tris-urea 7 in very good yield (Scheme 2). Again, characterization by FAB(+) and also by ESI(+) mass spectrometry was straightforward; intense [M + H]$^+$ (m/z 697) and, respectively,

$^1$H NMR spectra of compounds 5 and 6 gave no hint to such impurities.

![Scheme 2](image-url)
[M + Na]+ (m/z 719) peaks were in accord with the product of threefold addition. Interestingly, the 1H NMR spectra of compound 7 showed pronounced solvent effects on the NH proton resonances. The outer protons were found to resonate at δ 8.86 in DMSO-d6 but at δ 6.80 in C2D2Cl4, and the inner protons to resonate at δ 6.57 in DMSO-d6 and at δ 5.37 in C2D2Cl4. Thus, the change from the polar to the low-polarity solvent gives rise to a high-field shifts of Δδ = 2.3 and 1.4 ppm, respectively. This effect was attributed to the preferred formation of hydrogen-bound dimers of the tris-urea 7 in tetrachloroethane solution.

As outlined above, the potential dimerization of the threefold urea-functionalized TBTQ derivatives 5, 6 and 7 was one motif for this work. In fact, some evidence for the formation of neutral dimers in low-polarity solvents was found. As mentioned above, the strong high-field shift observed in dichloroethane solution for the NH protons of the bridgehead TBTQ tris-urea 7 suggests the formation of a dimeric aggregate. Specifically, the particularly strong shielding of the external NH protons (Δδ = −2.3) may point to the formation of face-to-face dimers in which the arene units of the opposite TBTQ core exert pronounced magnetic anisotropy effects.

In the case of the peripheral TBTQ tris-ureas 5 and 6, 1H NMR measurements in DMSO and tetrachloroethane revealed a surprising effect which may also be attributed to aggregation of these molecules in low-polarity solvents. Both of these compounds were found to be only poorly soluble in C2D2Cl4 and their 1H NMR spectra in this solvent showed very broad signals for the NH protons. Therefore, assignment of NH and arene proton resonances was impossible even at elevated temperatures (≤ 120 °C). Nevertheless, the 1H resonances of the three benzhydrylic and the single central methyl groups were clearly detectable in both DMSO-d6 and C2D2Cl4. However, in the latter solvent, the protons of the benzhydrylic methyl groups of 5 and 6 were found to resonate at significantly higher fields than those of the central methyl group (Table 1). This is in stark contrast to the relative chemical shifts found with DMSO-d6 as a solvent and with all TBTQ derivatives studied so far.

| Compound | Bridgehead methyl groups | Solvent DMSO-d6 | C2D2Cl4 |
|----------|--------------------------|----------------|----------|
| 5        | 12d-CH3                  | 1.30           | 1.34     |
|          | 4b-, 8b-, 12b-CH3        | 1.57           | 1.17     |
| 6        | 12d-CH3                  | 1.30           | 1.26     |
|          | 4b-, 8b-, 12b-CH3        | 1.56           | 1.07     |

It is noteworthy that the 1H resonance of the central methyl group is hardly affected by the solvent, whereas the resonance of the outer bridgehead methyl groups is shifted to a higher field by Δδ ≈ 0.4 ppm for the C3-symmetrical compound 5 and by even Δδ ≈ 0.5 ppm for the C1-symmetrical compound 6. This difference may be due to the different molecular symmetry. The FAB(+) mass spectrum of the bridgehead tris-urea 7 (obtained from pure methanol solution) showed abundant peaks for (singly charged) [M + Na]+ (m/z 784) and [2 M + 2 Na]+ ions. This difference may be due to the different molecular symmetry. The FAB(+) mass spectrum of the bridgehead tris-urea 7 (obtained with NBA as a matrix) and the corresponding ESI(+) mass spectrum obtained from pure methanol exhibited the [M + H]+ (m/z 697) and the [M + Na]+ (m/z 719) peaks, respectively, but only very minor dimeric quasi-molecular ion peaks.

No more detailed analysis of the dimerization of the new TBTQ-based tris-urea 5, 6, and 7 has been carried out so far. Also, it remains open whether homo- or heterochiral association is preferred with the peripheral tris-urea 5 and 6 and whether the C3-symmetrical isomer 6 behaves differently from the C1-symmetrical isomer 6. Nevertheless, the observations reported in the present work point to the existence of TBTQ tris-urea dimers in non-polar solvents. Congeners bearing three longer-chain aliphatic groups in place of the phenylcarbamoyl residues appear to be good candidates for further studies in this field.

Experimental

2,6,10-Trinitro-4b,8b,12b,12d-tetramethyl-4b,8b,12b,12d-tetrahydrodibenzo-[2,3;4,5]pentaleno-[1,6-ab]indene (8) and 2,6,11-trinitro-4b,8b,12b,12d-tetramethyl-4b,8b,12b,12d-tetrahydrodibenzo[2,3;4,5]-pentaleno-[1,6-ab]indene (9) (mixture of isomers). A suspension of tetramethyltribenzo(triquinacene 3 (250 mg, 743 mmol) in trifluoroacetic acid
(40 mL) was stirred while sodium nitrate (756 mg, 8.92 mmol) was added in small portions. The reaction mixture became a yellow solution after several hours and stirring was continued for a total of 48 h at ambient temperature. After the addition of water (25 mL), the mixture was cooled with ice and the pH was adjusted to ≈ 10 by the careful addition of sodium hydroxide (solid or 6 M aqueous solution). Repeated extraction with chloroform, drying of the combined extracts with sodium sulfate and removal of the solvent under reduced pressure followed by column chromatography (silica gel/CHCl3) gave a mixture of the trinitro compounds 8 and 9 (350 mg, 100%) in a ratio of ca. 1:3 as a yellow, amorphous solid; Rf (CHCl3) 0.74; mp >360 °C; IR (KBr): ν = 3073, 2974, 2934, 1594, 1530, 1475, 1453, 1396, 1345, 1207, 1153, 1112, 1097, 1067, 1032, 907, 887, 825, 742, 674 cm⁻¹; 1H NMR (500 MHz, CDCl3); integral data given as parts of a total of 9 H and 12 H: δ = 8.240 (d, J = 2.1 Hz, 1.5 H), 8.193 (d, J = 2.1 Hz, 1.5 H), 8.125 (d, J = 8.5 Hz, with fine splitting of J ≈ 1.9 Hz, 1.5 H), 8.105 (br d, J ≈ 8.5 Hz, 1.5 H), 7.575 (d, J = 8.5 Hz, 1.5 H), 7.506 (d, J = 8.5 Hz, 1.5 H), 1.77 (s, 2.25 H), 1.75 (s, 4.5 H), 1.73 (s, 2.25 H), 1.43 (s, 3H); 13C NMR (75 MHz, acetone-d6): δ = 155.9, 155.6, 150.8, 150.7, 149.9 (all C), 125.6, 124.8, 124.7, 119.7 (all CH), 72.3, 64.1, 63.94, 63.86 (all C), 25.54, 25.46, 16.1 (all CH3); MS (EI, 70 eV) m/z (%): 471 (9 [M⁺]), 456 (100), 410 (9), 395 (7); accurate mass (EI, 70 eV): m/z: 456 ([M – CH3]⁺), C35H18N2O6, calc 456.1196; found 456.1196.

2,6,10-Tris[(N-phenylcarbamoyl)amino]-4b,8b,12b,12d-tetramethyl-4b,8b,12b,12d-tetrahydrodibenzo[2,3;4,5]pentalenol[1,6-ab]indene (10) and 2,6,11-triamino-4b,8b,12b,12d-tetramethyl-4b,8b,12b,12d-tetrahydrodibenzo[2,3;4,5]pentalenol[1,6-ab]indene (11). Procedure A. A suspension containing the mixture of trinitromtrobenzimidaziniquinacenes 8 and 9 described above (230 mg, 488 mmol), charcoal (32 mg) and iron(III) chloride hexahydrate (10 mg) in methanol (10 mL) was heated to reflux for 15 min and then hydrazine hydrate (97%, 120 mg) added. After heating for 4 h further FeCl3•6H2O (10 mg) and 120 mg Na2H4•H2O (120 mg) were added. The mixture was heated under reflux for a total of 24 h, allowed to cool and filtered through silica with ethyl acetate. Removal of the solvent gave a mixture of the triamines 10 and 11 (151 mg, 81%) as a yellow solid, the components of which can be separated by chromatography. Procedure B. The mixture of isolates 8 and 9 described above (500 mg, 1.06 mmol) and Pd/C (10%, 100 mg) were suspended in ethanol (100 mL) and shaken under a hydrogen atmosphere at 5 bar at ambient temperature for 24 h. The catalyst was removed by filtration through silica gel and the solvent removed under reduced pressure. The triamines 10 and 11 were separated by column chromatography (silica gel, EtOAc) to yield the C3-symmetrical isomer 10 (115 mg, 20%) and the C1-symmetrical isomer 11 (399 mg, 70%).
ν 7 give the tris-urea form (70 mL) at 50 °C for 6 h and then treated as above to μ (125 mol) and phenylisocyanate similar to the procedures given above, triaminotribenzotriquinacene 11. The mixture was heated at 50 °C for 6 h and then treated as above to yield the tris-urea 6 (173 mg, 86%) as a colorless, amorphous solid, mp 257 °C (decomp.; IR (KBr): ν = 3381, 2968, 2928, 1664, 1598, 1549, 1491, 1443, 1415, 1372, 1311, 1227, 1079, 752, 692 cm⁻¹.1H NMR (500 MHz, DMSO-d₆), see Supporting Information File 1 for 600 MHz spectrum: δ = 8.62 (s, 1H), 8.60 (s, 2H), 8.57 (s, 1H), 8.56 (s, 1H), 8.54 (s, 1H), 7.53 (d, J = 1.4 Hz, 1H), 7.49 (d, J = 1.3 Hz, 1H), 7.46 (d, J = 1.4 Hz, 1H), 7.42 (d, J = 7.9 Hz, 6H), 7.38 (d, J = 8.4 Hz, 1H), 7.37 (d, J = 8.3 Hz, 1H), 7.25 (m, 9H), 7.16 (dd, J = 8.3 Hz, J = 1.5 Hz, 1H). 1H NMR (500 MHz, C₂D₂Cl₂): δ = 7.32 (br s), 7.22 (br s), 7.20 (br s), 7.19 (br s), 7.03 (br s), 7.00 (br s), 6.91 (br s), 1.26 (br s, 3H), 1.07 (br s, 9H); MS [FAB(+), NBE] m/z (%): 1478 (2) [2 M + H⁺], 739 (73) [M + H⁺], 723 (7); MS [ESI(+), MeOH] m/z (%): 1500.6 (19) [2 M + Na⁺], 1136.4 (4), 767.3 (9), 761.3 (100) [M + Na⁺], 756.3 (28), 739.3 (12), 670.3 (6), 620.3 (7); MS [ESI(+), MeCO₂H] m/z (%): 1500.7 (9) [2 M + Na⁺], 761.4 (100) [M + Na⁺]; MS [ESI(+), MeOH, (CH₃)₂NPF₆] m/z (%): 1518.5 (5) [2 M + (CH₃)₂N⁺], 1499.6 (5), 848.6 (17), 812.4 (70) [M + (CH₃)₂N⁺], 761.3 (76), 756.3 (100), 739.3 (54), 663.4 (20), 620.3 (27), 529.4 (30); combustion analysis: C₄H₂H₂N₂O₂H₂O₂ calecd C 74.58, H 5.86, N 11.07, found C 74.51, H 5.92, N 10.98.

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Supporting Information
Supporting Information File 1
1H NMR spectra including magnifications of compounds 5, 6 and 7 (600 MHz) and of compounds 8, 9, 10 and 11 (500 MHz).

[http://www.beilstein-journals.org/bojc/content/supplementary/1860-5397-7-43-S1.pdf]

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