Control of stilbene conformation and fluorescence in self-assembled capsules

Mark R. Ams¹, Dariush Ajami¹, Stephen L. Craig², Jye-Shane Yang³ and Julius Rebek Jr*¹

Abstract
The extensively studied trans-stilbene molecule is known to give only weak fluorescence in solution and inside loosely-fitting synthetic capsules. However, trans-stilbene has been recently studied in the context of antibody interiors, where binding results in strong blue fluorescence. The present research was undertaken to understand the spatial factors that influence stilbene fluorescence. trans-Stilbene was encapsulated in the snug, self-assembled complex 1.1 and exhibited fluorescence quenching due to the distortion of its ground-state geometry. When the complex is elongated by incorporating glycouril spacers, trans-stilbene is allowed to adapt a fully coplanar arrangement and fluorescence returns.

Introduction
The fluorescence of trans-stilbene has been extensively researched [1], and weak fluorescence occurs in aqueous solutions or typical organic solvents. In a highly structured environment such as an antibody interior [2-4], recent studies show that nearby tryptophans can transfer electrons to the stilbene excited state and an intense blue fluorescence develops. Inside the tight-fitting capsule 1.1 [5,6] (Figure 1) where it is surrounded by 16 aromatic panels, trans-stilbene’s fluorescence is reduced to only 2% of what is observed in bulk solution. In contrast, normal fluorescence is observed in a loose-fitting capsule [7] although the photostationary trans-/cis-isomerization equilibria are altered in the limited space [8]. Isomerization of trans- to cis-stilbene is not possible in 1.1 but little else is known about the photophysics of guests in this capsule. This research was undertaken to understand what controls the behavior of stilbenes in this and related constrained environments.
Results and Discussion
The space inside capsule 1.1 is defined by two pyramids comprising resorcinarenes at the ends and square prisms of the heterocyclic walls near the middle. The quenching of stilbene fluorescence may be an effect of the fixed aromatics of the resorcinarene or the heterocyclic walls, but we surmised that there was a subtler cause. As seen in the skeletal model 1.1 of the space within depicted in Figure 1A, the square prisms are twisted by about 45° along their long axes. A typical aromatic guest such as benzene fits best when it is nestled diagonally in a prism’s space. Accordingly, the aromatic rings of longer molecules such as biphenyls and stilbenes cannot be coplanar in their lowest energy conformations inside 1.1. Rather, they must be twisted by 45° or so along their rotatable internal bonds.

Among guests of 1.1, 4,4'-dimethylstilbene (2) provides an excellent fit; it fills about 53% of the capsule’s space and the methyl groups of the guest can access the tapered ends of the host. The homologue, 4-ethyl-4'-methylstilbene (3), is also encapsulated (see Supporting Information File 1 for NMR spectrum), but the slightly longer 4,4'-diethyl derivative 4 simply does not fit. Figure 2 illustrates the effect of encapsulation on stilbene 3 (assembly 6). The fluorescence is 96% quenched when $\lambda_{exc} = 318$ nm. For comparison, the emission of the permanently twisted, $o$-substituted stilbene 5 ($\lambda_{exc} = 300$ nm) is also shown [9].

Can the fluorescence of the encapsulated stilbene be restored? When suitable guests are present, addition of glycolurils such as 7 to solutions of 1.1 generates extended capsule 1.7.4,1 (Figure 3) [10]. The glycolurils are arranged in a chiral manner, in either cycloenantiomer of the extended capsule. The glycolurils force the two square prisms of 1.7.4,1 into registry (Figure 3A). That is, the square prisms are now aligned and stilbene as well as related guests may be found either in the fully coplanar arrangement favored by extended resonance stabiliza-
Figure 2: Room temperature fluorescence spectra at $\lambda_{\text{exc}} = 318$ nm for 10 µM mesitylene solutions of 4,4′-dimethylstilbene (2); 4-ethyl-4′-methylstilbene (3); 4,4′-diethylstilbene (4); 2,4,4′,6-tetramethylstilbene (5) and assembly 6 ($\lambda_{\text{exc}} = 300$ nm).

Earlier we showed that it is possible to reversibly interconvert capsules 1.1 and 1.7.4.1. The weakly basic glycoluril spacer is protonated by addition of gaseous HCl and precipitates in typical organic solutions. The remaining components reassemble to the original capsule 1.1 [11]. Subsequent addition of NEt$_3$ releases the glycoluril into solution and restores the extended capsule 1.7.4.1. This was shown previously to be a fully rever-

tion (Figure 3B) or at another minimum with a 90° dihedral angle between their aromatic rings. 4-Ethyl-4′-methylstilbene (3) is a guest for 1.7.4.1 when CD$_2$Cl$_2$ is a co-guest (see Supporting Information File 1 for NMR spectrum). In the fluorescence experiments at $\lambda_{\text{exc}} = 318$ nm, the fluorescence of 4-ethyl-4′-methylstilbene 3 and assembly 8 is restored, as shown in Figure 4. As a control experiment to test for the possibility of any fluorescence contribution from assembly 1.7.4.1, the fluorescence of the complex of 1.7.4.1 with alkyl chain C$_{17}$H$_{36}$ was also investigated. No additional fluorescence was observed when C$_{17}$H$_{36}$ was a guest (see Supporting Information File 1 for fluorescence and NMR spectra).

Figure 3: (Top) Glycouril 7, the extended capsule 1.7.4.1, (only one enantiomeric arrangement is shown) and its cartoon representation. The shape of the space inside (A) a schematic view along the central axis with two aromatic guests (B) and an end-on view of an energy-minimized complex of stilbene inside (C).
Figure 4: Room temperature emission spectra for 10 µM solutions of 4-ethyl-4′-ethylstilbene (3) in the capsule 1.1 (6) and 1.7 (4). λ<sub>exc</sub> = 318 nm.

sible process with long chain alkane guests. We are currently pursuing this application with stilbenes and studying the exchange of subunits in the process [12,13].

Many fluorescent sensors have been reported in the literature [14-17], however they usually respond to chemical changes rather than purely geometrical ones. Here, self-assembly of an external host system is responsible for turning on and off stilbene fluorescence through geometrical control of the stilbene’s surroundings.

Supporting Information

Supporting Information File 1
Control of stilbene conformation and fluorescence in self-assembled capsules.
[http://www.beilstein-journals.org/bjoc/content/supplementary/1860-5397-5-79-S1.pdf]

Acknowledgements
We are grateful to the National Institutes of Health and the Skaggs Institute for support. The authors would also like to recognize the help of Prof. David P. Millar and Dr. Goran Plevačič.

References

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The definitive version of this article is the electronic one which can be found at: doi:10.3762/bjoc.5.79