SUPPORTING MATERIAL

Gold-coated micellar composites as colorimetric probes for the determination of low molecular weight thiols in biological fluids using consumer electronic devices

Elli Akrivi,^{1,2+} Athanasios G. Vlessidis,³ Dimosthenis L. Giokas,^{3*} Nikolaos Kourkoumelis^{1*}

¹ Department of Medical Physics, School of Health Sciences, University of Ioannina, Greece

² Neurology Clinic, University Hospital of Ioannina, Greece

³ Department of Chemistry, School of Natural Sciences, University of Ioannina, Greece

^{*} Corresponding authors. Email: <u>nkourkou@uoi.gr</u>, <u>dgiokas@uoi.gr</u>

Preparation of artificial and simulated body fluids

Artificial urine solution (AUS) was prepared by mixing 12 mM Na₂SO₄, 1.5 mM uric acid, 2.45 mM trisodium citrate dihydrate, 7.8 mM creatinine, 250 mM urea, 31 mM KCl, 30 mM NaCl, 1.66 mM CaCl₂, 23.67 mM NH₄Cl, 0.19 mM potassium oxalate hydrate, 4.4 mM MgSO₄·7H₂O, 18.7 mM NaH₂PO₄·2H₂O and 4.7 mM Na₂HPO₄·2H₂O and adjusting the pH at the value of 6.0 ± 0.2 [S1,S2]. Artificial blood plasma (ABP) contained 137.5 mM NaCl, 4.2 mM sodium hydrogen carbonate, 3.0 mM KCl, 0.5 mM disodium hydrogen phosphate, 0.5 mM MgCl₂, 2.64 mM CaCl₂ and 0.5 mM NaSO₄ in distilled water and adjusting the pH at the value of 7.4 [S2]. Simulated blood plasma (SBP) was prepared by enrichment of ABP with bovine serum albumin (40 g/L), glucose (5.0 mM), urea (3.0 mM), uric acid (220 µM) and a mixture of common amino acids found in blood plasma (0.4 mM of glutamine, glycine, valine, arginine, lysine and alanine; total concentration of 2.4 mM) [S3]. All artificial and simulated body fluids were fortified with cysteine (at variable concentrations depending on the needs of the analysis), which the most abundant biothiol species in biological fluids.

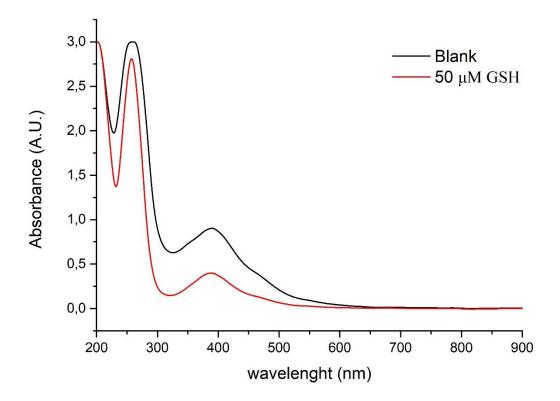


Figure S1. Absorbance spectra of Au-CTAB complex (black line) and Au-CTAB in the presence of 50 μ M of glutathione. No peaks above 500 nm are observed suggesting that gold has not been reduced to its respective gold nanoparticle species under the optimum experimental conditions (0.25mM AuCl4⁻, 10 mM CTAB, 50 μ M GSH, sodium acetate/acetic acid buffer pH 6, 15 min incubation time at room temperature).

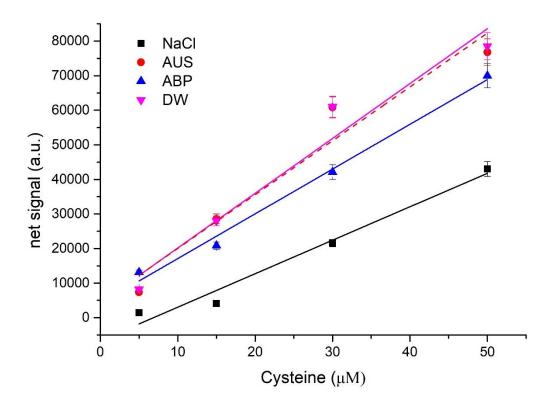


Figure S2. Response of the colorimetric assay in various artificial biofluids. The linear curves are the result of linear regression while error bar represent the standard error calculated for triplicate samples. AUS: Artificial urine solution, ABP: artificial blood plasma, DW: distilled water.

References for the Supporting Material

[S1] T. Brooks, C.W. Keevil, A simple artificial urine for the growth of urinary pathogens, Lett. Appl. Microbiol. 24 (1997) 203-206.

[S2] N. Sarigul, F. Korkmaz, İ. Kurultak, A new artificial urine protocol to better imitate human urine, Sci. Rep. 9 (2019) 20159.

[S3] L. Liu, C.L. Qiu, Q. Chena, S.M. Zhang, Corrosion behavior of Zr-based bulk metallic glasses in different artificial body fluids, J. Alloys Comp. 425 (2006) 268–273.

[S4] E, Gyori, I. Fábián, I.Lázár, Effect of the chemical composition of simulated body fluids on aerogel-based bioactive composites, J. Compos. Sci. 1 (2017) 15.