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A Vitamin C Fuel Cell with a Non-bonded Cathodic Interface

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Table S1- Comparison of vitamin-C oxidation on different electrodes



Figure S1: Chronoamperometry curves on Pt, GC, and CoPc at an overpotential of 340 mV.

Table S2- Comparison of charge transfer resistance (R_{CT}) on different electrodes.

Electrode	R _{cT} (kohm.cm²)	
Carbon	7.778	
Platinum	2.477	
СоРс	0.102	



Figure S2- a) Vitamin C oxidation at different concentrations on CoPc modified carbon electrode in phosphate buffer of pH 7 at a scan rate of 10 mV/s, b) plot of peak current vs. concentration of vitamin C, c) scan rate dependence of vitamin C oxidation at 3 mM concentration on a CoPc electrode and d) plot of peak current vs. square root of scan rate, and e) plot of log (i) vs. log



(v).

Figure S3- Rotating disk electrode measurements for vitamin C (5 mM) oxidation on a CoPc modified glassy carbon disk electrode in phosphate buffer of pH = 7.

Table S3: Rate constants and number of electrons of Vitamin C oxidation on CoPc electrode from different methods.

Parameters	Method	Values
Number of electrons	Koutecky-Levich	2.0
Rate constant K ⁰ x10 ⁻⁴ (cm sec ⁻¹)		5.1



Figure S4- a) Cyclic voltammogram for the redox reaction of 10 mM ferricyanide (in phosphate buffer of pH = 7) on a glassy carbon based electrode at 10 mV/s scan rate, b) rotating disk electrode measurements for ferricyanide (10 mM in phosphate buffer of pH = 7) on a glassy carbon disk electrode, c) K-L plots at different potentials and d) plot of over potential vs. log of kinetic current.



Figure S5- a) Plot of oxidation and reduction peak currents for ferricyanide (10 mM) vs. square root of scan rate, b) plot of log (i) vs. log of scan rate, (c) cyclic voltammogram for the redox reaction of ferricyanide at different concentration and (d) corresponding peak current vs. concentration plot.

Table S4- Rate constants and number of electrons of ferricyanide on carbon electrode from different methods.



Figure S6. a) Cyclic voltammogram for ferricyanide acquired during UV-Visible spectroelectrochemistry, b) corresponding in-situ UV-Vis spectra during reductive scan and c) during oxidative scan. The spectrum of ferricyanide was taken as the background signal throughout the measurement.



Figure S7- a) Cell voltage at different discharge current densities and b) Stability of the CoPc electrode before and after long term stability tests.