



Analytical Methods

ARTICLE

Square wave voltammetry enables fast quantification and evaluation of Bi³⁺ extraction from eye shadow samples

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Electronic Supplementary Information

Table S1. Description of eye shadow samples used in this work.

Brand	Code	Color name	Tone	Origin country
A	1	Lead gray	Dark	Brazil
	2	Plum	Dark	
B	3	Carmine	Dark	China
	4	Light green	Light	
C	5	Purple	Dark	China
	6	Lilac	Dark	
D	7	Violet	Dark	Brazil
	8	Salmon	Light	
E	9	Red	Dark	Brazil
	10	Beige	Light	
F	11	Dark red	Dark	Brazil
	12	Pale green	Light	

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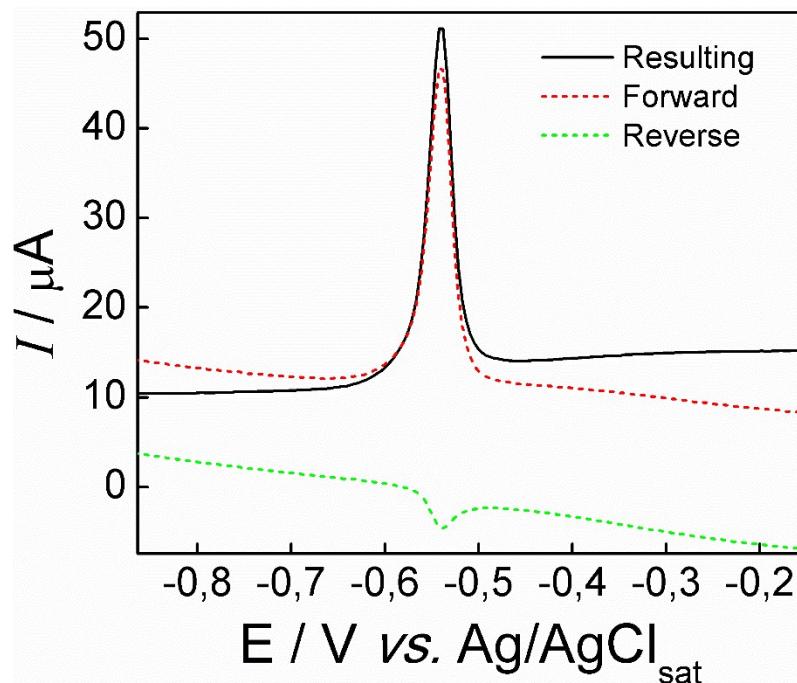


Figure S1. SW-ASV voltammograms recorded with GCE in NaOH 1.0 mol L⁻¹ in presence of 500 nmol L⁻¹ Bi³⁺. Voltammetric conditions: $E_{\text{dep}} = -1.3 \text{ V}$; $t_{\text{eq}} = 5 \text{ s}$; $f = 50 \text{ Hz}$; $\Delta E = 25 \text{ mV}$ and $\Delta E_s = 4 \text{ mV}$. Electrochemical cleaning at +0.5 V for 30 s.

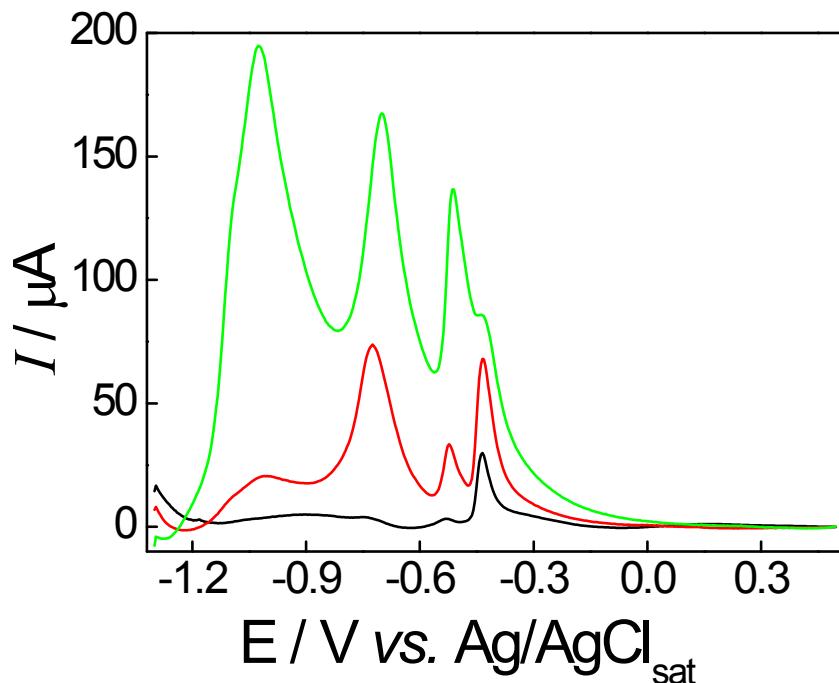


Figure S2. SW-ASV voltammograms recorded in $\text{NaOH} 1.0 \text{ mol L}^{-1}$ with CPE containing the immobilized particles of sample 2 with different deposition times. (—) 0; (—) 5 and (—) 180 s. Voltammetric conditions:

$E_{\text{dep}} = -1.3 \text{ V}$; $t_{\text{eq}} = 5 \text{ s}$; $f = 50 \text{ Hz}$; $\Delta E = 25 \text{ mV}$ and $\Delta E_s = 4 \text{ mV}$. Electrochemical cleaning at $+0.5 \text{ V}$ for 30 s.

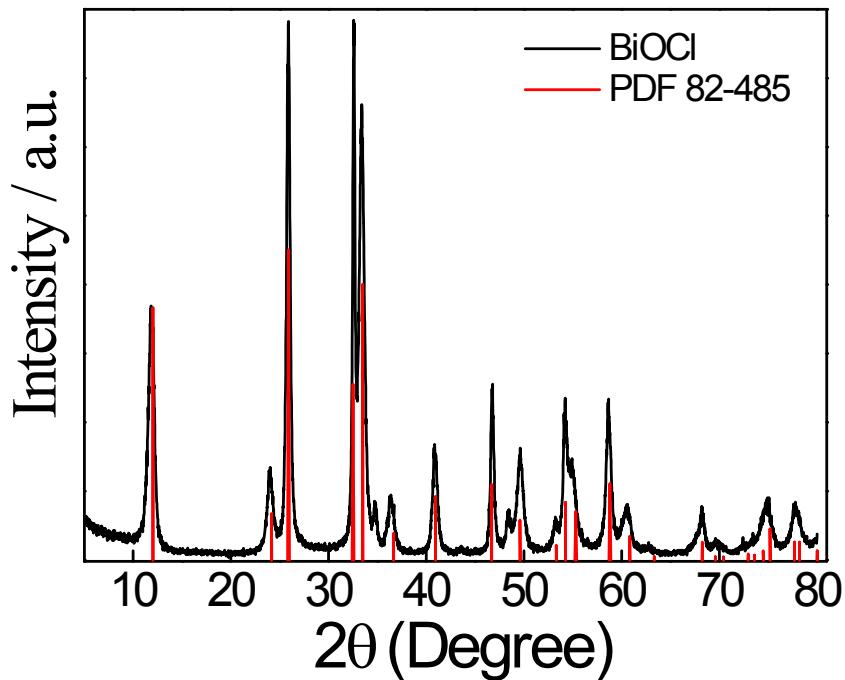


Figure S3. XRD pattern obtained for the synthesized BiOCl.

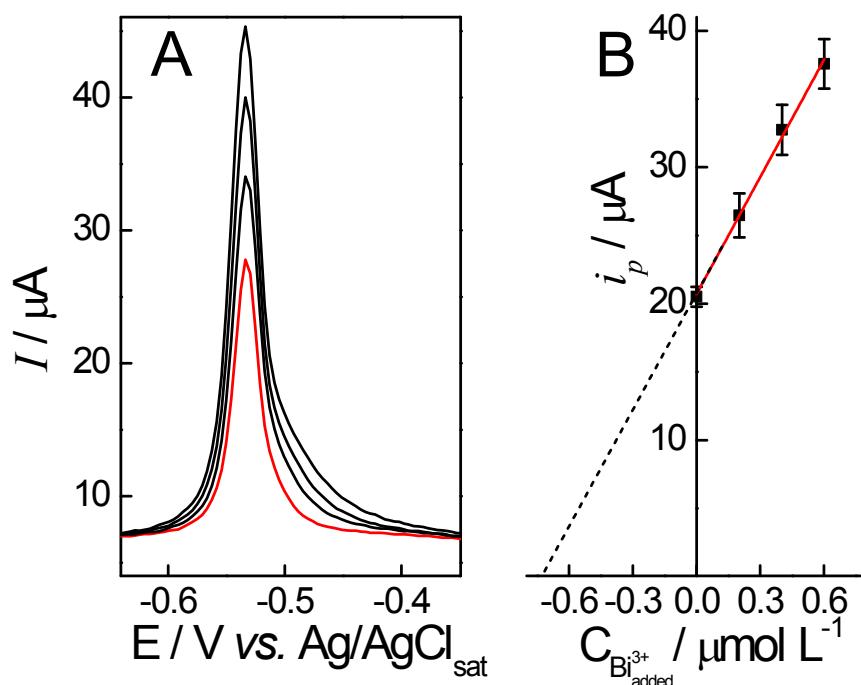


Figure S4. A) SW-ASV recorded with GCE in 1.0 mol L⁻¹ NaOH. (—) Aliquot of 10 μL of the BiOCl solution. (—) Successive standard additions of 0.2 μmol L⁻¹ Bi³⁺. Voltammetric conditions: $E_{\text{dep}} = -1.3$ V; $t_{\text{dep}} = 180$ s, $t_{\text{eq}} = 5$ s; $f = 50$ Hz; $\Delta E = 25$ mV and $\Delta E_s = 4$ mV. Electrochemical cleaning at +0.5 V for 30 s. B) Bi³⁺ standard addition curve.

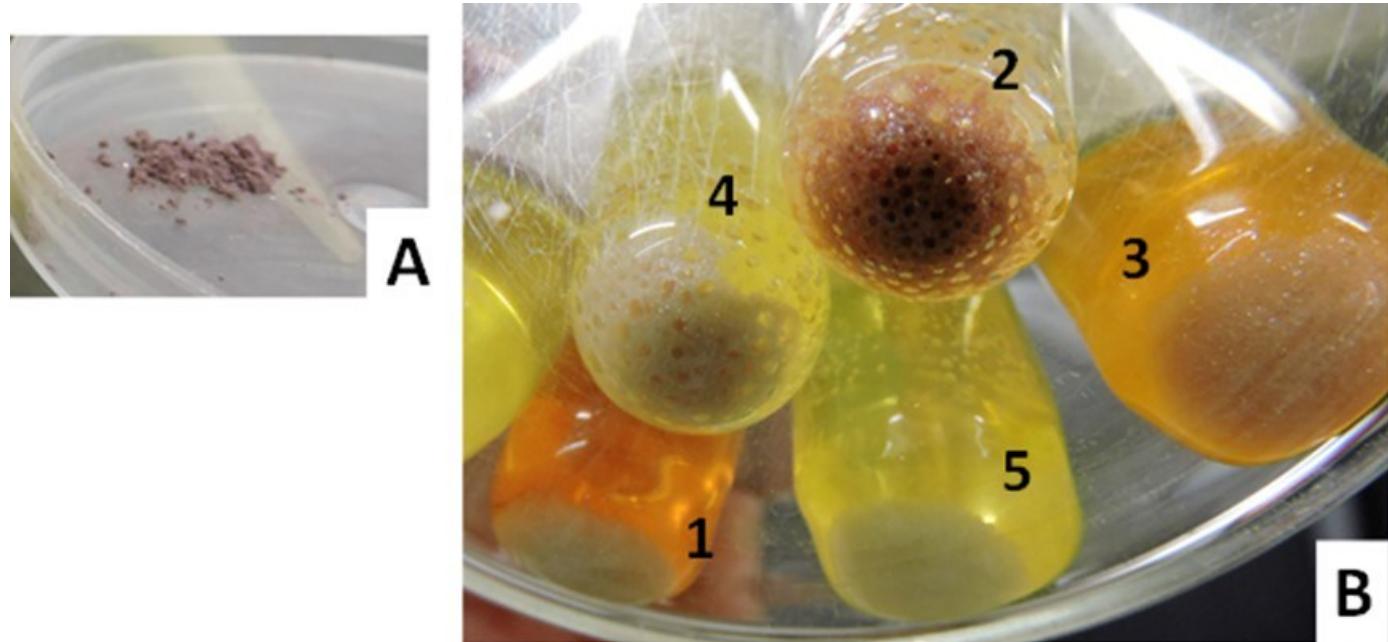


Figure S5. A) Image of the original sample 9. B) Images of the residues of sample 9 after the different sample pretreatment procedures.

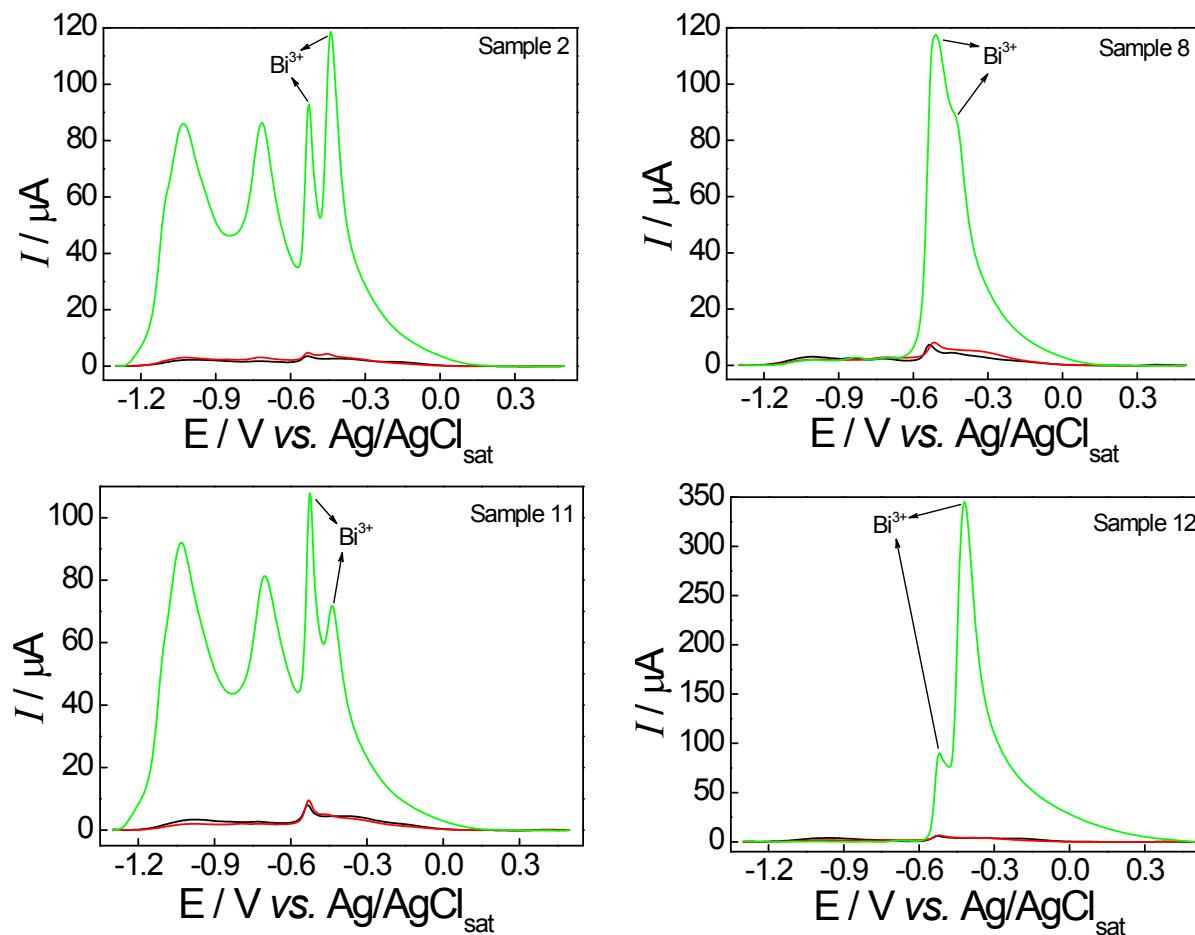


Figure S6. Baseline-corrected SW-ASV voltammograms recorded with CPE in NaOH 1.0 mol L⁻¹. (—) Without the immobilized particles. (—) After the immobilization of the particles of untreated samples. (—) After the immobilization of the particles of the samples treated with procedure 5*. Voltammetric conditions: $E_{dep} = -1.3$ V; $t_{eq} = 5$ s; $f = 50$ Hz; $\Delta E = 25$ mV and $\Delta E_s = 4$ mV. Electrochemical cleaning at +0.5 V for 30 s. Samples originally containing Bi³⁺.

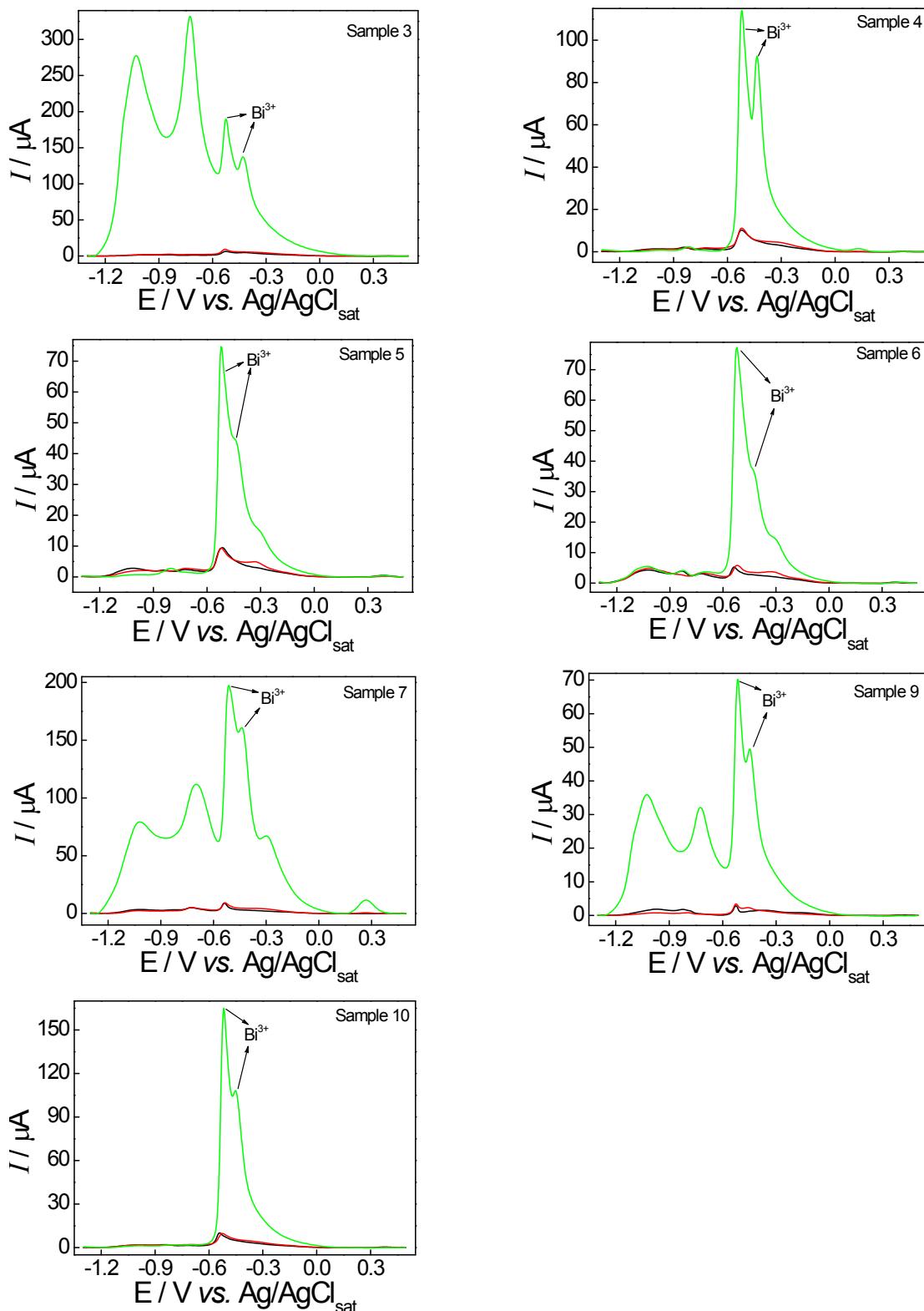


Figure S7. Baseline-corrected SW-ASV voltammograms recorded with CPE in $\text{NaOH } 1.0 \text{ mol L}^{-1}$. (—) Without the immobilized particles. (—) After the immobilization of the particles of the untreated samples. (—) After the immobilization of the particles of the samples treated with procedure 5*. Voltammetric conditions: $E_{\text{dep}} = -1.3 \text{ V}$; $t_{\text{eq}} = 5 \text{ s}$; $f = 50 \text{ Hz}$; $\Delta E = 25 \text{ mV}$ and $\Delta E_s = 4 \text{ mV}$. Electrochemical cleaning at $+0.5 \text{ V}$ for 30 s. Samples spiked with known amounts of Bi^{3+} .



Figure S8. Images of the original solid eye shadow samples and the respective residues from sample pretreatment with procedure 5*.

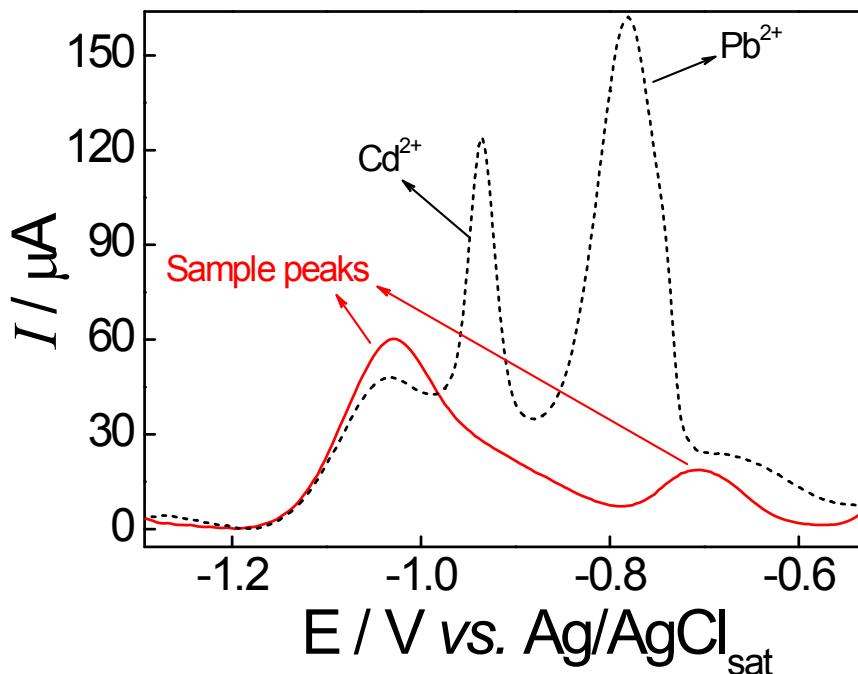


Figure S9. SW-ASV voltammograms recorded with GCE in NaOH 1.0 mol L⁻¹. (—) 40 µL of the supernatant from procedure 5* (sample 3). (---) 40 µL of the supernatant from procedure 5* (sample 3) + 36 µmol L⁻¹ Cd²⁺ and 10 µmol L⁻¹ Pb²⁺. Voltammetric conditions: $E_{\text{dep}} = -1.3$ V; $t_{\text{dep}} = 180$ s; $t_{\text{eq}} = 5$ s; $f = 50$ Hz; $\Delta E = 25$ mV and $\Delta E_s = 4$ mV. Electrochemical cleaning at +0.5 V for 30 s.

*Procedure 5 consisted in treat the eye shadow samples with 1.5 mL of concentrated HCl + 10 min in an ultrasound bath followed by addition of 0.5 mL of H₂O₂ (30 % v:v) + 5 min in an ultrasound bath.