

Supporting information

Sensitive immunosensing of α -Synuclein protein in human plasma samples using gold nanoparticle conjugated with graphene: An innovative immuno-platform towards early stage identification of Parkinson disease using point of care (POC) analysis

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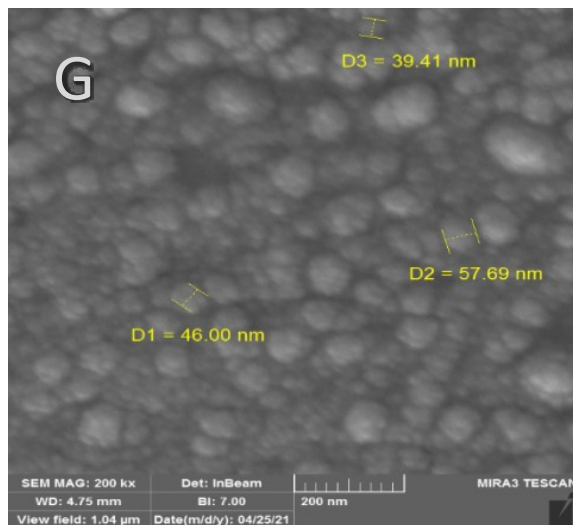
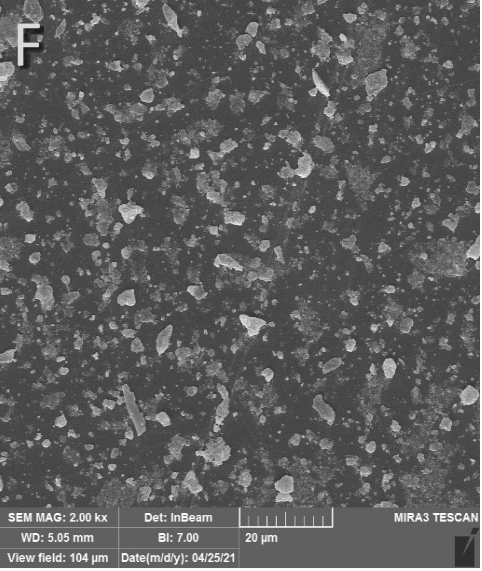
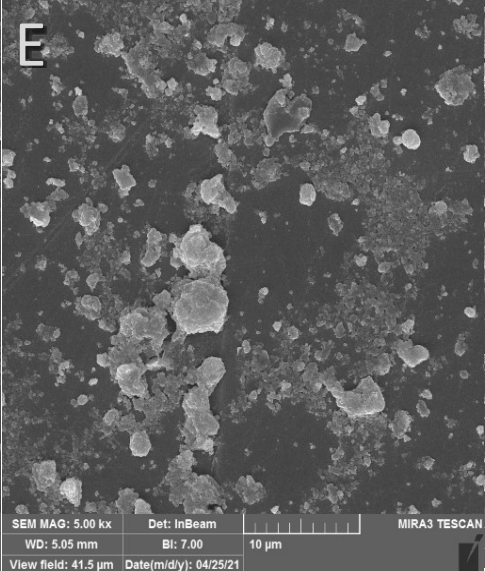
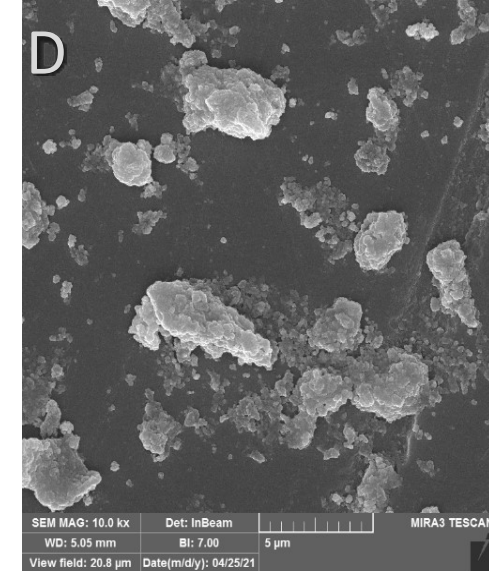
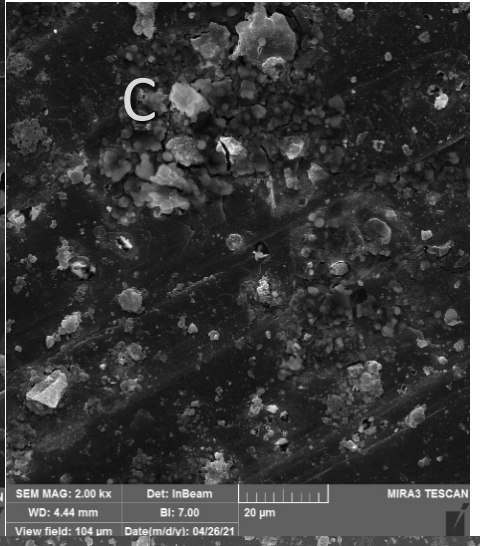
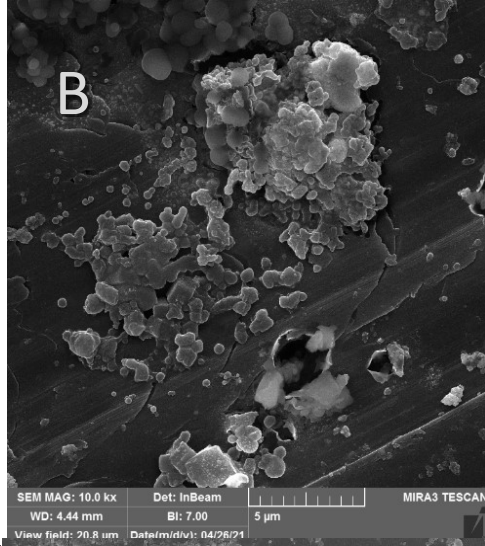
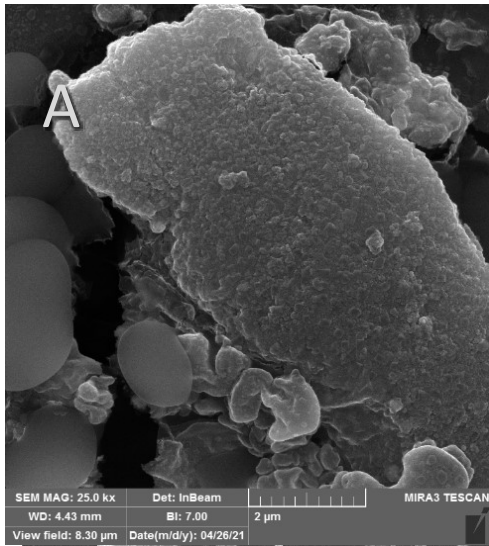
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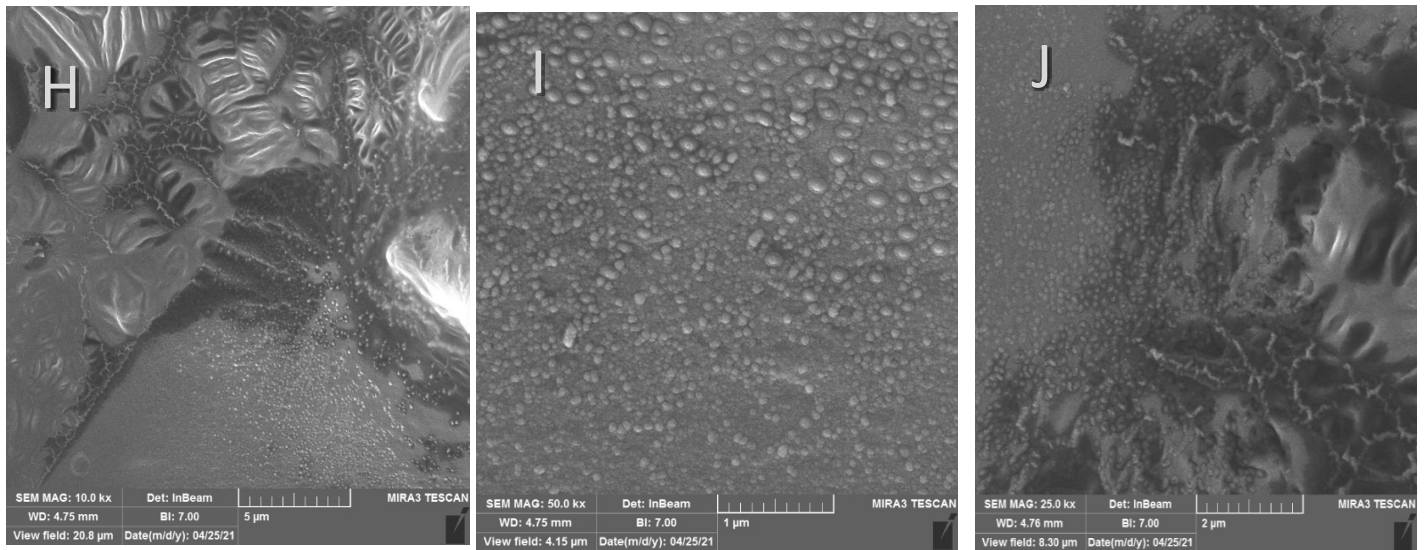
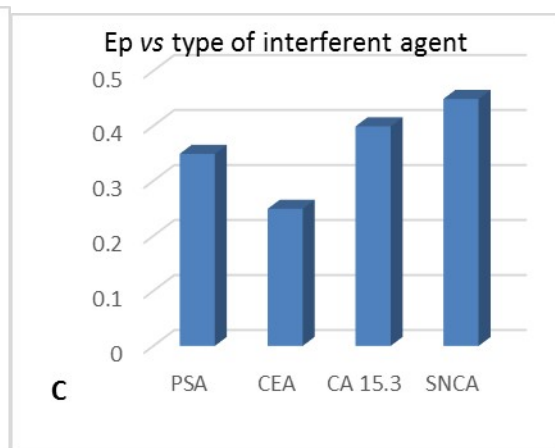
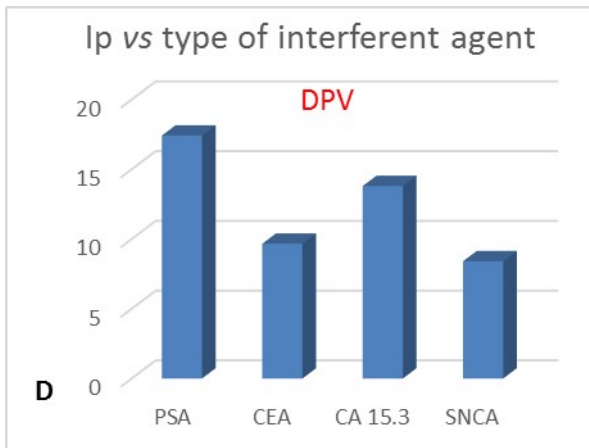
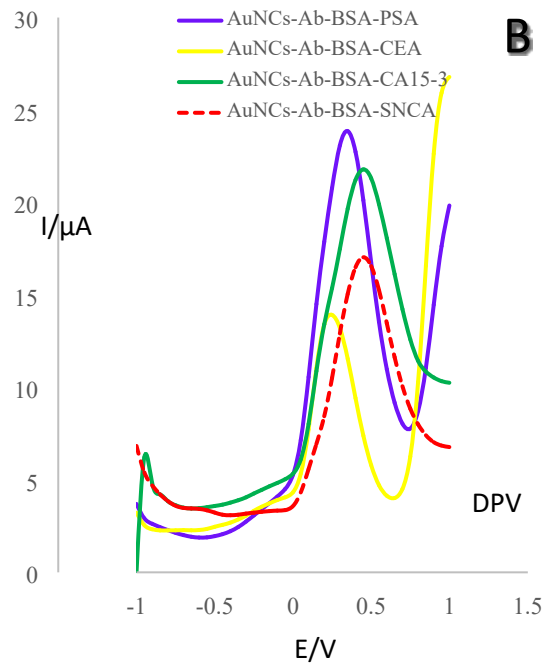
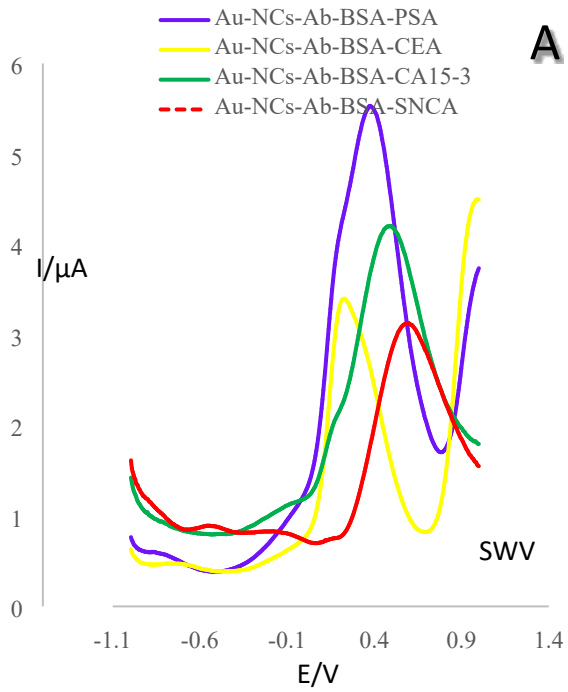


Figure S1. FE-SEM images of **A-C)** AuNCs, **D-F)** GC electrode/AuNCs, **G-J)** GC electrode/AuNCs/Ab and **K-M)** GC electrode/AuNCs/Ab/BSA/Antigen in different magnification.



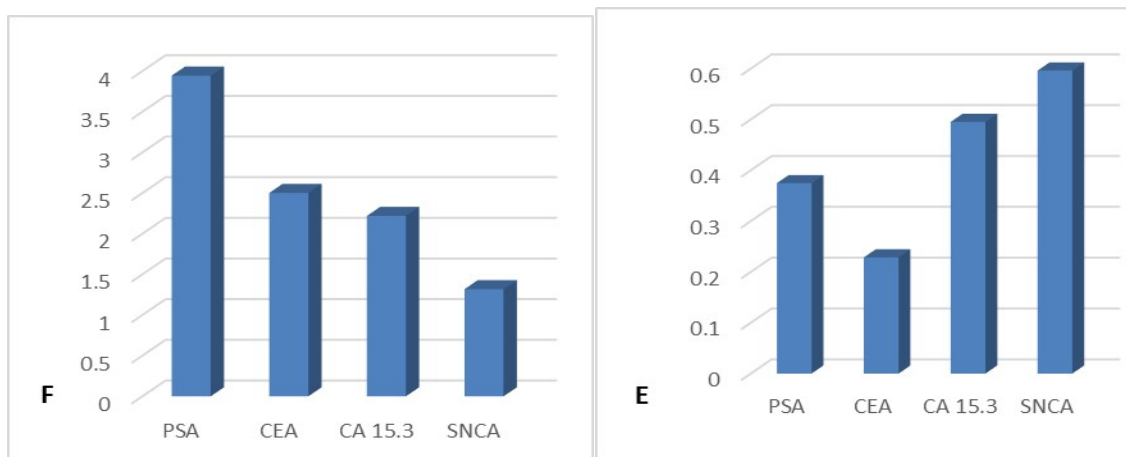
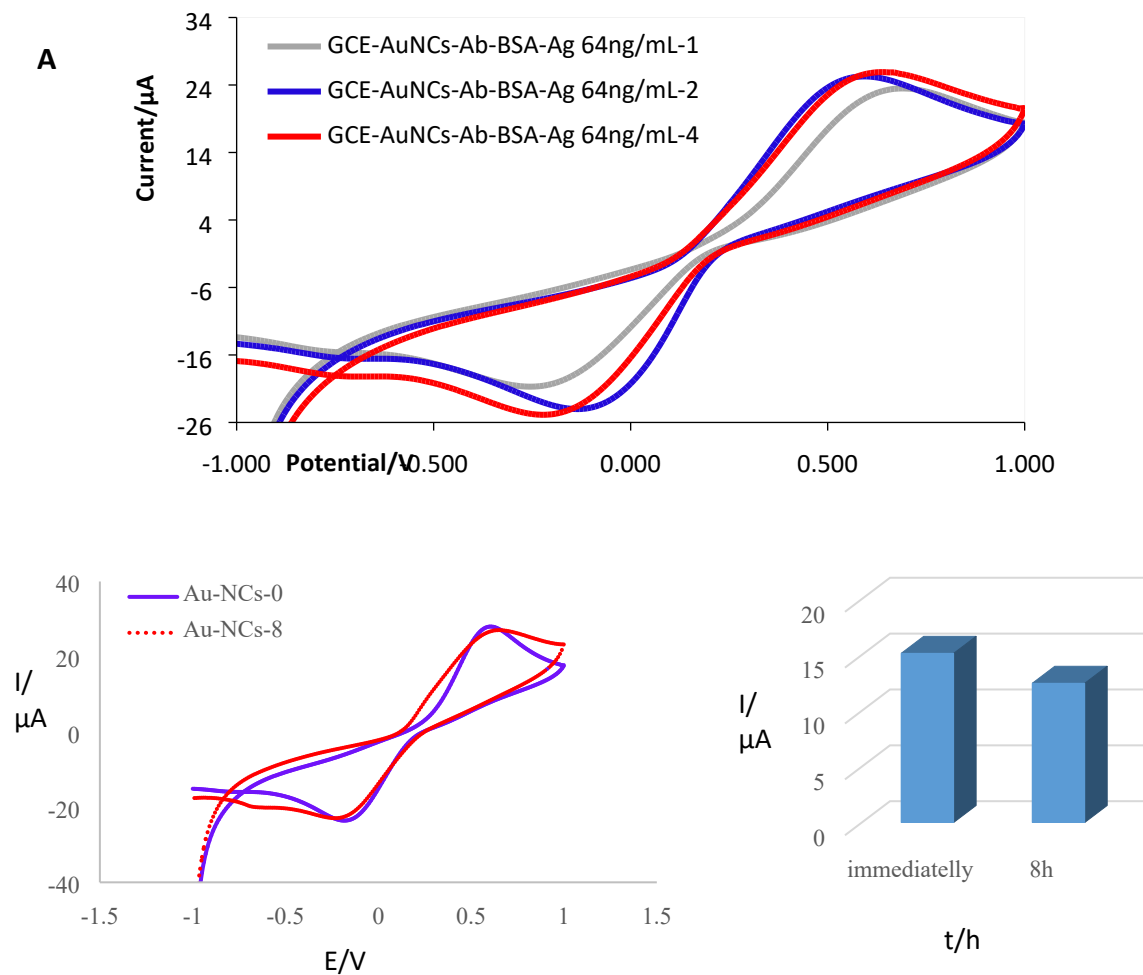


Figure S2. A) SWVs and **B)** DPVs of the designed immunosensor for hybridization by SNCA, CA 15.3, CEA and PSA in 0.01M $[\text{Fe}(\text{CN})_6]^{3-/4-}/\text{KCl}$ solution **C-F)** and related histograms. ($n=3$, $Sd=2.53$).



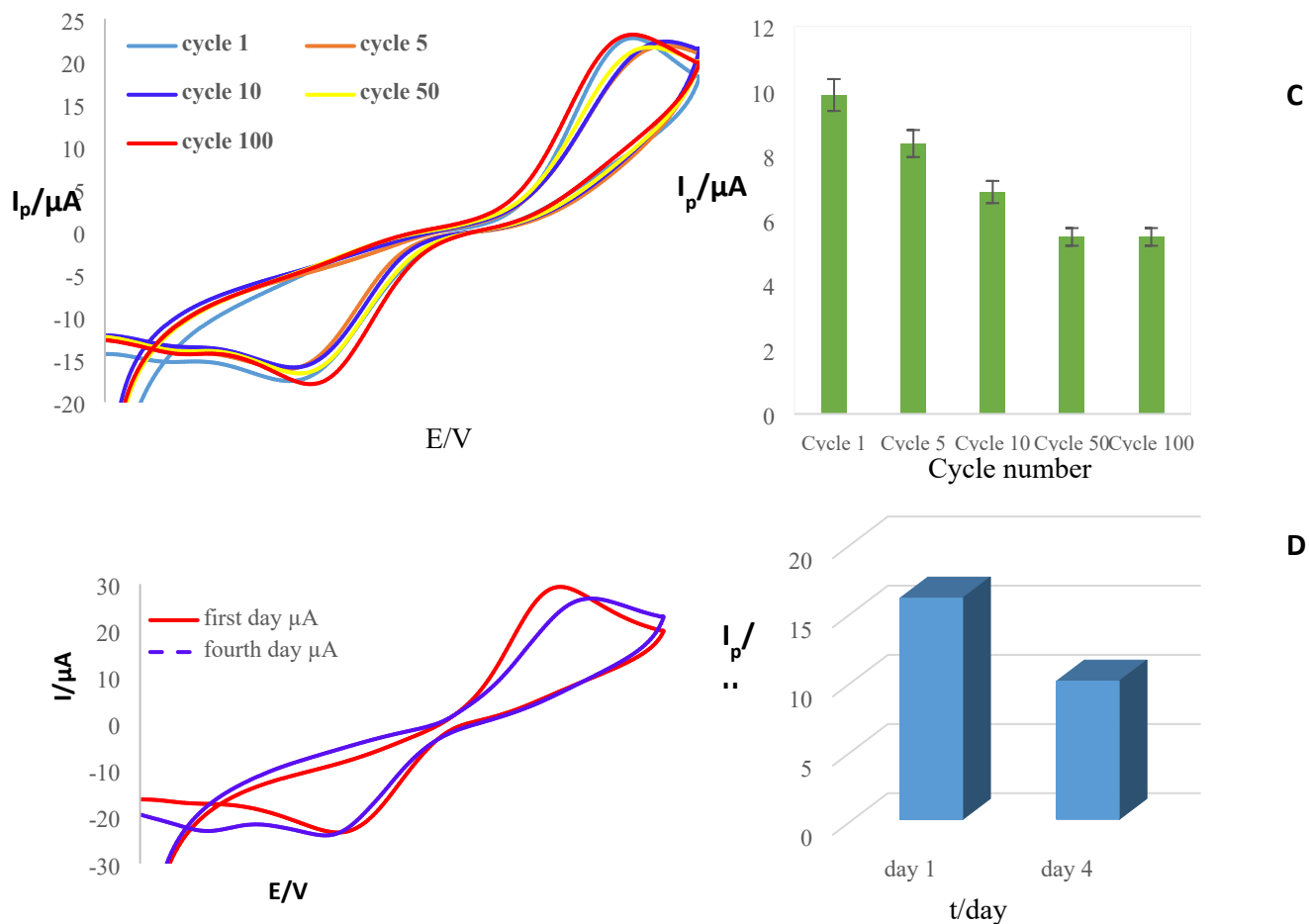


Figure S3. A) Reproducibility of immunosensor in 0.01M $[\text{Fe}(\text{CN})_6]^{3-/4-}/\text{KCl}$ solution (as supporting electrolyte) in a potential range of -1 to +1 V and scan rate of 100 mV/s. B) Inter-day stability of AuNCs on the surface of GC electrode. C) CVs of immunosensor in different cycle number and related histogram D) Intraday stability of Ab/AuNCs/GC electrode. ($n=3$, $Sd=2.85$).

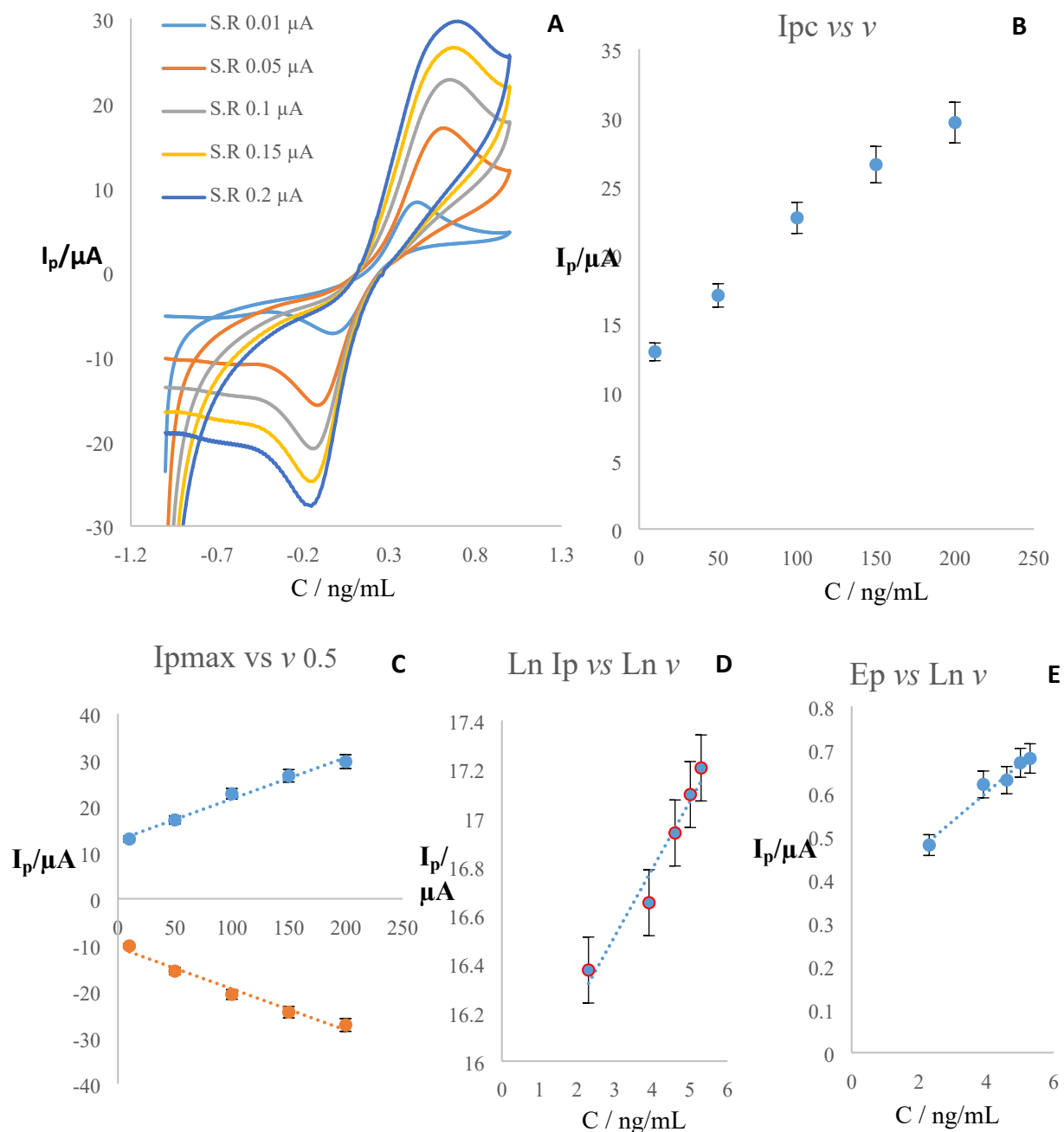


Figure S4. **A)** CVs of AuNCs/GC electrode in different scan rates (10, 50, 100, 150 and 200 mV/s). **B)** Variations of oxidation peak current versus different sweep rate. **C)** Calibration curve of oxidation peak current versus square root of different sweep rate. **D)** Calibration curve of Neperian logarithm of oxidation peak current versus Neperian logarithm of scan rate. **E)** Calibration curve of reduction peak position versus sweep rate in term of Neperian logarithm.