

### Description

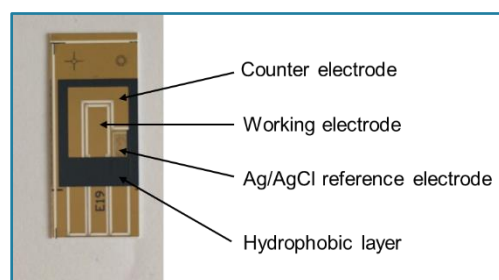
Part No. Au-5-001



Our thin-film planar gold electrodes are fabricated using high-resolution laser ablation of sputtered gold on a PET substrate with the cell volume delimited by screen-printed hydrophobic ink. Precision manufacture ensures excellent consistency and sensor-to-sensor reproducibility.

These are ideal for electrochemical research and analysis in environmental, clinical or agri-food areas and for electrochemical biosensor development.

With counter electrode, working electrode and screen-printed reference electrode made of Ag/AgCl, our electrodes are designed for robust electrochemistry research, with sensor-to-sensor reproducibility a key requirement.



### Applications

Key properties of these thin-film electrodes include their low-cost, high-resolution fabrication, disposability, and low reagent consumption. This results in an easy-to-use base for multiple research applications:

#### Biosensors

- Electrochemical transducers
- Environmental sensors
- Point-of-care tests

#### Electrochemistry

- Electrochemical research/analysis
- Student research projects/practicals
- *In vivo* diagnostics research

#### Flow systems & microfluidics

- Microfluidics research
- Flow systems
- Bioassays

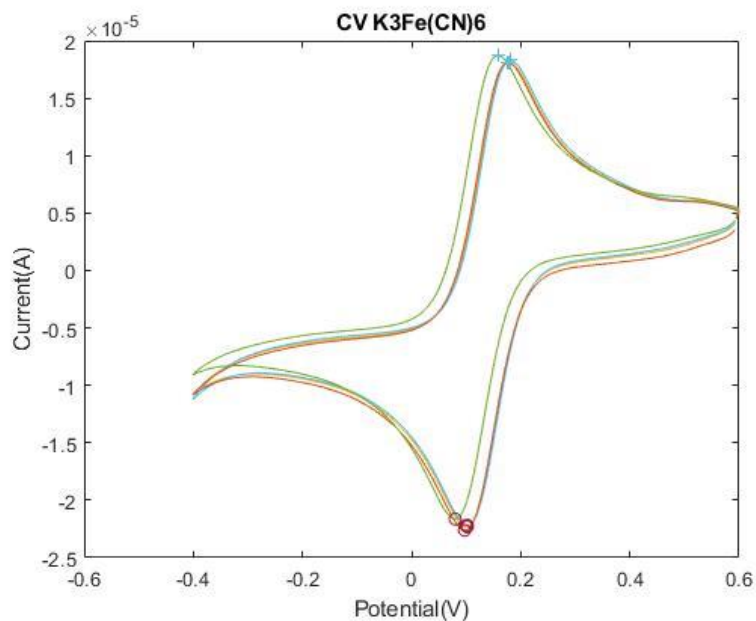
#### Nanotechnology

- Modified electrodes
- Nanostructure research
- Nanomaterial research

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## Technical data

## Reproducibility

Cyclic Voltammetry of Potassium Ferricyanide ( $K_3Fe(CN)_6$ )Fig.1: Cyclic voltammograms for 1 mM  $K_3Fe(CN)_6$ , for 5 different sensors.

## Sensitivity

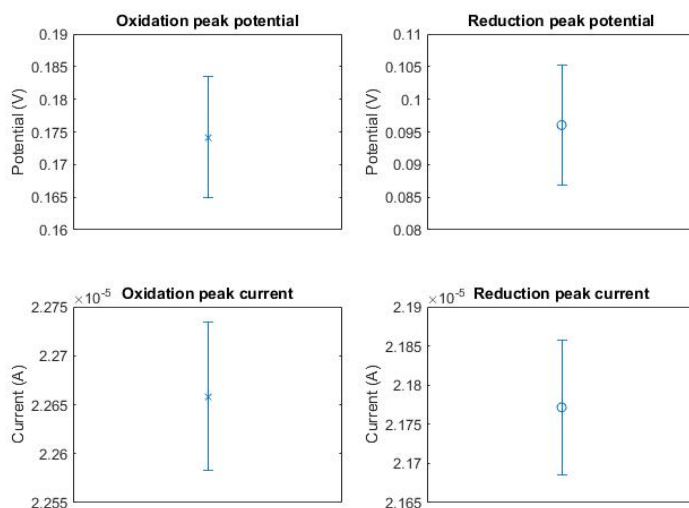


Fig.2: Oxidation (x) and reduction (o) peak potential and peak height variation for 5 sensors.

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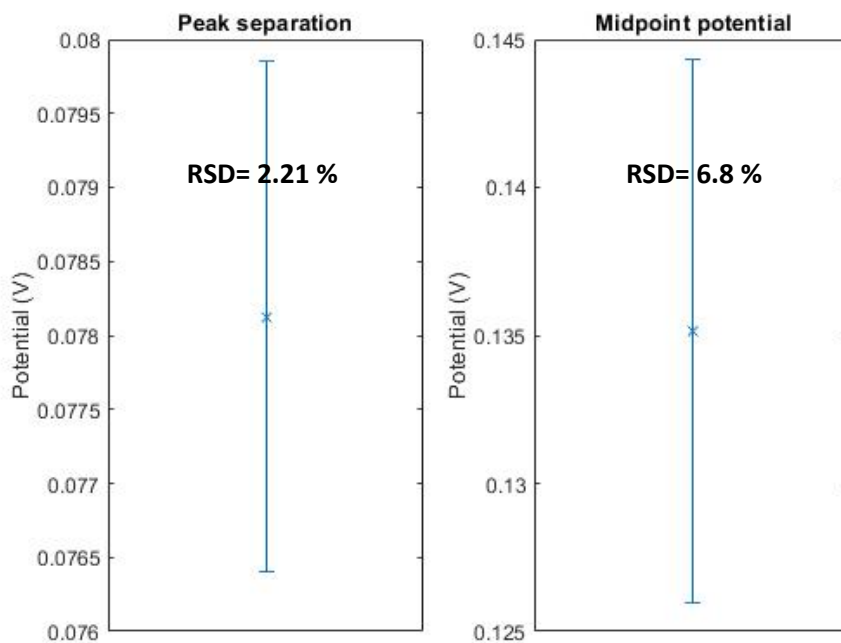


Fig.3: Average redox peak separation and equilibrium potential for 5 electrodes. (\*RSD= Relative Standard Deviation)

## Variable scan rate

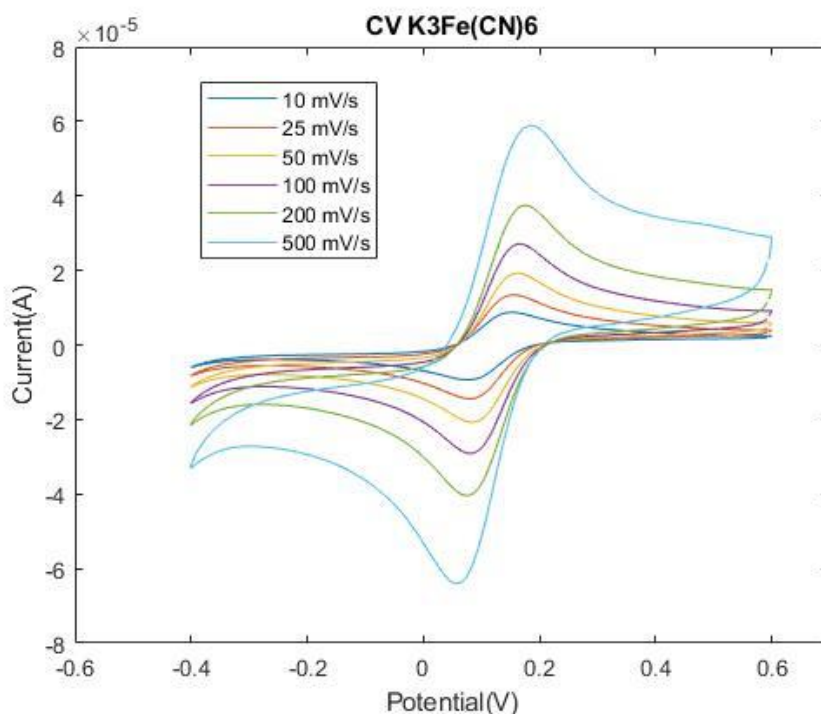


Fig.4: Cyclic voltammograms for 1 mM K<sub>3</sub>Fe(CN)<sub>6</sub>, for 1 sensor at different scan rates.

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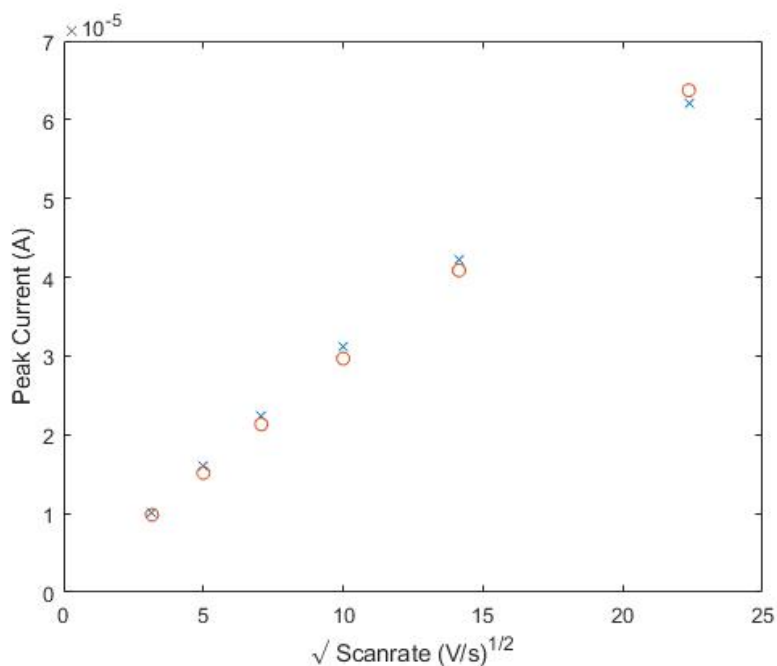


Fig.5: Peak height Vs  $\sqrt{\text{Scan rate}}$  for oxidation (x) and reduction (o) peaks.

### Cyclic Voltammetry of Potassium Ferricyanide ( $\text{K}_3\text{Fe}(\text{CN})_6$ )

#### Protocol

Pre-cleaning protocol:

- 1) Prepare 0.5M  $\text{H}_2\text{SO}_4$  (99.9%)
- 2) Dispense 150uL on the electrode
- 3) Perform a CV

CV settings

Scan rate: 100mV/s  
 Start E: 0V  
 Upper E: 1.3V  
 Lower E: -0.5  
 # of scans: 10

Potassium ferricyanide  $\text{K}_3\text{Fe}(\text{CN})_6$  protocol:

- 1) Prepare 1mM  $\text{K}_3\text{Fe}(\text{CN})_6$  & 10mM phosphate (pH=7) & 100mM KCl
- 2) Dispense 150uL on the electrode
- 3) Perform a CV

CV settings

Scan rate: 50mV/s (Variable)  
 Start E: 0.6V  
 Upper E: 0.6V  
 Lower E: -0.4V  
 # of scans: 3

Procedure:

- 1) Perform an acid clean followed by washing the electrode with distilled ultra-pure ( $18 \text{ M}\Omega \text{ cm}^{-1}$ ) water & dry with nitrogen gas
- 2) Once the electrode surface is dried perform a CV with potassium ferricyanide ( $\text{K}_3\text{Fe}(\text{CN})_6$ )
- 3) Repeat steps 1 and 2 for each electrode

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