

Supporting Information

**A label-free fluorescent sensor based on silicon quantum dots-MnO₂ nanosheets
for the detection of α -glucosidase and its inhibitor**

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Reagents

α -glucosidase was purchased from Sigma Reagents Company. Tetramethylammonium (TMA) was purchase from Beijing Chemical Works. L-ascorbic acid-2-O- α -D-glucopyranosy (AAG), (3-aminopropyl) trimethoxysilane (APTES) and acarbose were purchased from Shanghai Aladdin Co. Ltd. Hydrogen peroxide (H_2O_2), ascorbic acid (AA), glucose, aspartic acid, histidine, and glutamate were purchased from Beijing Dingguo Biotechnology Co. Ltd. $MnCl_2 \cdot 4H_2O$, NaCl, $CaCl_2$, $ZnCl_2$ were purchased from Tianjin Guangfu Institute of Fine Chemicals. Tyrosinase (TYR), glucose oxidase (GO_x), pepsin, trypsin and protein kinase (PKA) were purchased from Sino-American Biotechnology Co. Ltd. Trisodium citrate was purchased from Sinopharm Chemical Reagent Co. Ltd. All chemicals used were of analytical reagent grade without further purification. Phosphate buffered saline (PBS) was prepared by mixing different ratios of Na_2HPO_4 and NaH_2PO_4 solution. The resistivity of distilled water used in all experiments was higher than $18 M\Omega cm^{-1}$.

Instruments

Scanning electron microscope (SEM) experiment was performed on a JF6700 scanning electron Microscope (JEOL Ltd, Japan). Fluorescence spectra were collected with a Shimadzu RF-5301 PC spectrofluorophotometer (Shimadzu Co.Ltd, Kyoto, Japan). UV-vis absorption spectra were collected on Varian GBC Cintra 10e UV-visible Spectrophotometer (Shimadzu Co.Ltd, Kyoto, Japan). FT-IR spectra was recorded by using a Bruker IFS66V FT-IR Spectrometer (Bruker Corporation, Germany). All pH measurements throughout the study were performed with a PHS-3C pH meter (INESA Scientific Instrument Co. Ltd, Shanghai, China).

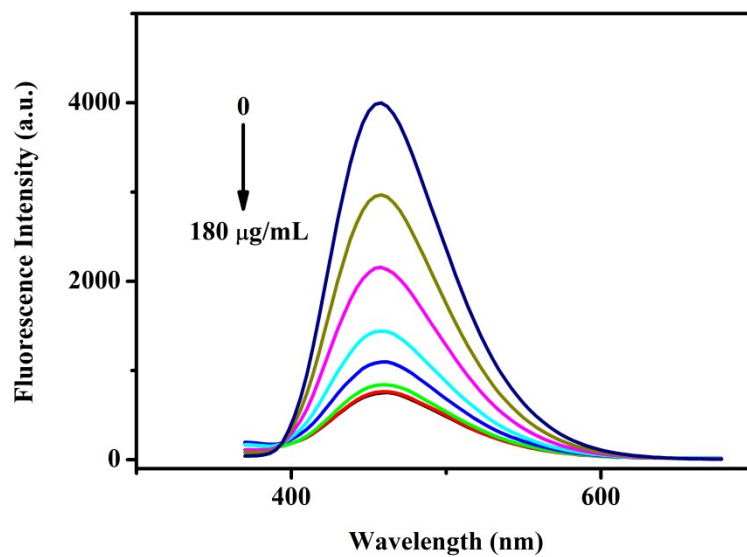


Fig. S1. Fluorescence emission spectra of SiQDs with different concentrations of MnO₂ nanosheets (0, 20, 40, 80, 120, 140, 160 and 180 μg mL⁻¹).

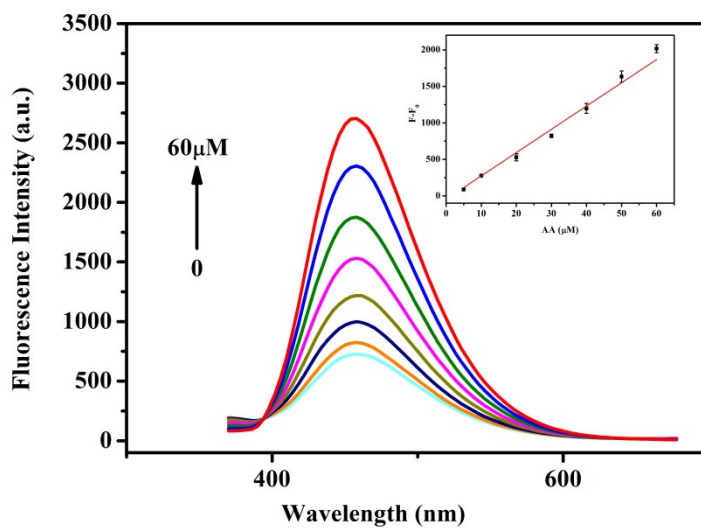


Fig. S2. Fluorescence emission spectra of SiQDs/MnO₂ nanosheets with different concentrations of AA (0, 5, 10, 20, 30, 40, 50 and 60 μM).

Table S1. Detection of α -glucosidase in human serum samples

Sample	Added (U/mL)	Founded (U/mL)		Recovery (%)	RSD (n=3,%)
		Colorimetry	Our method		
serum	0	0.060	0.063	-	-
	0.050	0.13	0.11	94.00	0.75
	0.10	0.16	0.17	107.0	3.24
	0.45	0.47	0.50	97.11	2.50

Table S2. Comparison of our method with the previous methods for the detection of α -glucosidase

Methods	Materials	Linear range (U/mL)	LOD (U/mL)	Reference
Colorimetric assay	Gold nanoparticles	0.05 – 1.1	0.004	[1]
Fluorescence assay	N-doped CDs	0.2 – 10	0.01	[2]
Fluorescence assay	Conjugated polymer and PNPg	0.1 – 0.5	0.01	[3]
Electrochemical assay	Gold nanoparticle-modified gold electrode	0.1 – 1.1	0.04	[4]
Electrochemical assay	AgNPs/DA and MNPs/ pAPG with PBA/GE	0 – 1.1	0.04	[5]
Fluorescence assay	SiQDs/MnO ₂ nanosheets	0.02 – 2.5	0.007	This work

Table S3. Comparison of our method with the previous methods for the detection of acarbose

Methods	Materials	Linear range (μM)	IC50 (μM)	Reference
Fluorescence assay	N-doped CDs	0.1 – 1000	58.68	[2]
Colorimetric assay	Gold nanoparticles	–	5.87	[6]
Fluorescence assay	β -CD-CQDs nanoprobe	50 – 500	319	[7]
Fluorescence assay	N,B-CDs	0.03–5000	58	[8]
Fluorescence assay	SiQDs/MnO ₂ nanosheets	1–1000	33.88	This work

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