Screen-printed cells CS-4.0

Pretreatment: only rinse with distilled water and stir for 1 min in the probe solution before measurement.

Probe solution: K₃Fe(CN)₆ 5mM/KCl 0.1M, pH 7

 $E_{\text{start}} = -0.8 \text{ V}$; $E_{\text{end}} = +0.8 \text{ V}$

scan rates range: 0.01-1 V s⁻¹

Active area: 3.47(2) mm² (computed by Randles–Sevick's equation)



Pretreatment: wash with ethanol, rinse with distilled water and stir for 1 min in the probe solution before measurement.

Probe solution: K₃Fe(CN)₆ 5mM/KCl 0.1M, pH 7

 $E_{\text{start}} = -0.8 \text{ V}; E_{\text{end}} = +0.8 \text{ V}$

scan rates range: 0.01-1 V s⁻¹

Active area: 3.1(5) mm² (computed by Randles–Sevick's equation)



✓ Repeatability and reproducibility tests

Pretreatment: only rinse with distilled water and stir for 1 min in the probe solution before measurement.

Probe solution: $K_3Fe(CN)_6$ 5mM/KCl 0.1M, pH 7 $E_{start} = -0.8 V$; $E_{end} = +0.8 V$

scan rate: 0.1 V s⁻¹

- * 3 replicates of 10 CV scans with the same screen-printed cell
 - 1st replicate: $I_{pA} = 32(1) \ \mu A$; $I_{pC} = -41(1) \ \mu A$ (*n* = 10 scans)
 - \circ 2nd replicate: $I_{pA} = 34.1(5) \ \mu A$; $I_{pC} = -41.7(6) \ \mu A$ (*n* = 10 scans)
 - 3^{rd} replicate: $I_{pA} = 34.8(4) \ \mu A$; $I_{pC} = -42.2(6) \ \mu A$ (*n* = 10 scans)





✓ Example of application: DPV for Dopamine (DA) detection Pretreatment: only rinse with distilled water and stir for 1 min in the solution before measurement. Electrolyte solution: 10 mL of PBS 0.1 M/KCl 0.1 M at pH 7 DPV parameters: E_{start} = -0.3 V; E_{end} = +0.6 V; E_{step} = 0.01 V; E_{pulse} = 0.025 V; t_{pulse} = 0.2 s;

scan speed = 0.02 V/s.

3 repetitions for each concentration, stirring 20 s before registering the voltammogram



Calibration curve: $I_p [\mu A] = 0.01252(1) \cdot C_{DA} [\mu M] + 0.176(4)$ R² = 0.9999 Detection limit = 0.8 μ M; Quantification limit = 2.4 μ M