

Electronic supplementary information

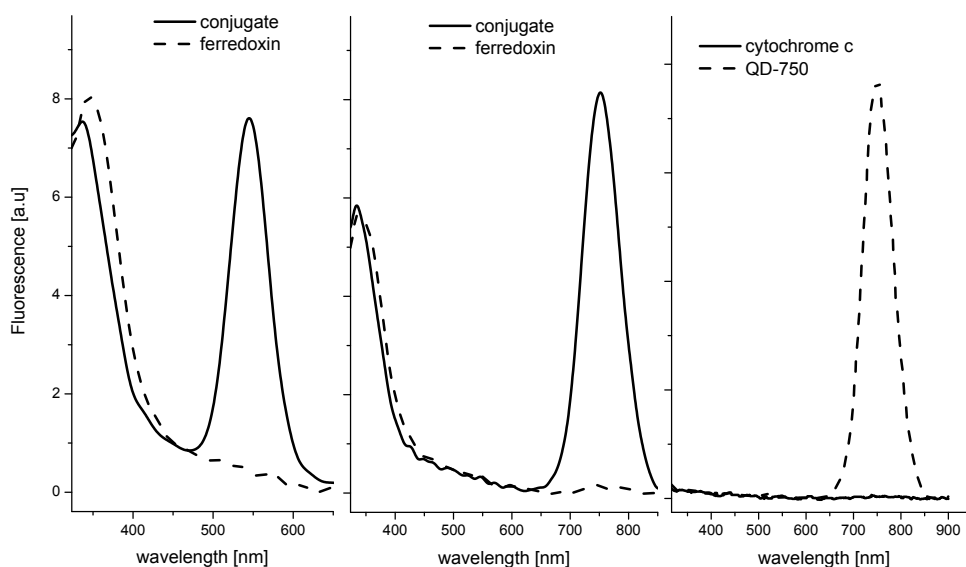


Fig.S-1. Fluorescence emission spectra (extinction at 280 nm) for 1 ml fractions collected during separation of Fd incubated with QD-550 (a) or QD-750 (b) (under illumination), respectively at 13 ml/12 ml (conjugate) and at 16.5 ml (ferredoxin), and during separation of Cyt c incubated with QD-750 (c), respectively at 13 ml (QD-750) and at 17 ml (cytochrome c) For details, see Fig.3

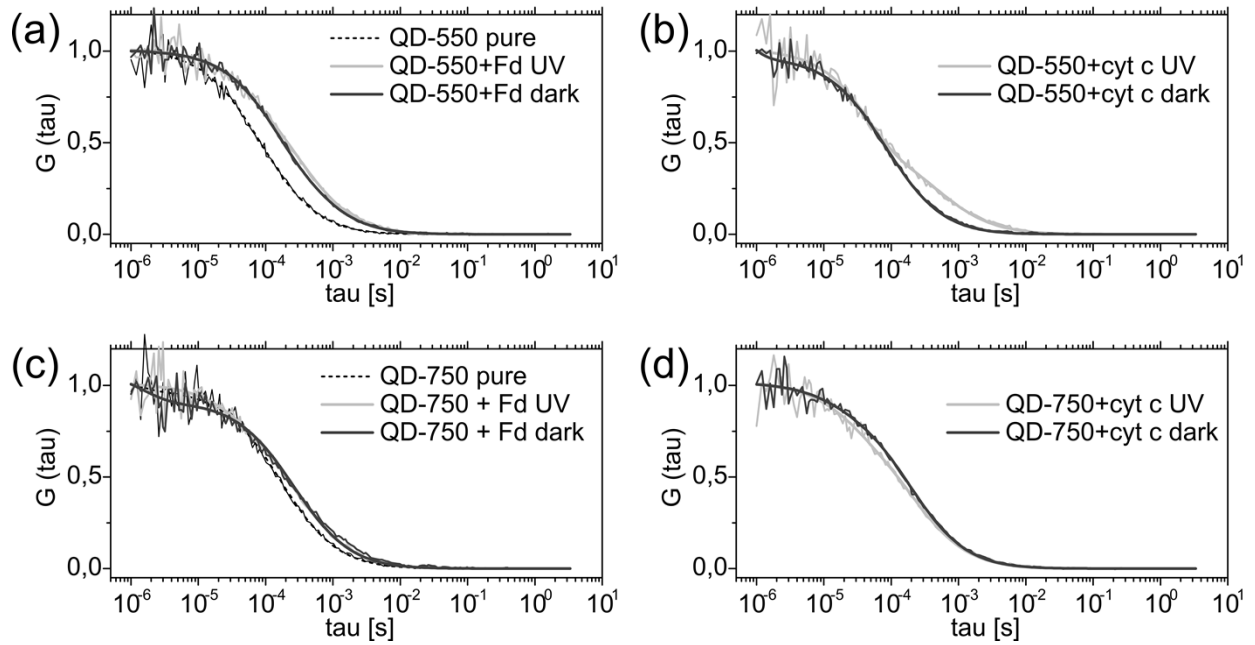


Fig.S-2 Examples of FCS autocorrelation curves, recorded for different experimental variants for pure QDs and after its incubation for 30 min with respective proteins, in darkness or at illumination (a) QD-550 + Fd, (b) QD-550+ Cyt c, (c) QD-750+Fd, (d) QD-750+Cyt c.

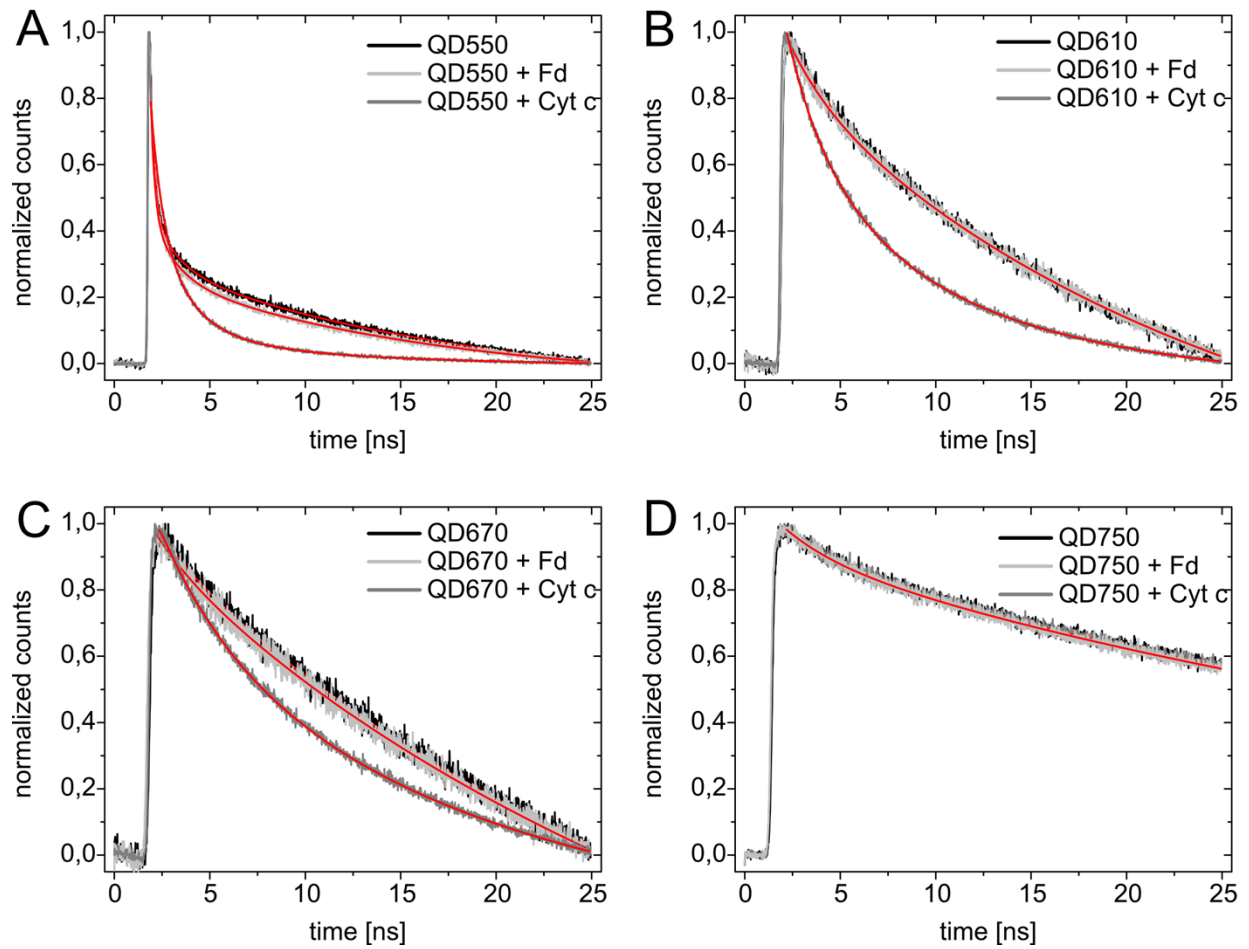


Figure S-3. Normalized luminescence decay curves (representative examples) for pure QDs and their mixture with Fd or Cyt c. Solid lines presents fitted decay.

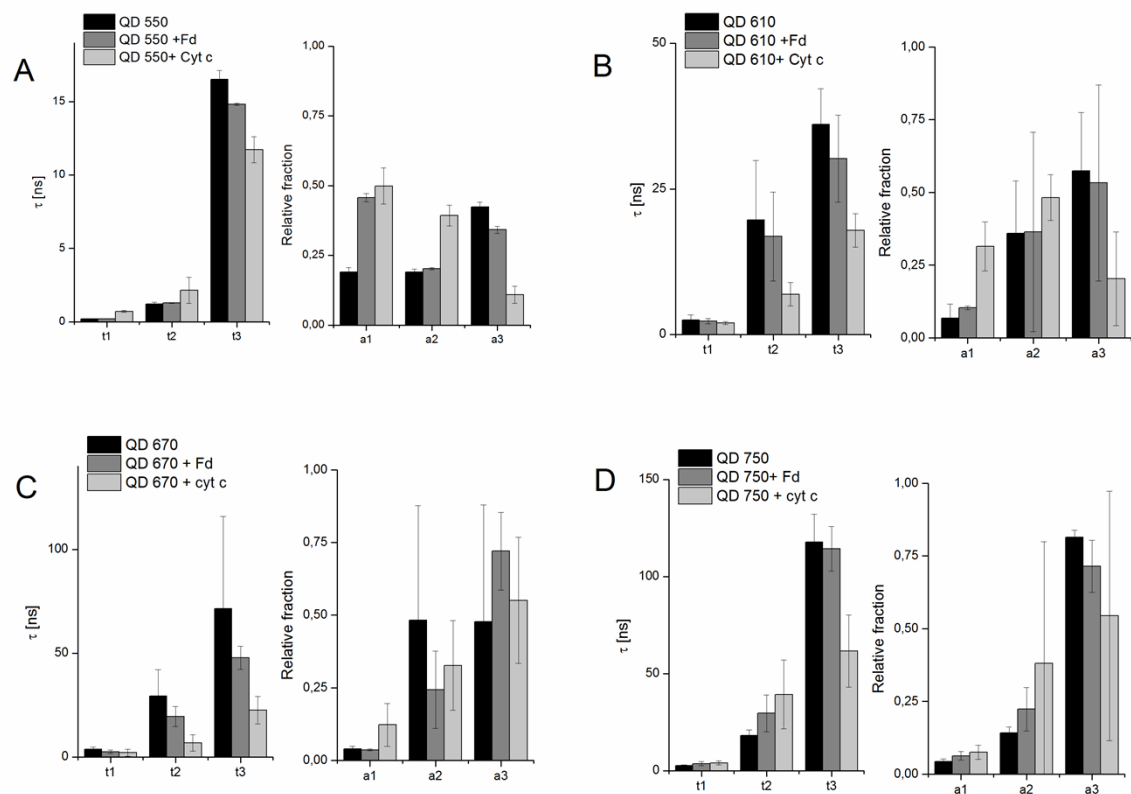


Figure S-4. Detailed characteristics of luminescence decay curves QDs and its mixtures with Fd or Cyt c. Lifetime values (t_1 , t_2 , t_3) and corresponding relative fractions (a_1 , a_2 , a_3) were averaged from fitting of decay curves of three independent measurements. Error bars represent SD. Note that values were averaged due to τ , ordered by its increasing length (shortest, medium, longest) and high SD for some experimental cases corresponds to possibility of more than one good fitting.