Supplementary Data:

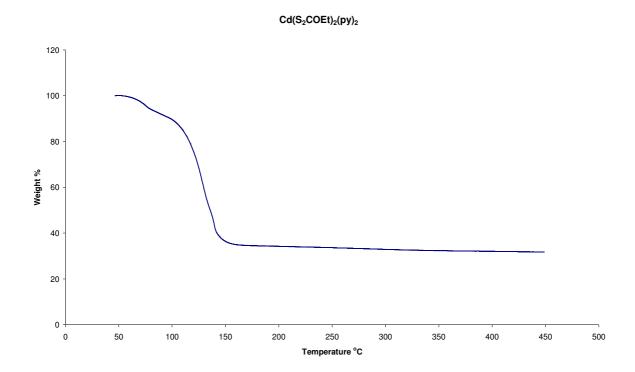
 1 H NMR (CDCl₃): 1.44 (t, 3H, CH₃; 2 J = 7.1 Hz), 4.48 (q, 2H, CH₂; 2 J = 7.1 Hz), 7.42 (m, 2H, m-CH), 7.82 (m, 1H, p-CH), 8.76 (m, 2H, o-CH).

¹³C (CDCl₃): 14.2 (CH₃), 74.4 (CH₂), 125.0, 138.4, 149.6 (C₅H₅N), 231.5 (CO)

Micro Analysis: Found (Calc) for C16 H20 N2 S4 O2 Cd: C 37.3 (37.5), H 3.95 (3.93), N 5.34 (5.46)

Transient absorption spectroscopy measurements:

Transient absorption data were collected using a highly sensitive microsecond absorption systems under N_2 . Data was collected using an excitation wavelength of 567 nm, with pulse width 0.6 ns and pulse energy density of 60-80 μ J cm-2 at 4 Hz repetition from a dye laser (Photon Technology International Inc. GL-301) pumped by a nitrogen laser (Photon Technology International Inc. GL-3300). Samples were probed using a quartz halogen lamp (Bentham, IL1) with a stabilized power supply (Bentham, 605). Probe light was detected by a silicon photodiode and the signal subsequently amplified and passed by electronic band-pass filters to improve signal to noise.



Crystallography:

Crystallographic data for the structural analysis (in CIF format) has been deposited with the Cambridge Crystallographic Data Center, CCDC no. 753046. Copies of this information may be obtained from the Director, CCDC, 12 Union Road, Cambridge, CB21EZ, UK (Fax: +44-1233-336033; e-mail: deposit@ccdc.cam.ac.uk or www.ccdc.cam.ac.uk).

Table 1. Crystal data and structure refinement for 1.

h09kcm32
C16 H20 Cd N2 O2 S4
512.98
150(2) K
Monoclinic
P 2 ₁ /c
a = 10.1957(3)Agst alpha = 90deg
b = 13.1149(3)Agst beta = 104.450(2)deg
c = 16.4045(3)Agst gamma = 90deg
2124.15(9) Agst@3
4
1.604 Mg/m@3
1.433 mm@-1
1032
0.25 x 0.20 x 0.20 mm
4.13 to 27.51 deg.
-13<=h<=13; -17<=k<=17; -21<=l<=21
40865
4864 [R(int) = 0.0936]
3647
0.995
0.7626 and 0.7159
Full-matrix least-squares on F@2
4864 / 0 / 255
1.055
$R \sim 1 = 0.0335$ w $R \sim 2 = 0.0660$
$R\sim1 = 0.0577 \text{ wR}\sim2 = 0.0733$
0.445 and -0.755 e.Agst@-3