Voltammetric Determination of Pb(II) by a Ca-MOF-Modified Carbon Paste Electrode Integrated in a 3D-Printed Device

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Figure S1. TGA curve of Pb-MOF obtained by immersing Ca-MOF in a PbCl₂ aqueous solution for 30 min, followed by filtration and drying over silica gel in a desiccator overnight.

Determination of the Active Area of the Electrode

The effective working area of the electrode was determined using cyclic voltammetric conditions (scan rate 0.05 V s^{-1}) for the one-electron reduction of K₃[Fe(CN)₆] [1.0 mmol L⁻¹ in 0.5 mol L⁻¹ KCl (Figure S2)] using the Randles-Sevcik equation.

$i_p = 0.446 nFAC (nFDv/RT)^{1/2}$

where i_p is the peak current (A), n (=1) is the number of electrons transferred, A is the effective area of the electrode(cm²), D is the diffusion coefficient of [Fe(CN)₆]³⁻ (taken to be 7.60 × 10⁻⁶cm²s⁻¹), C is the concentration (mol cm⁻³), v is the scan rate (Vs⁻¹), F is the Faraday constant (C mol⁻¹) R is the universal gas constant, T is the temperature in Kelvin.



Figure S2. Cyclic voltammogram of K_3 [Fe(CN)₆] (1.0 mmol L⁻¹ in 0.5 mol L⁻¹ KCl) at Ca-MOF/GP. Scan rate: 0.05 V s⁻¹



Figure S3. Potential stability of the 3D-printed carbon pseudo-reference electrode. The measurements were conducted in a solution containing 50 µg L⁻¹ Pb(II) in 0.1 mol L⁻¹ acetate buffer (pH 4.5)