SUPPORTING INFORMATION

Photochromism of Novel Chromenes Constrained to be Part of [2.2]Paracyclophane: Remarkable 'Phane' Effects on the Colored *o*-Quinonoid Intermediates

Jarugu Narasimha Moorthy,* Susovan Mandal and Amrit Kumar

Department of Chemistry, Indian Institute of Technology, Kanpur 208016, INDIA

moorthy@iitk.ac.in

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compound	СР-Н
molecular formula	$C_{31}H_{26}O$
formula weight	414.52
solvent for crystallization	CHCl ₃
crystal system	Triclinic
<i>a</i> (Å)	7.8416(18)
<i>b</i> (Å)	10.779(2)
<i>c</i> (Å)	13.485(3)
α (deg)	95.885(4)
β (deg)	98.485(5)
γ (deg)	103.936(4)
volume (Å ³)	1082.8(4)
Temperature (K)	100 (2)
space group	<i>P</i> - <i>l</i> (No. 2)
Ζ	2
reflections measured	3264
independent reflections	2907 [<i>R</i> (int) = 0.0141]
calculated density (mg/m ³)	1.271
absorption coefficient (mm ⁻¹)	0.075
F (000)	440
goodness-of-fit on F ²	1.016
final <i>R</i> indices $[I \ge 2\sigma(I)]$	$R_1 = 0.0770$, $wR_2 = 0.2302$
R indices (all data)	$R_1 = 0.0928$, $wR_2 = 0.2529$

 Table S1. Crystal Data for Chromenes CP-H.



Figure S1. The decay kinetics for bleaching of the colored intermediates of **CP-H** (a) and **CP-OMe** (b) as monitored by change in the absorbance at 524 nm at 298 K.



Figure S1. ¹H NMR (CDCl₃, 500 MHz) spectrum of *rac*-4-formyl[2.2]paracyclophane in CDCl₃.



Figure S2. ¹H NMR (CDCl₃, 500 MHz) spectrum of *rac*-4-hydroxy[2.2]paracyclophane in CDCl₃.



Figure S3. ¹H NMR (CDCl₃, 500 MHz) spectrum of CP-H in CDCl₃.



Figure S4. ¹³C NMR (CDCl₃, 125 MHz) spectrum of cyclophanochromene CP-H in CDCl₃.



Figure S5. ¹H NMR (CDCl₃, 500 MHz) spectrum of 4,16-diformyl[2.2]paracyclophane in CDCl₃.



Figure S6. ¹H NMR (CDCl₃, 500 MHz) spectrum of 4,16-dihydroxy[2.2]paracyclophane in CDCl₃.



Figure S7. ¹H NMR (CDCl₃, 500 MHz) spectrum of 4-hydroxy-16-methoxy[2.2]paracyclophane in CDCl₃.



methoxy[2.2]paracyclophane in CDCl₃.



Figure S9. ¹H NMR (CDCl₃, 500 MHz) spectrum of CP-OMe in CDCl₃.



Figure S10. ¹³C NMR (CDCl₃, 125 MHz) spectrum of CP-OMe in CDCl₃.