Fluorescence anisotropic probe for sensing cardiac troponin-I antigen through

Target-specific antibody-conjugated gold nanoclusters

^aS Madanan Anju, ^aSusan Varghese, ^aK Abraham Merin, ^aAli Ibrahim Shkhair, Greeshma

Rajeevan, Geneva Indongo, Arathy B.K and *aSony George

Corresponding author,

Dr. Sony George

Associate Professor, Department of Chemistry, School of Physical and Mathematical Science, University of Kerala, Thiruvananthapuram, Kerala, India- 695581

Mob: +919446462933

Email: emailtosony@gmail.com

^a Department of Chemistry, School of Physical and Mathematical Science, University of Kerala, Thiruvananthapuram, Kerala, India- 695581

Supporting documents



Fig.S1. (A) UV- Vis absorption spectrum of glutathione-capped gold nanoclusters (AuNC@GSH). **(B)** FTIR spectra of glutathione-capped gold nanoclusters and their precursors.



Fig.S2. (A) UV- Vis absorption spectrum and fluorescence excitation spectrum of glutathione-capped gold nanoclusters (AuNC@GSH) and (B) Fluorescence emission spectra of glutathione-capped gold nanoclusters with different excitation wavelengths ranging from 360 to 480nm, Fluorescence emission spectrum of AuNC@GSH at 400 nm excitation(inset).



Fig.S3. Fluorescence anisotropy response (Δr) of 0.05 µg/mL cTnI specific monoclonal antibody conjugated glutathione capped gold nanoclusters (AbcTnI@ AuNC@GSH) towards sequential addition of cTnI antigen (from 0 to 45ng/mL) at different incubation times.



Fig.S4. Fluorescence anisotropy response of (Δr) of 0.05 µg/mL cTnI specific monoclonal antibody conjugated glutathione capped gold nanoclusters (AbcTnI@AuNC@GSH) towards sequential addition of cTnI antigen (from 0 to 45ng/mL) spiked serum and its diluted samples.

Table S1: Table of comparison of the analytical performance of the current FA method of detection of cTnI using antibody-conjugated gold nanoclusters with other previous reports.

SI.	Probe	Method	LoD	Referenc
No.				e
1.	Fluorescent europium (III) chelate-dyed nanoparticle	Fluorescence	0.0020µg/L	[32]
2.	ECL functionalized metal- organic framework	Electrochemiluminescence	0.48fg/mL	[33]
3.	Gold nanoparticle labelled antibody platform	Fluorescence Anisotropy	0.50nM	[34]
4.	Carbon nanofiber nanoelectrode array	Electrochemical	0.20ng/mL	[35]
5.	Antibody-conjugated gold nanocluster	Fluorescence Anisotropy	0.91ng/mL	This work

Table S2. Table showing the recovery percentage of cTnI spiked serum sample with FA assay.

SI.	Sample id	Spiked	Recovered	Recovery
No.		Conc. (ng/mL)	Conc.(ng/mL)	percentage (%)
1.	Conc. Serum sample 1	2.70	2.68	99.25
2.	Conc. Serum sample 2	20.00	20.19	100.91
3.	10 Times dil. Serum sample 1	10.85	10.32	94.90
4.	10 Times dil. Serum sample 2	28.50	29.20	102.45
5.	100 Times dil. Serum sample 1	30.86	31.5	102.07
6.	100 Times dil. Serum sample 2	35.65	34.45	96.63