

Controlled diastereoselectivity at the alkene-geometry through selective encapsulation: *E-Z* photoisomerization of oxazolidinone-functionalized enecarbamates within hydrophobic *nano*-cavities†

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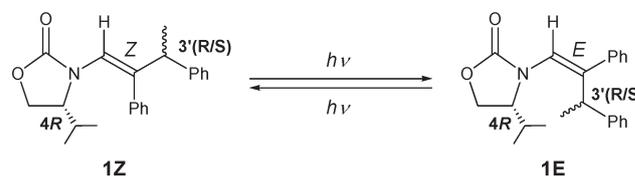
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Photoisomerization of encapsulated *Z*-enecarbamates within the hydrophobic chiral cavities of γ -CD showed higher diastereoselectivities in the photoproducts than those obtained in solution. The selective encapsulation of the enecarbamates and the following isomerization process are both diastereoselectively controlled by γ -CD.

Understanding the intricacies involved in photoisomerization within confined cavities has provided insights into the excited state processes that occur in biological systems.¹ Supramolecular assemblies such as micelles, zeolites and cyclodextrins have been shown to be very effective in controlling the excited state processes during phototransformation.² Immense progress has been made in recent decades to achieve high stereoselection in photoreactions in which the chiral information in the photoproducts is imprinted within the intervening short-lived excited species/reactive intermediate.^{3–6} Recently we have shown that oxazolidinone-functionalized enecarbamates are versatile systems for the study of conformational, electronic, stereoelectric, and steric effects.^{7–11} Enantioselectivity as high as 97% was observed with these systems during photooxygenation. Further, it was clearly demonstrated that the alkene geometry is crucial to control the approach of the singlet oxygen. We have now carried out a systematic study on how to control the photoisomerization process that will provide insights into how the alkene geometry may be fine-tuned in isotropic media.¹² Herein, we report the influence of cyclodextrin nano cavities¹³ on the diastereoselectivity of the photoisomerization process.

Diastereoselective photoisomerization of oxazolidinone-functionalized enecarbamates **1** (Scheme 1) in solution gave low diastereoselectivities (5–15%) upon direct and sensitized irradiation.¹² Even chiral sensitizers did not significantly alter the



Scheme 1 Photoisomerization of 4-isopropylloxazolidinone functionalized 1'-*Z*-4*R*(*Pr*),3'-*R/S*-enecarbamates **1Z**.

observed selectivity in these systems.¹² In this context, we investigated cyclodextrin nanocavities^{13,14} as host to bias stereoselection during the photoisomerization processes. 4-Isopropylloxazolidinone functionalized 1'-*Z*-4*R*(*Pr*),3'-*R/S*-enecarbamates (**1Z**) complexed readily with γ -cyclodextrin (γ -CD). The γ -CD/**1Z** complex was formed by adding **1Z** (0.03 mmol) in 12.5 mL CD₃OD to 12.5 mL D₂O solution of γ -CD (0.03 mmol).¹⁵ FTIR analysis of the γ -CD/**1Z** complex in a 1 : 1 v/v of D₂O/CD₃OD mixture showed the ν (CO) band of the oxazolidinone ring at 1699 cm⁻¹ compared to that of the uncomplexed **1Z** in CD₃OD at 1732 cm⁻¹ (Fig. 1). The observed shift [$\Delta\nu$ (CO) = -33 cm⁻¹] strongly suggests that a host-guest complex is formed between **1Z** in the hydrophobic chiral cavity of the cyclodextrin; the carbonyl moiety of oxazolidinone ring is stabilized through a hydrogen bond with a proton provided by the hydrophobic inner cavity of γ -CD.¹⁶ ¹H-NMR analysis is consistent with the FTIR results, in which an

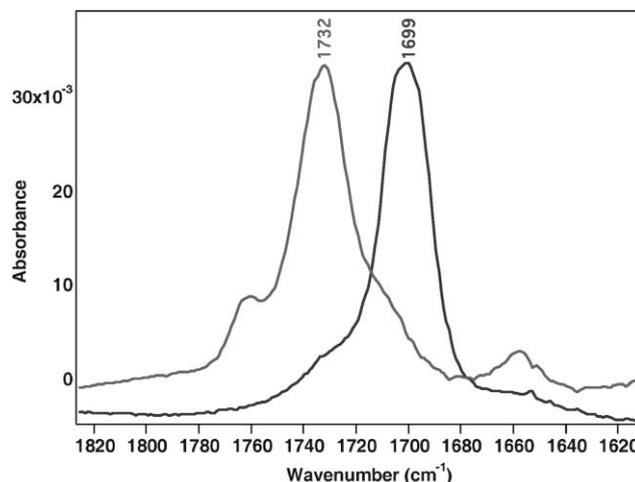


Fig. 1 FT-IR spectra of **1Z** in CD₃OD (left) and the complex of **1Z** with γ -CD (right) in 1 : 1 v/v of D₂O/CD₃OD.

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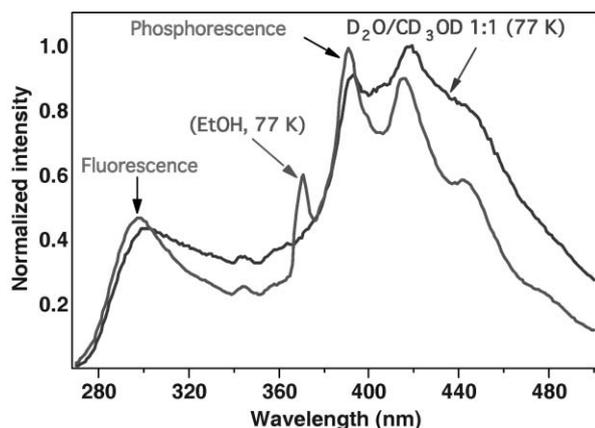


Fig. 2 Steady state emission spectra at 77 K (λ_{ex} : 260 nm) of **1Z** (5.7×10^{-5} M) in ethanol and γ -CD/**1Z** ($[\mathbf{1Z}] = [\gamma\text{-CD}] = 4.8 \times 10^{-5}$ M) in 1 : 1 v/v $\text{D}_2\text{O}/\text{CD}_3\text{OD}$.

upfield shift in the vinylic hydrogen of the enecarbamate was observed.¹⁵

Steady-state emission spectra of the γ -CD/**1Z** complex were recorded in 1 : 1 v/v of $\text{D}_2\text{O}/\text{CD}_3\text{OD}$ at 77 K, and were compared to the steady-state emission spectra of **1Z** in an ethanol glass under identical conditions (Fig. 2). The phosphorescence emission of the γ -CD/**1Z** complex was broadened and slightly redshifted, which indicates that there are different orientations of the guest within the γ -CD cavity of the host. Time-resolved phosphorescence measurements¹⁵ were performed at 77 K by multichannel scaling using light pulses at 260 nm for excitation and monitoring the emission at 390 nm. The γ -CD/**1Z** host-guest complex showed a phosphorescence lifetime of ~ 27 ms compared to 6.6 ms in ethanol glass. It is clear that the increase in the lifetime was caused by encapsulation of **1Z** within the hydrophobic cavity of γ -CD. A similar increase in lifetime upon complexation by γ -CD has previously been reported.¹⁷

Photoisomerization of the γ -CD/**1Z** complex in 1 : 1 v/v of $\text{D}_2\text{O} : \text{CD}_3\text{OD}$ ($[\gamma\text{-CD}] = [\mathbf{1Z}] = 1.4$ mM) was carried out for different time intervals and the observed diastereomeric ratio (dr) in the product *E*-enecarbamates and the starting *Z*-isomer was determined both by ¹H-NMR spectroscopy and by gas chromatography (Table 1).¹⁵ All the photoirradiations were performed on a 50 : 50 mixture of the 3'*R* and 3'*S* epimers for both *Z*- and *E*-enecarbamate, unless noted otherwise. The diastereomeric ratio

Table 1 Photoisomerization of **1Z** with γ -CD in a $\text{D}_2\text{O}/\text{CD}_3\text{OD}$ solution

Irradiation/min ^a	<i>Z</i> : <i>E</i>	dr (1Z) ^b	
		3' <i>R</i> : 3' <i>S</i>	3' <i>R</i> : 3' <i>S</i>
0.5	79 : 21	57 : 43	29 : 71
1	67 : 33	56 : 44	39 : 61
2	51 : 49	53 : 47	42 : 58
3	51 : 49	54 : 46	46 : 54
5	52 : 48	52 : 48	49 : 51

^a $[\mathbf{1Z}] = [\gamma\text{-CD}] = 1.4$ mM. Photoirradiation was performed at ambient temperature in a 1 : 1 v/v of $\text{D}_2\text{O}/\text{CD}_3\text{OD}$ with an excimer laser at 308 nm; 20 Hz, 100 mJ pulse⁻¹ (2W). The diastereomeric mixture (50 : 50) of the starting material was used. ^b Diastereomeric ratio (dr) of the starting *Z*-isomer after photoreaction. Mass balance >95%. ^c Diastereomeric ratio (dr) of the product.

(dr) were enhanced by encapsulation within the hydrophobic cavity, compared to the direct irradiation of **1Z** without γ -CD in MeOH; the latter showed practically no diastereoselectivity.¹² As can be seen from Table 1, high dr were observed for short irradiation (0.5 min) and a decrease upon prolonged irradiation; the dr (favoring the C-3'*S* epimer) of **1E** decreased from 29 : 71 at 0.5 min to 49 : 51 at 5 min, that is probably due to the fast isomerization of the *Z*- to *E*-isomer.^{18–20}

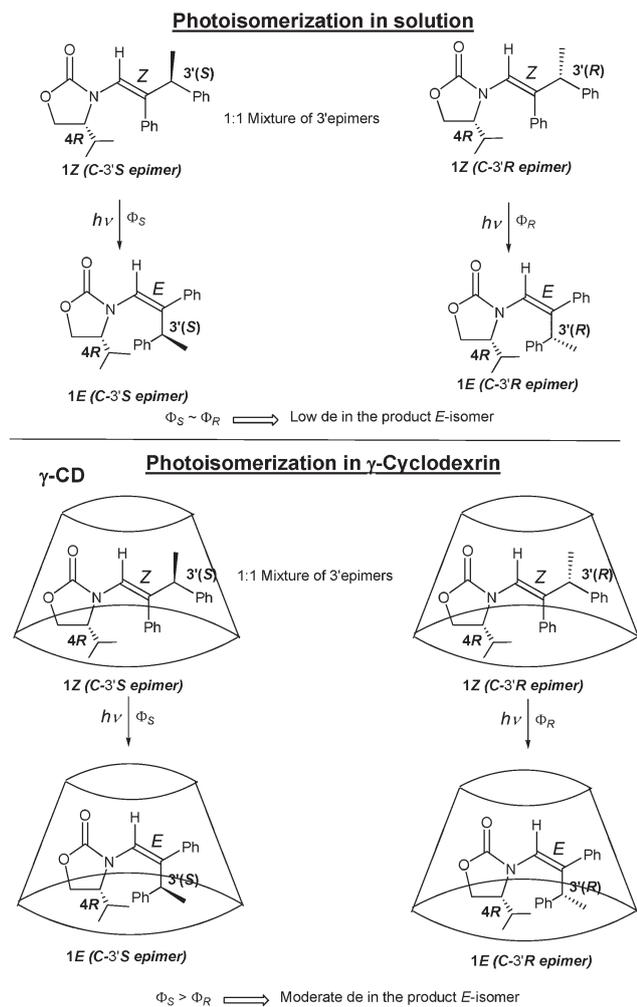
To gain more details on the origin of the selectivity observed within γ -CD, the photoisomerization was investigated in the solid state. The complexation of γ -CD with **1Z** was achieved by adding **1Z** (1.5×10^{-5} mol) in 5 mL ether and γ -CD (1.5×10^{-4} mol) in 10 mL deionized water. A white precipitate of γ -CD/**1Z** complex formed immediately. For easy handling of the sample, excess amount of γ -CD was used during complexation; the ratio of host : guest prepared were 10 : 1 and 20 : 1, respectively. The solid complexes were collected by filtration and washed thoroughly with ether, and the residue was dried in vacuum overnight. Diffuse reflectance Fourier transform IR spectroscopy (DRIFTS) was performed on these samples in a KBr matrix. The carbonyl stretch of the oxazolidinone moiety was observed at 1727 cm^{-1} for both 10 : 1 and 20 : 1 ratio of the γ -CD/**1Z** complexes, compared to 1753 cm^{-1} for the uncomplexed **1Z** (in KBr matrix). The shift [$\Delta\nu(\text{CO}) = -26$ cm^{-1}] of the complex in the solid state was similar to that observed in a $\text{D}_2\text{O}/\text{MeOD}$ solution, which indicates the formation of similar host-guest complexes in both cases.

After the extraction of the *Z*-enecarbamates from the solid complex by dissolving the samples in CD_3CN , ¹H-NMR analysis was carried out to determine the preferential diastereoselectivity of the complexation of **1Z** by γ -CD. Preferential complexation of the 1'*Z*,4*R*(¹Pr),3'*S*-isomer was observed within the hydrophobic chiral cavity compared to 1'*Z*,4*R*(¹Pr),3'*R*-isomer. It should be noted, therefore, that the starting γ -CD/**1Z** complex is preferred for the C-3'*S* epimer, with the dr of (3'*R* : 3'*S*) 42 : 58 and 45 : 55 for the 20 : 1 and 10 : 1 complexes, respectively (Table 2). The solid complexes were then sandwiched between two quartz glass plates and irradiated at ambient temperature. Efficient photoisomerization was observed and the *E*-3'*S* epimer was enhanced to a dr value of 29 : 71 in the solid state (Table 2). The observed selectivity upon photoisomerization in the solid phase indicates the difference in the rate of the photoisomerization of the *Z*-3'*R* and *Z*-3'*S* epimeric enecarbamates, compared to that in CD_3OD solution, for which they have similar photoisomerization rates as

Table 2 Photoisomerization of the γ -CD/**1Z** complex in the solid state^a

Entry	Mixture ratio ($\gamma\text{-CD}:\mathbf{1Z}$)	Irradiation/min	dr (1Z) ^b		
			<i>Z</i> : <i>E</i>	3' <i>R</i> : 3' <i>S</i>	3' <i>R</i> : 3' <i>S</i>
1	20 : 1	0	100 : 0	42 : 58	—
2		15	67 : 33	41 : 59	30 : 70
3		30	60 : 40	40 : 60	32 : 68
4	10 : 1	0	100 : 0	45 : 55	—
5		15	78 : 22	43 : 57	32 : 68
6		30	66 : 34	43 : 57	35 : 65

^a Irradiation was performed at 254 nm at ambient temperature in a Rayonet reactor. The diastereomeric mixture (50 : 50) of the starting material was used. ^b Diastereomeric ratio (dr) of the starting *Z*-isomer after photoreaction and the product *E*-isomer. Mass balance >95%; dr was determined by ¹H-NMR spectroscopy in CD_3CN as solvent.



Scheme 2 Photoisomerization of 4-isopropylloxazolidinone functionalized in solution compared to that within γ -cyclodextrins.

shown by their low dr (Scheme 2-top). The photoisomerization process of *Z*-3'*S* to *E*-3'*S* epimer is clearly enhanced (Scheme 2-bottom) by the complexation compared to that of the *Z*-3'*R* to the *E*-3'*R*, which may be attributed to the less hindered structure of the *E*-3'*S* epimer than the *E*-3'*R* epimer in the confined cavity of γ -CD.

Our current study of diastereoselective photoisomerization of oxazolidinone-functionalized enecarbamates **1Z** within the nano cavities of γ -CD shows that the rate of photoisomerization of the 1'*Z*4*R*(*i*Pr),3'*R* and 1'*Z*4*R*(*i*Pr),3'*S* epimers may be altered by supramolecular assemblies like cyclodextrin. The 1'*Z*4*R*(*i*Pr),3'*S* epimer photoisomerizes faster than the corresponding C-3'*R* epimer upon complexation, which presumably reflects the

conformational effects in the two complexed epimers with γ -CD. This rate difference manifests itself in the observed difference in the isomerization rates that is reflected in the diastereoselectivity of the product *E*-isomer. Our current results, coupled with our photo-oxygenation studies,⁷⁻¹¹ demonstrate that not only the bimolecular singlet-oxygen reaction be controlled, but also the unimolecular photoisomerization process, especially by utilizing chiral confined media such as γ -CD. Evidently, the C-3' position in the enecarbamates is critical in dictating the photoreaction within confined *nano* cavities.⁷⁻¹¹

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Notes and references

- C. Dugave and L. Demange, *Chem. Rev.*, 2003, **103**, 2475; R. S. H. Liu and G. S. Hammond, *Acc. Chem. Res.*, 2005, **38**, 396.
- N. J. Turro, *Proc. Natl. Acad. Sci. U. S. A.*, 2005, **102**, 10766.
- Y. Inoue and V. Ramamurthy, *Chiral Photochemistry*, CRC Press, Boca Raton, FL, 2004.
- A. G. Griesbeck and U. J. Meierhenrich, *Angew. Chem., Int. Ed.*, 2002, **41**, 3147.
- Y. Inoue, *Chem. Rev.*, 1992, **92**, 741.
- J. Sivaguru, A. Natarajan, L. S. Kaanumalle, J. Shailaja, S. Uppili, A. Joy and V. Ramamurthy, *Acc. Chem. Res.*, 2003, **36**, 509.
- W. Adam, S. G. Bosio, N. J. Turro and B. T. Wolff, *J. Org. Chem.*, 2004, **69**, 1704.
- W. Adam, S. G. Bosio and N. J. Turro, *J. Am. Chem. Soc.*, 2002, **124**, 14004.
- W. Adam, S. G. Bosio and N. J. Turro, *J. Am. Chem. Soc.*, 2002, **124**, 8814.
- T. Poon, J. Sivaguru, R. Franz, S. Jockusch, C. Martinez, I. Washington, W. Adam, Y. Inoue and N. J. Turro, *J. Am. Chem. Soc.*, 2004, **126**, 10498.
- J. Sivaguru, T. Poon, R. Franz, S. Jockusch, W. Adam and N. J. Turro, *J. Am. Chem. Soc.*, 2004, **126**, 10816.
- H. Saito, J. Sivaguru, S. Jockusch, Y. Inoue, W. Adam and N. J. Turro, *Chem. Commun.*, 2005, 3424.
- R. Breslow and S. D. Dong, *Chem. Rev.*, 1998, **98**, 1997.
- J. Szejtli, *Chem. Rev.*, 1998, **98**, 1743.
- Refer to the electronic supplementary information.
- K. Tanhuanpää, K. H. Cheng, K. Anttonen, J. A. Virtanen and P. Somerharju, *Biophys. J.*, 2001, **81**, 1501.
- S. Monti, G. Koehler and G. Grabner, *J. Phys. Chem.*, 1993, **97**, 13011.
- B. Giese, *Angew. Chem., Int. Ed. Engl.*, 1977, **16**, 125.
- C. D. Johnson, *Chem. Rev.*, 1975, **75**, 755.
- A. Pross, *Adv. Phys. Org. Chem.*, 1977, **14**, 69.