

Thermodynamic Investigation of the Interaction between Ionic Liquid Functionalized Gold Nanoparticles and Human Serum Albumin for Selective Determination of Glutamine

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SUPPLEMENTARY MATERIALS

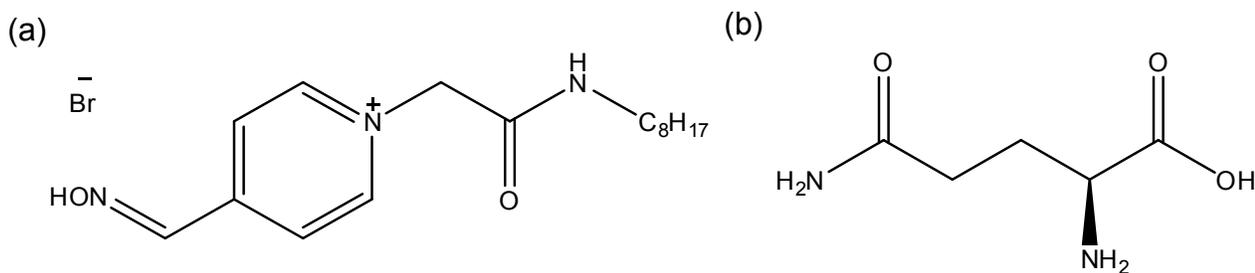
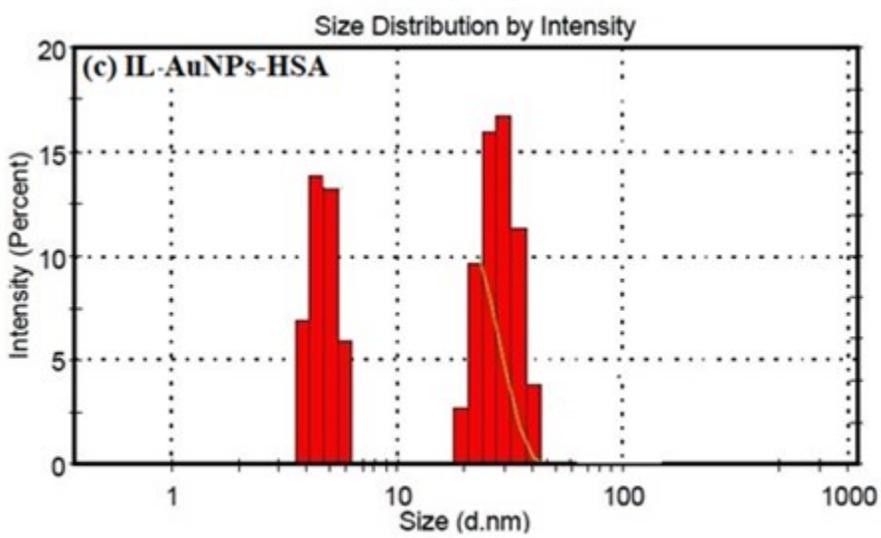
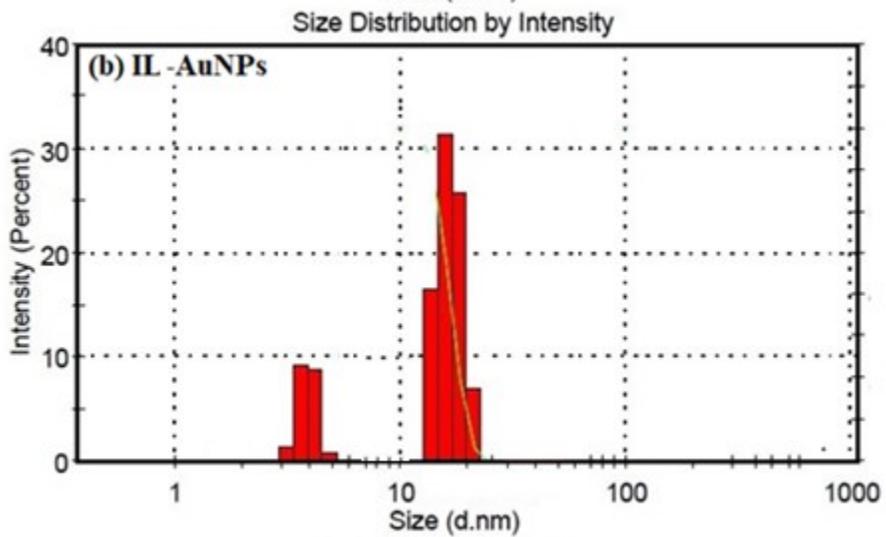
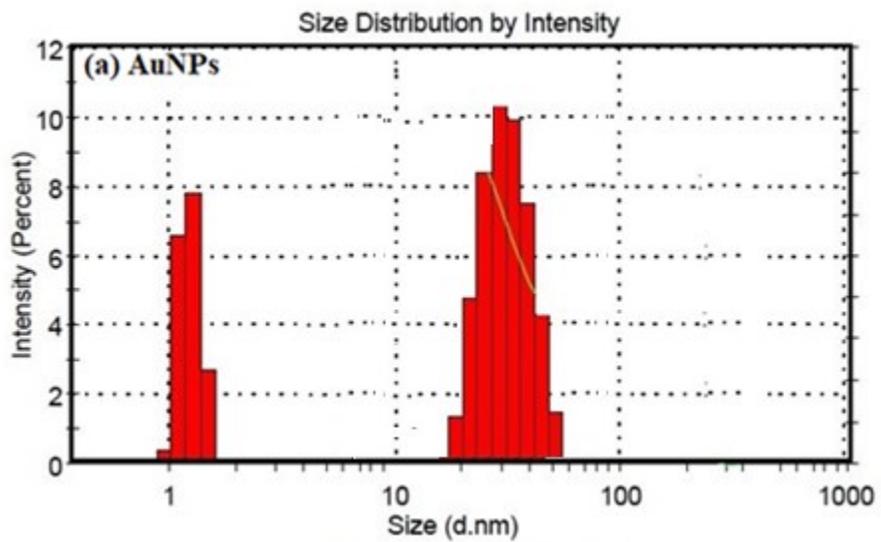


Fig. S1 Chemical structure of (a) amino based ionic liquid [4-((hydroxiimino)methyl)-1-(2-(octylamino)-2-oxoethyl)pyridine-1-iumbromide] and (b) glutamine.



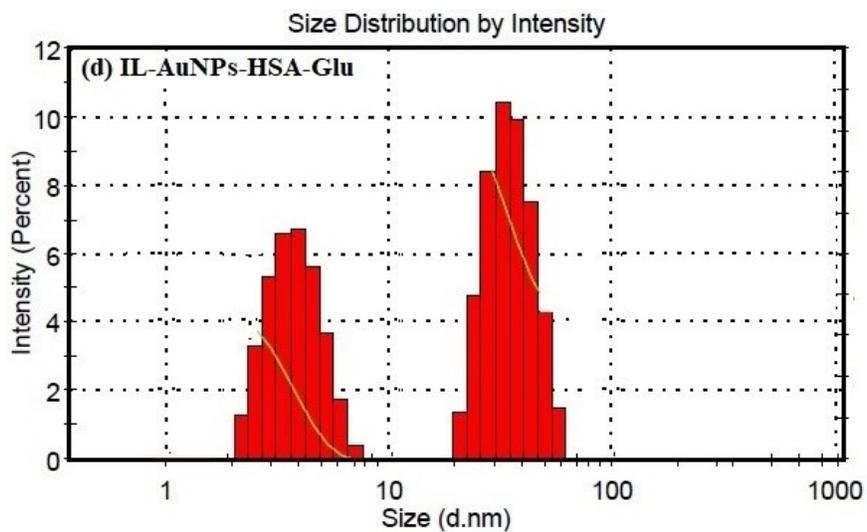


Fig. S2 DLS size distribution plots of (a) AuNPs, (b) IL-AuNPs (c) IL-AuNPs-HSA and (d) IL-AuNPs-HSA with Glu.

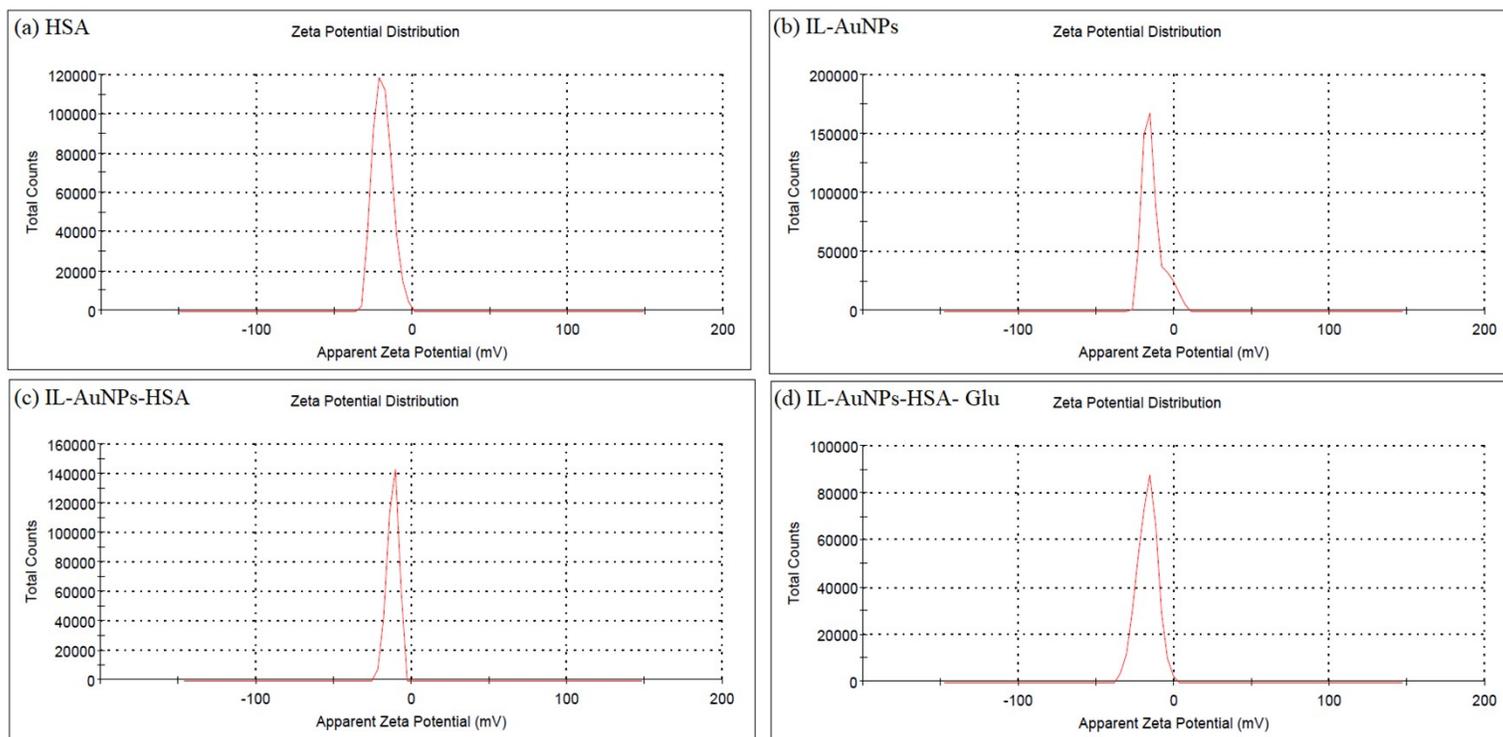


Fig. S3 Zeta potential measurements of (a) HSA, (b) IL-AuNPs (c) IL-AuNPs-HSA and (c) IL-AuNPs-HSA with Glu.

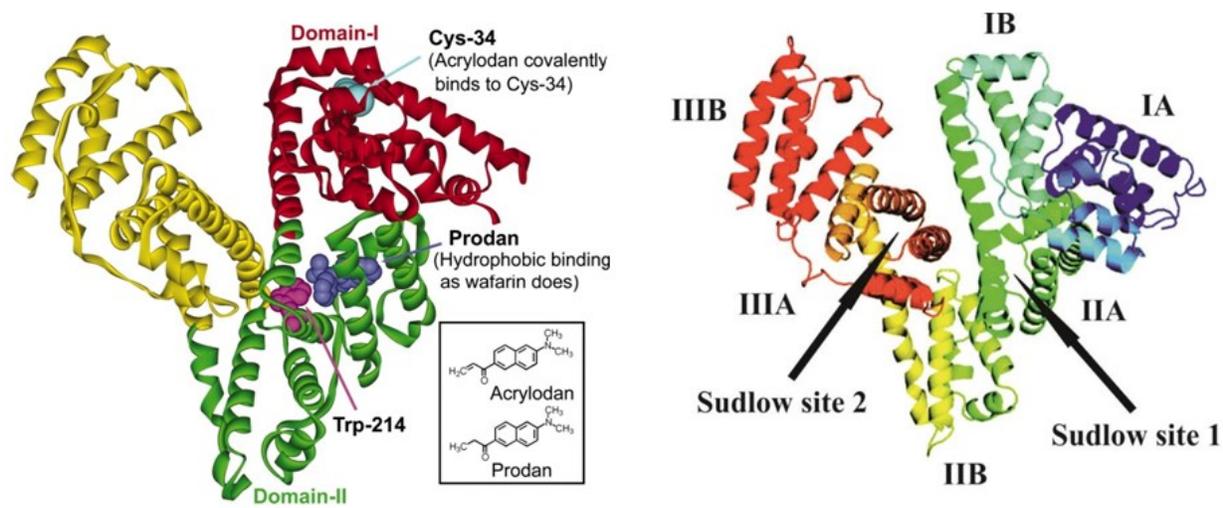


Fig. S4 Structure of HSA showing three different domains.

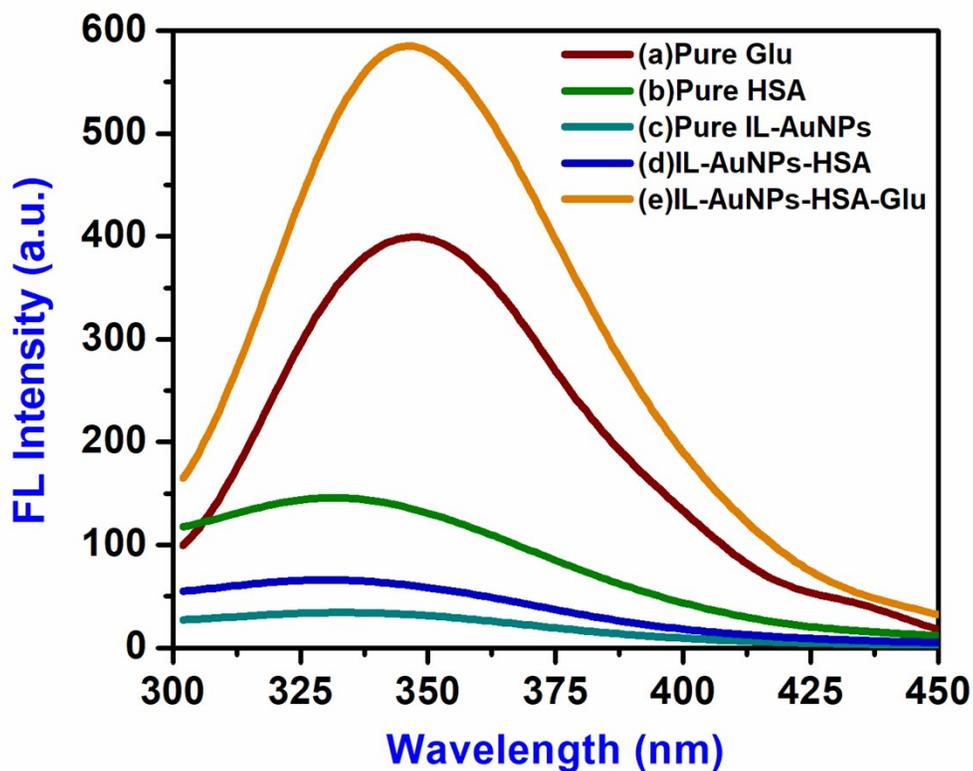


Fig. S5 Fluorescence spectra for (a) pure Glu, (b) pure HSA, (c) pure IL-AuNPs, (d) IL-AuNPs-HSA and (e) IL-AuNPs-HSA with Glu.

Table S1. Comparison of sensitivities of detection techniques with the proposed methodology

Method	Analyte	Concentration	Correlation Coefficient	Limit of Detection	Ref
Colorimetry	Lysozyme	100-1000 nM	0.9890	50 nM	52
Colorimetry	Lysozyme	20 nM to 2 μ M	0.99	20 nM	53
Impedimetric	MWCNT-SPEs-aptamer	0-28.6 μ M	0.9984	862 nM	54
Capillary Electrophoresis	Lysozyme	0.005-0.3 μ M	0.9930	8 nM	55
Fluorescence	FITC-BSA	0.0-0.10 μ g/ml	0.94		56

Colorimetry	Glutamine	10-100 nM	0.99627	0.67	Present Method
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