

Supporting Information for:
**“Distribution and Orientation Study of Dyes Intercalated into Single
Sepiolite Fibers. A Confocal Fluorescence Microscopy Approach”**

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Fig. S1. Typical intensity profiles for dye/Sep particles: (A) along the main C-axis and (B) along the short b-axis.

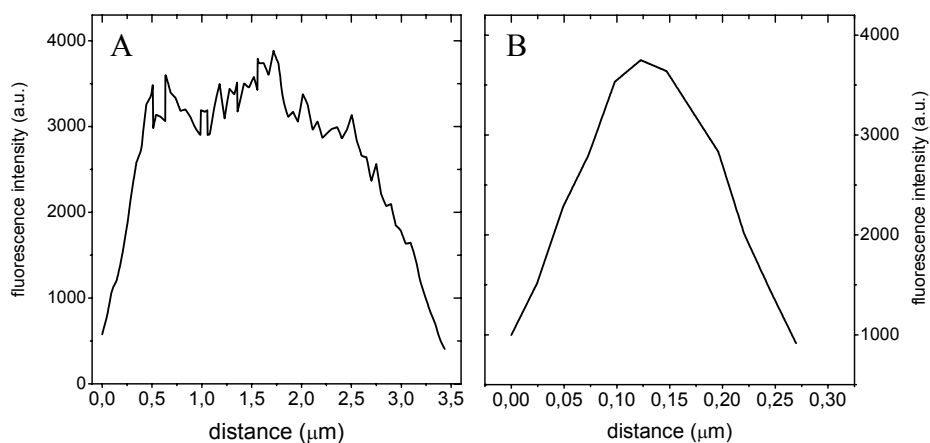


Fig. S2. Polarized intensity fluorescence images of 50% CEC R6G/Sep fiber. Arrows indicates the direction of polarization registered in each detection channel.

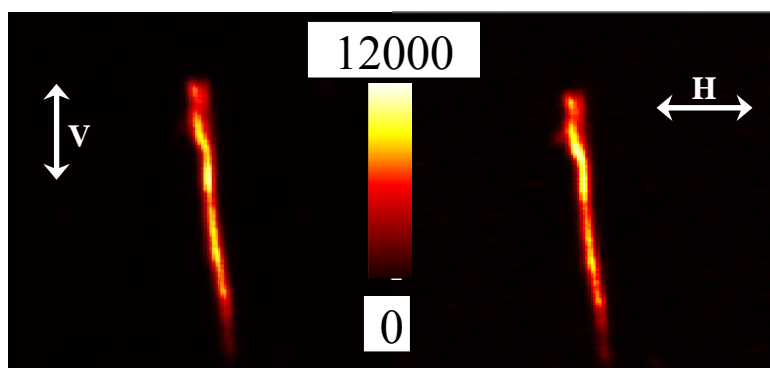


Fig. S3. (A) FLIM image of a single Sepiolite fiber dopped with LDS 698; (B) Fluorescence decay curve of the dye/Sep fiber (inset: residuals); (C) Histogram of the fitted lifetimes.

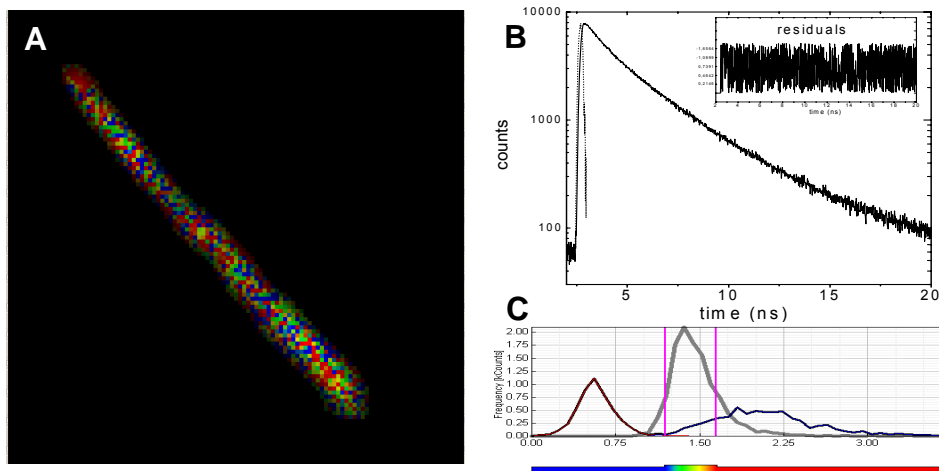


Fig. S4. Absorption and fluorescence spectra of LDS 698 (green) and LDS 722 (blue) in ethanol; solid lines indicates diluted dye solutions ($C = 10^{-6}$ M) measured with cuvettes of 1 cm of optical path) and dashed lines concentrated dye solutions ($C = 10^{-3}$ M) measured in cuvettes with 0.001 cm of optical path)

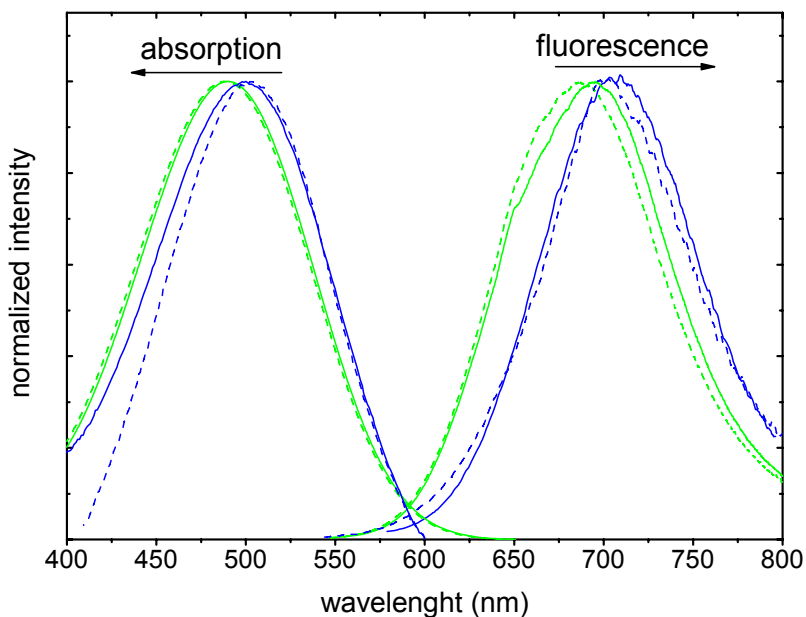


Fig. S5. (A) Fluorescence spectra and (B) Fluorescence decay curves of LDS 698 in ethanol (red) and in HEMA polymer matrix(Hydroxyethylmethacrylate).

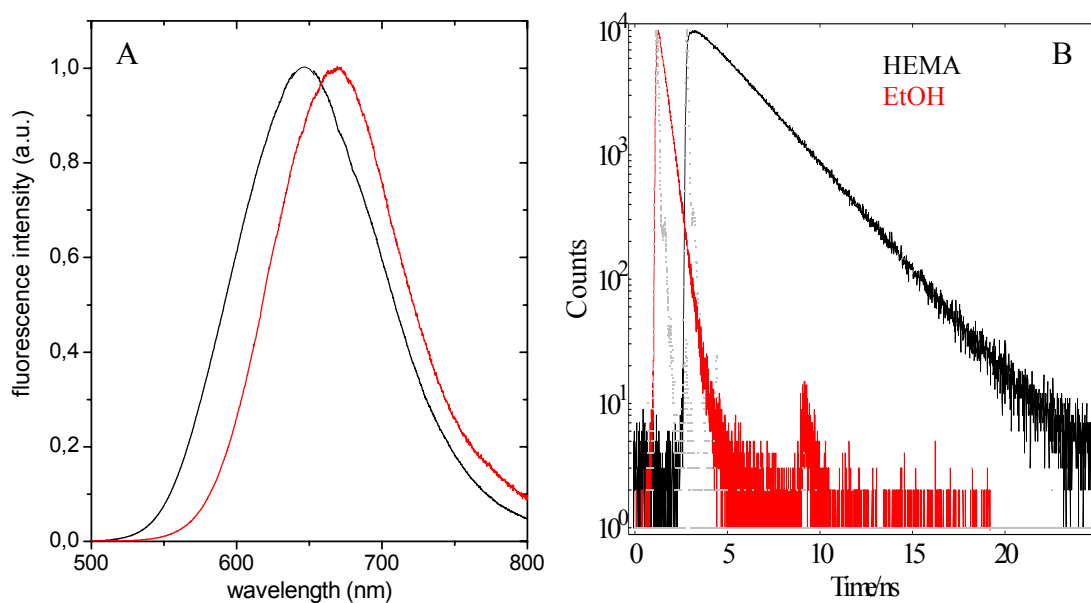


Table S1: Spectroscopic data of LDS 698 and LDS 722 in ethanol and embedded in the matrix of HEMA (Hydroxyethylmethacrylate) polymer.

	$\lambda_{ab}(nm)$	$\epsilon_{max} (10^4 M^{-1} cm^{-1})$	$\lambda_{fl} (nm)$	ϕ	$\tau (ns)$
LDS698	490.5	3.1	670.0	0.093	0.36
LDS698/HEMA	486.5	2.7	647.0	0.22	2.67
LDS722	504.5	3.3	695.0	0.14	0.58
LDS722/HEMA	507.5	2.3	665.0	0.35	2.54

Fig. S6. Fluorescence decay curves registered in both detectors: parallel (black) and perpendicular (red) with respect to the long axis of LDS 698/Sep fiber. Dichroic ratio, $D = I_{\parallel} / I_{\perp}$, analyzed at different time windows; markers corresponds to the part of the decay fitted for short and long lifetime of the LDS 698 adsorbed in Sep.

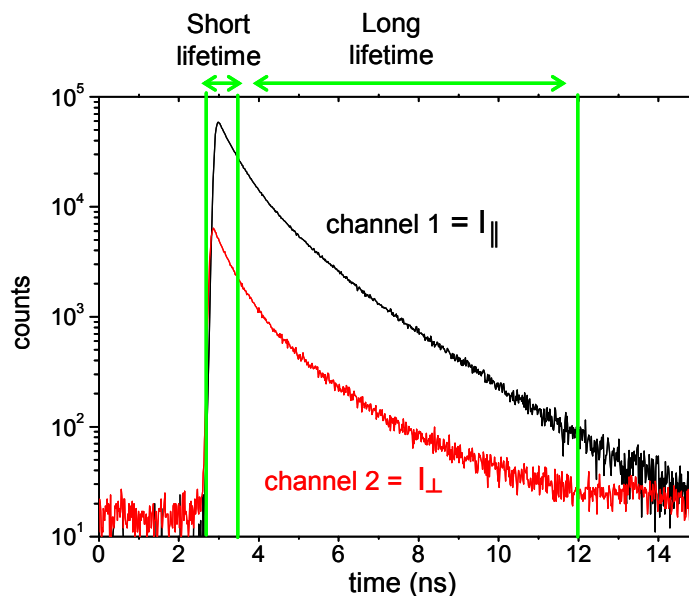


Fig. S7. FLIM image of LDS722/Sep 130% CEC together with the histogram of lifetimes.

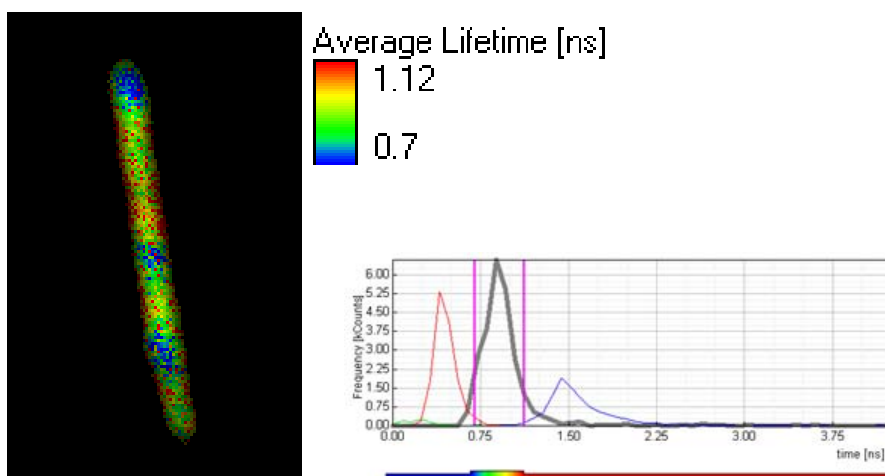


Fig. S8. Emission spectra of LDS 722 in solution (black line) and LDS 722 in single Sepiolite particle (red), measure in the same conditions in a Fluorescence Confocal Microscope by a spectrograph and a CCD camera mounted in a side port.

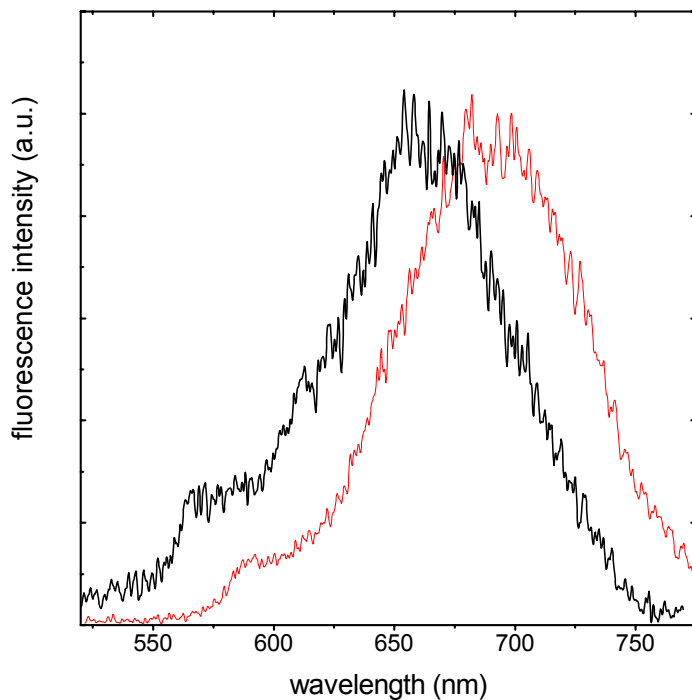


Fig. S9. FLIM image (left) of 6% CEC PY/Sep; Fluorescence decay curve along the particle (top right) and histogram of lifetimes (bottom right)

