Molecular solubilization of fullerene C\textsubscript{60} in water by \(\gamma\)-cyclodextrin thioethers

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Abstract
Various hydrophilic \(\gamma\)-cyclodextrin (CD) thioethers, containing neutral or ionic side arms were found to form molecular disperse solutions of C\textsubscript{60} in water reaching concentrations of 15 mg/L. Equilibrium state was approached after seven days without the use of organic cosolvents. The 1:2 stoichiometry of the C\textsubscript{60}/\(\gamma\)-CD thioether complexes was demonstrated by a parabolic phase-solubility diagram. In contrast, native \(\gamma\)-CD forms nanoparticles with C\textsubscript{60}. Particle sizes of C\textsubscript{60} were determined by dynamic light scattering.

Introduction
Since the first spectroscopic discovery of buckminsterfullerene, C\textsubscript{60}, by Kroto, Heath, Curl and Smalley in 1985 [1], and its first macroscopic synthesis by Krätschmer et al. in 1990 [2], this third allotropic modification of carbon has been the subject of more than 10,000 publications giving rise to many interesting potential applications both in the biomedical field [3-5] and materials science [6-9].

A good solubility of C\textsubscript{60} in water is especially required for biological applications; however, this is not the case at all. Solubility of C\textsubscript{60} was estimated to be as low as \(10^{-8}\) ng/L, equivalent to 10 C\textsubscript{60} molecules per millilitre of water [10,11]. Therefore, hydrophilic derivatives of C\textsubscript{60} have been synthesized and utilized for the inhibition of therapeutically important enzymes, such as HIV-1 protease [12], for the prevention of bacterial growth [13,14], or for photodynamic therapy of cancer by scission of DNA [3]. Despite these successes, there are still several issues relating to the chemical modification of C\textsubscript{60}. The regioselectivity of derivatization is difficult to control [15], and derivatization reduces aromaticity, which leads to a change of the distinct electronic and photonic properties of C\textsubscript{60}.

Dispersions of C\textsubscript{60} nanoparticles in water have been discussed as alternatives for molecular solutions. Such C\textsubscript{60} dispersions are
generally obtained by the so-called solvent-exchange method [16,17], where $C_{60}$ is dissolved primarily in an organic solvent, such as benzene, THF, or acetone, and afterwards diluted with water. After evaporation of the organic solvent, clusters of $C_{60}$ in water remain, which are temporarily stable. The biological activities of those dispersions strongly increase with decreasing size of the $C_{60}$ nanoparticles [13]. Also micellar dispersions of $C_{60}$ in water stabilized by detergents such as Tween, Triton, or SDS are known [18]. Applications of these $C_{60}$ dispersions are hampered by the toxicities of the employed organic solvents or surfactants.

The most successful strategy to carry the extremely hydrophobic $C_{60}$ molecule into water is the use of appropriate water-soluble carriers that can form host–guest complexes, such as calixarenes [19,20] and cyclodextrins (CDs) [21,22]. CDs are cyclic oligosaccharides consisting of six, seven, eight or more glucose subunits connected through $\alpha$-, $\beta$-, and $\gamma$-CD, respectively. CD molecules resemble truncated cones comprising a hydrophilic outer surface and a relatively hydrophobic cavity [23-25]. CDs form water-soluble inclusion complexes with many hydrophobic or amphiphilic guest molecules [26], mainly driven by hydrophobic interactions [27]. Among the commercially available CDs, $\gamma$-CD with a clear width of $d = 0.74$ nm [28] is only large enough to partially accommodate $C_{60}$, which has a still greater van der Waals diameter of 1.0 nm [2]. Molecular dynamics studies strongly favour a sandwich-like structure of the complex, in which two $\gamma$-CD molecules tightly interact through hydrogen bonds between their secondary rims and in which $C_{60}$ is situated in the middle at the widest sites of both CDs (see Figure 1, [29]).

Andersson et al. reported that $C_{60}$ formed a water-soluble 1:2 inclusion complex with $\gamma$-CD after heating under reflux in water [21]. Mixing of an aqueous solution of $\gamma$-CD with a methanolic solution of $C_{60}$ led to a $C_{60}$ dispersion with a concentration of ca. 70 mg/L [30]. Even higher concentrations of up to 1 g/L were reached by high-speed vibration milling of $C_{60}$ in aqueous solutions of $\gamma$-CD [31]. The main drawbacks of these aqueous systems are still (a) their lack of stability, leading to crystallization of the $C_{60}$ complex after some days [31]; and (b) their pronounced tendency to form nanoparticular aggregates [21,32], both of which limit their practical application.

Recently, we developed a new class of highly water-soluble per-6-deoxy-thioethers of $\beta$- and $\gamma$-CD (>20% w/w), which showed exceptionally high solubilization abilities for several aromatic molecules, such as anthracene and acenaphthylene, in water [33-35]. In this work, we investigated solubilization of $C_{60}$ by these $\gamma$-CD thioethers (compounds 1–7 in Scheme 1) with the hope of achieving high concentrations of solubilized $C_{60}$. The hydrophilic substituents at the primary face of $\gamma$-CD should increase solubility and avoid aggregation, because they cannot form intermolecular hydrogen bonds like the primary hydroxyls of native $\gamma$-CD.

**Results and Discussion**

**UV–vis spectra of $C_{60}$ solutions**

Powdered $C_{60}$ was stirred at 25 °C in 6 mM aqueous solutions of $\gamma$-CD and $\gamma$-CD thioethers 1–7 giving rise to clear, dark yellow solutions of $C_{60}$, which show a narrow absorption band at $\lambda_{\text{max}} = 335$ nm quite similar to that of a solution of $C_{60}$ in THF ($\lambda_{\text{max}} = 327$ nm), shown in Figure 2. Extensive centrifugation caused only a small reduction of signal intensity (less than
Dissolution kinetics
After the dissolution of C₆₀ by γ-CD thioethers had been demonstrated, we were interested in how long it takes to reach equilibrium, within experimental error. Therefore a thinly casted film of C₆₀ was incubated with an aqueous solution of γ-CD thioether 7 at 50 °C and stirred according to Kuroda et al. [37]. The slow increase of C₆₀ concentration was monitored by the increase in absorption intensity at 335 nm. The observed first-order dissolution kinetics of C₆₀ came nearly to an end after 7 d, as shown in Figure 3. The obtained rate constant, \( k = 0.021 \, \text{h}^{-1} \) was somewhat higher than the one already found for native γ-CD, \( k = 0.011 \, \text{h}^{-1} \), which also showed first-order dissolution kinetics [37].

Dissolution of C₆₀ can be described by a two-step process, originally proposed by Kuroda et al. [37]. Alternatively, a one-step process, in which a C₆₀ molecule is trapped by two CD rings at the same time, appeared reasonable to us. Both models are shown in Scheme 2.

The observed simple first-order kinetics was puzzling for us, because the concentration of the CD host does not go down significantly (<<1%) over the course of the dissolution of C₆₀. Therefore other reasons for the observed continuous decrease of dissolution rate had to be found.

The two-step model (a) employs a slow, rate-determining complexation of the guest C₆₀ by the first CD molecule and a fast further complexation by the second. The second step is much faster than the first due to the strong stabilization exerted by multiple hydrogen bonds between both CD rings. For the back process, the first dissociation step should be slow and rate determining, because of the necessary cleavage of these intermolecu-

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8%, Figure 2), which clearly precludes the existence of aggregates, which would otherwise sediment to the bottom. Therefore the molar concentration of C₆₀ could be calculated by using the known molar extinction coefficient of C₆₀ molecularly dissolved in n-hexane at 328 nm, i.e., \( \varepsilon = 52,000 \, \text{M}^{-1} \cdot \text{cm}^{-1} \) as published previously [36].

**Figure 2**: UV–vis spectra of (a) C₆₀ solution in THF, (b) aqueous solutions of C₆₀ with 6 mM γ-CD thioether 5 before centrifugation, and (c) after centrifugation (13,000 rpm) for 60 min. The absorbance intensities of (a) and (b) are shifted by 0.2 and 0.1 AU, respectively.

**Figure 3**: Isothermal kinetics of the dissolution of C₆₀ in the presence of 10 mM CD 7 in water. Curve: best fit of first-order kinetics \( k = 0.5 \, \text{d}^{-1} = 0.021 \, \text{h}^{-1} \).
ular hydrogen bonds. These assumptions lead to an apparent equilibrium constant $K'$ in which the CD concentration is only present in first order. The initial rate of complexation is predicted to be proportional to CD concentration, which was already experimentally found for native $\gamma$-CD [37].

On the other hand, the one step model (b) leads to the classical binding constant $K$. The initial complexation rate should be proportional to the square of the host concentration, as shown in Supporting Information File 1.

Both models have in common that the formation of the complex follows pseudo-zero-order kinetics whereas dissociation follows first-order kinetics. Consequently, the integrations of the rate equations for both models (described in Supporting Information File 1) lead to the same final kinetics (Equation 1), a simple first-order equation converging to the equilibrium solubility of the guest $[C_{60} \cdot CD_2]_{equ}$ in agreement with the observed experimental data. According to both new models, the obtained rate constants $k_2$ and $k_1$, respectively, are not due to formation of the complex as proposed previously [37], but due to its dissociation.

$$[C_{60} \cdot CD_2] = [C_{60} \cdot CD_2]_{equ} \left(1 - \exp(-k_2 t)\right) \quad (1)$$

### Dependence of the equilibrium concentration of $C_{60}$ on the host and its concentration

Equilibrium concentrations of $C_{60}$ in aqueous solutions of 6 mM $\gamma$-CD and $\gamma$-CD thioethers 1–7, determined from the absorptions ($\lambda_{max} = 335$ nm) after being stirred for 7 d, are listed in Table 1. $C_{60}$ concentrations obtained with $\gamma$-CD thioethers were up 35 times higher than the one obtained with native $\gamma$-CD, in accordance with previous results for other hydrophobic guests [38]. The improved solubilization potential of the thioethers was attributed to the higher hydrophobicity of sulfur compared to oxygen. The highest concentration of $C_{60}$ in water, 14.9 $\mu$M, was found for CD derivative 5 with attached neutral diol substituents. In general, neutral $\gamma$-CD thioethers 1, 5, and 7 performed better than the anionic ones 2, 3, 4. Coulomb repulsion between the anionic groups in between the two CD molecules was held responsible for the reduced binding affinity. Astonishingly, the amino derivative 1 also showed a high solubilization potential, which may originate from the addition of the amine to a double bond of $C_{60}$, as was already observed by Geckeler for other amino compounds [39].

The phase-solubility diagram of $C_{60}$ in the presence of $\gamma$-CD thioether 3, according to the method established by Higuchi and Connors [40], was obtained by plotting the concentration of the dissolved $C_{60}$ versus the concentration of the host, as depicted in Figure 4. The observed parabolic $A_P$-type phase concentration dependence is typical for the formation of complexes with 1:2 stoichiometry [40,41]. The equation for the best fit $[C_{60}] = 10^{-3}(0.21 + 0.17[CD] + 0.11[CD]^2)$ indicates that the two-step process (first order in [CD]) as well as the one-step process (second order in [CD]), discussed above, contribute to the dissolution of $C_{60}$.

### Preparation of $C_{60}$ complexes with the aid of organic solvents

Because equilibration took a long time (7 d) and occupancies of the $\gamma$-CD thioethers are still low (<0.3%), organic solvents, such as toluene, DMF or CS$_2$ were added to the aqueous solutions of $\gamma$-CD and $\gamma$-CD thioether 5, in the hope of accelerating and improving the dissolution of $C_{60}$ as reported by Murphy et al. [42]. The resulting solutions were characterized by UV–vis spectroscopy. The saturation concentrations of $C_{60}$ are listed in Table 2.

#### Table 1: $C_{60}$ concentration in 6.0 mM aqueous solutions of $\gamma$-CD and $\gamma$-CD thioethers 1–7.

<table>
<thead>
<tr>
<th>host</th>
<th>$[C_{60}]$ ((\mu)M)</th>
</tr>
</thead>
<tbody>
<tr>
<td>$\gamma$-CD</td>
<td>0.4</td>
</tr>
<tr>
<td>1</td>
<td>10.3</td>
</tr>
<tr>
<td>2</td>
<td>2.9</td>
</tr>
<tr>
<td>3</td>
<td>5.3</td>
</tr>
<tr>
<td>4</td>
<td>2.1</td>
</tr>
<tr>
<td>5</td>
<td>14.9</td>
</tr>
<tr>
<td>6</td>
<td>7.6</td>
</tr>
<tr>
<td>7</td>
<td>9.3</td>
</tr>
</tbody>
</table>

![Figure 4: Phase-solubility diagram of $C_{60}$ in aqueous solution in the presence of CD 3.](image-url)
Table 2: C₆₀ concentration in 6.0 mM aqueous solutions of γ-CD and γ-CD thioether 5.

<table>
<thead>
<tr>
<th>CD</th>
<th>[C₆₀] (μM) procedure a</th>
<th>procedure b</th>
<th>procedure c water/DMF/toluene [42]</th>
<th>procedure d water/CS₂</th>
</tr>
</thead>
<tbody>
<tr>
<td>γ-CD</td>
<td>0.4</td>
<td>7.5</td>
<td>4.9</td>
<td>10.8</td>
</tr>
<tr>
<td>5</td>
<td>14.9</td>
<td>21.4</td>
<td>6.3</td>
<td>18.6</td>
</tr>
</tbody>
</table>

Indeed, for native γ-CD much higher C₆₀ concentrations could be achieved in water with the aid of organic solvents. But careful examination of the UV–vis spectra of the C₆₀ solutions produced according to procedure c [42] before filtration, after filtration, and after centrifugation, as shown in Figure 5, revealed significant differences. The unfiltered solution of the C₆₀/γ-CD complex showed an additional absorption maximum 475 nm, which is typical for C₆₀ nanoparticles (nC₆₀) such as those formed by dilution of a solution of C₆₀ in THF with water [16,43]. After centrifugation nearly no C₆₀ was left. Consequently, apparent improvements of the solubilization obtained with the aid of organic solvents were mostly due to the formation of nanoparticle dispersions. Since these organic solvents are also hazardous for most cells, applications in biomedicine are prohibited. Therefore, organic solvents were avoided for the dissolution of C₆₀ by γ-CD thioethers, because they also did not improve the solubilization process significantly.

Investigation of aggregation by dynamic light scattering
Since aggregation of CDs and CD inclusion compounds was already found in previous work [44,45], dynamic light scattering (DLS) investigations were performed to check for any aggregation during solubilization of C₆₀ by γ-CD thioethers. DLS is a relatively fast method for the determination of the particle size distributions of proteins, polymers, micelles, and nanoparticles [46]. In particular, DLS is able to distinguish between a homogenous molecular solution and a dispersion of aggregates [47,48]. The size distribution of the solution of C₆₀ in 6 mM γ-CD thioether 5 before centrifugation, shown in Figure 6, comprised two peaks at particle sizes of 3 and 300 nm. The first peak was attributed to the molecular CD/C₆₀ complex, the second to aggregates of it. Since the intensity of the scattered light increases with the sixth power of the particle size, the content of aggregates is highly overestimated [45]. For getting the right picture, this intensity profile had to be transformed to the volume distribution profile, by using the Mie theory [49]. In the resulting volume distribution profile (Figure 6b) only one peak remains, which corresponds to the molecular complex. Consequently, this solution mainly consists of molecularly dissolved C₆₀. This finding is in accordance with the negligible decrease of C₆₀ absorption caused by centrifugation, shown in Figure 2.

After centrifugation (13,000 rpm) for 60 min, shown in Figure 6c, only a sharp peak at 1 nm was observed for both the intensity and the volume size distributions, which demonstrates the validity of the DLS measurement. For comparison, the corresponding intensity and volume size distributions of a freshly prepared C₆₀ solution in the presence of 6.0 mM native γ-CD according to procedure c [42] (Figure 7) only showed one peak at a diameter of 166 nm, very similar to nC₆₀ prepared from THF. Consequently, the DLS investigations confirmed our previous finding that solubilization of C₆₀ in water with the aid of organic solvents results in dispersions of C₆₀ nanoparticles.

Conclusion
Neutral γ-CD thioethers are especially well suited as solubilizers for C₆₀. Sandwich-like 1:2 complexes are formed at room temperature without the necessity of adding organic cosolvents. These complexes show a much lower aggregation tendency than the corresponding ones of native γ-CD. Molecular solubilization of fullerene C₆₀ in water, reaching concentrations as high...
as 15 μM, was achieved in the presence of 6.0 mM of a γ-CD thioether. The resulting aqueous molecular solutions of C_{60} free of toxic organic solvents will hopefully find interesting applications in biomedicine, such as in photodynamic therapy or HIV-protease inhibition.

**Experimental**

**General**

Unless otherwise stated, all chemicals were used as received. Powdered fullerene C_{60} (> 99%) was purchased from Sigma Aldrich. Teflon syringe filters from Roth, Karlsruhe, Germany (0.45 μm) were used to remove insoluble material before UV–vis spectrophotometric analysis. UV–vis spectra of aqueous samples were performed on a Perkin Elmer Lambda 2 spectrometer (λ: 200–600 nm), by using quartz cells with a 1 cm or 1 mm optical path at 298 K.

**Synthetic procedures**

Hydrophilic thioethers 1–7 at all primary carbon atoms of γ-CD were synthesized from octakis(6-deoxy-6-iodo)-γ-CD by nucleophilic displacement reaction with sulfur nucleophiles by using standard procedures described previously [38].

**Phase-solubility investigations**

Solubility measurements of C_{60} in the presence of γ-CD and γ-CD derivatives in water were carried out according to the method proposed by Higuchi and Connors [40]. In glass vials containing excess amounts of C_{60}, aqueous solutions of γ-CD or γ-CD derivatives with different concentrations were added. The vials were sealed, protected from light, and magnetically stirred at room temperature for seven days. The solid residues were removed by filtration with syringe filter. According to the Lambert–Beer law, the concentrations of C_{60} in pure water and in CDs solutions were determined from UV–vis extinctions at the absorption maxima (log ε = 4.717, λ_{max} = 335 nm) [18].

**Procedures for the solubilization of C_{60}**

Several procedures were employed for the solubilization of C_{60}. **Procedure a**: C_{60} stirred in water in the presence of 6 mM γ-CD thioether at rt for 7 d. **Procedure b** was modified from a previously described method [21]: heated under reflux in water/toluene (ν/ν 1:1) for 3 d (γ-CD thioether concentration 6 mM), and then the resulting mixture dissolved in water after evaporation of the solvents. **Procedure c** was taken from a previous paper [42]: C_{60} stirred in DMF/toluene (ν/ν 1:1) at rt for 7 d.
with 6 mM γ-CD thioether, and then the obtained inclusion complex dissolved in water after evaporation of the solvents. **Procedure d:** C₆₀ stirred in water/CS₂ (v/v 1:1) at rt for 7 d with 6 mM γ-CD thioether, and then the obtained inclusion complex dissolved in water after evaporation of the solvents. nC₆₀ was prepared following a method similar to that reported by Deguchi et al. (**procedure e**) [16]. nC₆₀: A saturated solution of C₆₀ in THF was prepared by adding an excess amount of solid C₆₀ (>2 mg) into THF (20 mL) and stirring overnight under a nitrogen atmosphere at room temperature. Excess solid was filtered off with a syringe filter. Saturated C₆₀/THF solution (500 mL) was placed in a flask and an equal volume of water was added at a rate of ca. 25 mL/min under vigorous stirring. A rotary evaporator was used to remove THF by using a stepwise evaporation approach. The start temperature was set at 30 °C. When the mixture volume had decreased to 500 mL, the temperature was increased at 1 °C/min to 70 °C and maintained at 70 °C until the volume had decreased to 250 mL, at which time an additional 250 mL volume of water was added. The last step was repeated once. The resulting nC₆₀ solution could be diluted by adding water or concentrated by evaporation as needed.

**Isothermal kinetic measurement**

The procedure for performing isothermal kinetic investigations was modified from a previously described method [37]: A solution of C₆₀ in chloroform (ca. 0.5 mg/mL, 0.5 mL) was carefully evaporated in a 1 cm quartz cell under nitrogen flow. After the addition of 3 mL of an aqueous solution containing the necessary amount of γ-CD derivative, the cell was sealed, protected from light, and maintained at 50 °C under gentle shaking. The UV–vis spectra of the resultant solution at appropriate time intervals were measured directly.

**Dynamic light scattering measurement**

Particle size distributions of the aqueous solution or dispersions of C₆₀ were determined by dynamic light scattering (DLS) with a ZetaSizer Nano ZS (Malvern Instruments Ltd., Malvern, United Kingdom). From the diffusion coefficient the radius of the particle was determined via the Stokes–Einstein equation. The samples were filtered through a 0.45 µm syringe filter prior to particle-size measurements. Both intensity and volume size distribution curves were calculated from the scattering data by using the software of the instrument.

**Supporting Information**

Supporting Information File 1

Detailed dissolution kinetics.

[http://www.beilstein-journals.org/bjoc/content/supplementary/1860-5397-8-188-S1.pdf]

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**References**
