

Carbon-dot-hydrogel for enzyme-mediated bacterial detection

Sagarika Bhattacharya,^a Sukhendu Nandi,^{a,#} and Raz Jelinek^{*a,b}

^a Department of Chemistry, Ben Gurion University of the Negev, Beer Sheva 84105, Israel.

E-mail: razj@bgu.ac.il; Fax: (+)972-8-6472943

^b Ilse Katz Institute for Nanotechnology, Ben Gurion University of the Negev, Beer Sheva 84105, Israel.

Present address: Ruhr Universitat Bochum, Universitaetsstrasse 150, D-44780, Bochum, Germany.

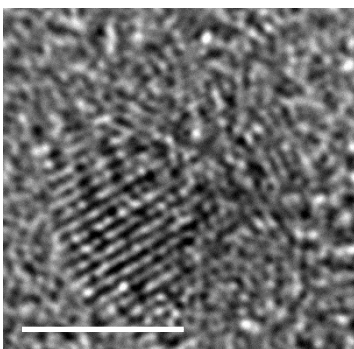


Fig. S1 High resolution transmission electron microscopy (HRTEM) image of a representative carbon dot embedded within the hydrogel, showing the graphite crystal planes. Scale bar is 2 nm. HRTEM images were recorded on a 200 kV JEM-2100F transmission electron microscope (JEOL, Peabody, MA). For the HRTEM measurements, 0.5 mg of C-dots synthesized as described above were dissolved in 500 mL of chloroform, and 10 mL of the solution was placed upon an ultrathin carbon-film-coated Cu grid, dried at room temperature for 2 h, and imaged.

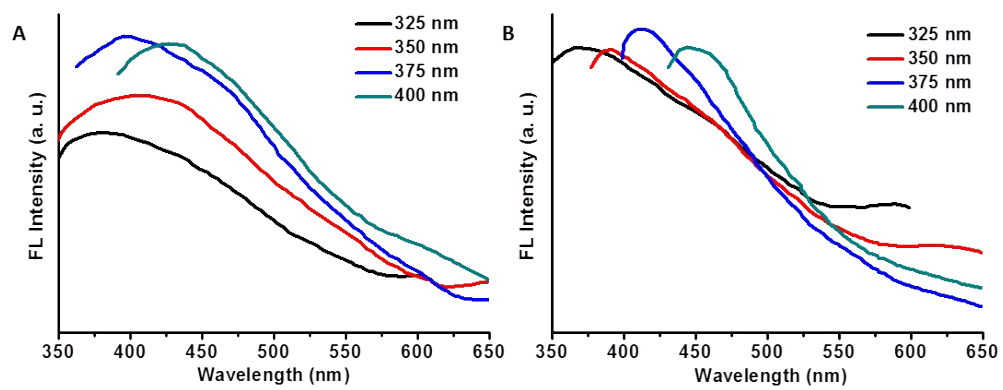


Fig. S2 Photoluminescence spectra of C-dot embedded in hydro gel (**A**) and in solution excited at different wavelengths (**B**) (10 mg mL^{-1}) were recorded on FL920 spectrofluorimeter.