Supporting Information

Distance-dependent metal-enhanced quantum dots fluorescence analysis in solution by capillary electrophoresis and its application to DNA detection

Yong-Qiang Li,^{†,‡} Li-Yun Guan,^{†,‡} Hai-Li Zhang,[†] Jun Chen,^{†,‡} Song Lin,^{†,‡} Zhi-Ya Ma,^{†,‡} and

Yuan-Di Zhao*,^{†,‡}

[†] Britton Chance Center for Biomedical Photonics, Wuhan National Laboratory for Optoelectronics-Huazhong University of Science and Technology, Wuhan 430074, P. R. China

[‡] Key Laboratory of Biomedical Photonics of Ministry of Education, Colleage of Life Science and Technology, Huazhong University of Science and Technology, Wuhan 430074, P. R. China



Figure S1. Hydrodynamic diameters of Au NPs (a) and Au-B_{3.4} conjugate (b) in water measured by

means of DLS.



Figure S2. Zeta potentials of Au NPs (a) and Au- $B_{3.4}$ conjugate (b) in water.



Figure S3. UV-Vis absorption spectra of filtrate collected during the preparation of $Au-B_{3.4}$

conjugate (a), and 3 OD of $B_{3,4}$ treated by the same dilution procedure in control experiment (b).



Figure S4. Hydrodynamic diameters of CdSe/ZnS QDs (a) and CdSe/ZnS QD-A_{3.4} conjugate (b) in

PBS buffer (10 mM, pH 7.4) measured by means of DLS.



Figure S5. Zeta potentials of CdSe/ZnS QDs (a) and CdSe/ZnS QD-A_{3.4} conjugate (b) in PBS

buffer (10 mM, pH 7.4).



Figure S6. Effect of polymer additive on the separation of hybridization mixture of QD-A_{3.4} and

Au-B_{3.4} conjugates. Sample: 50 μL QD-A_{3.4} + 200 μL Au-B_{3.4}. Separation buffers: (a) 25 mM
Na₂B₄O₇, pH 9.0; (b) 25 mM Na₂B₄O₇ + 0.5% HEC, pH 9.0. Coated capillary with 40 cm effective (60 cm total) length and 50 μm I.D. was used. Applied voltage was -20 kV, and hydrodynamic injection was carried out by siphoning at 15 cm height differences for 20 s. The relative standard deviations (%RSD) for the migration time of peaks 1 and 2 in curve a and b (three repeat experiments) were 0.54% and 0.55%, 0.57% and 0.59%, respectively, while the %RSD for their peak areas were 1.89% and 1.98%, 1.86% and 1.89%, respectively.



Figure S7. Electropherograms for the QD-A_{11.9}~B_{11.9}-Au system (50 μL QD-A_{11.9} + 250 μL Au-B_{11.9}, a), and the mixture of QD-A_{11.9}~B_{11.9}-Au system and 150 nM non-target DNA (b). The temperature and time of the competition reaction were 37 °C and 1.5 h, respectively. Other

conditions were same as described in Figure S6. The %RSD for the migration time of peaks in curve a and b (three repeat experiments) were 0.61% and 0.62%, respectively, while the %RSD for

their peak areas were 1.83% and 1.84%, respectively.