## **Supporting Information**

## Carbon Dots as Analytical Tools for Sensing of Thioredoxin Reductase and Screening of Cancer Cells

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Figure S1-S8



Scheme S1: Synthesis of DTDPA functionalized fCDs



Fig. S1: XPS Spectra of CDs



**Fig. S2** Linear Relationship Between the Absorbance of Ruhemann Purple with Alanine as Reference at 570 nm and concentration of amino group was found to be 1.30µg/mL



**Fig. S3 (A)** Zeta potential measurement of CDs (15 mV) and fCDs (4.5 mV) (**B**) FTIR spectra of CDs and fCDs



Fig. S4 Stern-volmer plot of fCDs with increase in concentration of  $Cu^{2+}$  and  $K_{sv}$  was calculated to be  $2.2 \times 10^4$ . Intercept:  $7.2 \times 10^{-1}$  and Slope:  $2.1 \times 10^{-5}$ 



**Fig. S5** Fluorescence decay time study of fCDs, fCDs-Cu<sup>2+</sup> and fCDs-Cu<sup>2+</sup>-TrxR. The average life time of fCDs was decreased from 7.1 ns to 3.7 ns in , fCDs-Cu<sup>2+</sup> indicates the ultrafast transfer of electron from CDs to Cu<sup>2+</sup> metal ion. After the addition of TrxR into sensor probe solution increased the average decay time (6.7 ns).

Analytes	Sensitivity Factor ( $\Delta$ F/F <sub>0</sub> )
TrxR (100 nM)	8.7
GSH (1 mM)	0.37
GSH (5 mM)	0.65
Cys (1 mM)	0.43
GSH reductase (0.1 µM)	0.35
NADPH (200 μM)	0.13
TrxR (100 nM) + EDTA	9.0

Table. S1 Sensitivity Factor calculated in the presence of different disulfide reducing biomolecules



Fig. S6 Km determination of fCDs-Cu<sup>2+</sup> (5.5  $\mu$ g)



**Fig. S7** Limit of detection calculation of sensor probe for TrxR (LOD =  $3 \times (0.017/0.0017) = 20$  nM)



Fig. S8 (A) Fluorescence response of fCDs-Cu<sup>2+</sup> in response to TrxR at different pH (B) Effect of salt concentration over sensitivity of fCDs-Cu<sup>2+</sup>



Fig. S9 MTT assay of CDs and fCDs against MCF-7 cells



Fig. 10 MTT assay of fCDs-Cu<sup>2+</sup> against MCF-7 cells

XEVO G2-XS QTOP





Fig. S12 Mass spectra of CDsM-Cu after addition of TrxR (Incubation time was 2 h)