

Supplementary Materials: Assessing the Potential of Co-Pt Bronze for Electrocatalysis in Acidic Media

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Table S1. Experimental set-ups for ORR test and accelerated durability test.

Component	Detail
Cell	Three electrode cell
Working electrode	Catalyst deposited on Glassy carbon
Counter electrode	Au mesh connected with a salt bridge
Reference electrode	RHE
Electrolyte	0.1 M HClO ₄ (Kanto Chemical, Ultrapure) 310 mL for OER and ORR measurements With 48 ppm Cl ⁻ (by addition of 35 μL of 12M HCl) for potential cycles
Temperature	30 °C for OER and ORR measurement 60 °C for potential cycles

Table S2. Experimental procedures for OER test.

Procedure	Details
1. Electrolyte deaeration	Ar bubbling (30 min)
2. Cyclic Voltammetry	Potential cycling 0.05–1.7 V, 50 mV s ⁻¹ , 400 rpm, 5 cycles
3 Potential step and holding	0.4 V for 10s to 1.7 V for 60 s, 400 rpm

Table S3. Experimental procedures for ORR test.

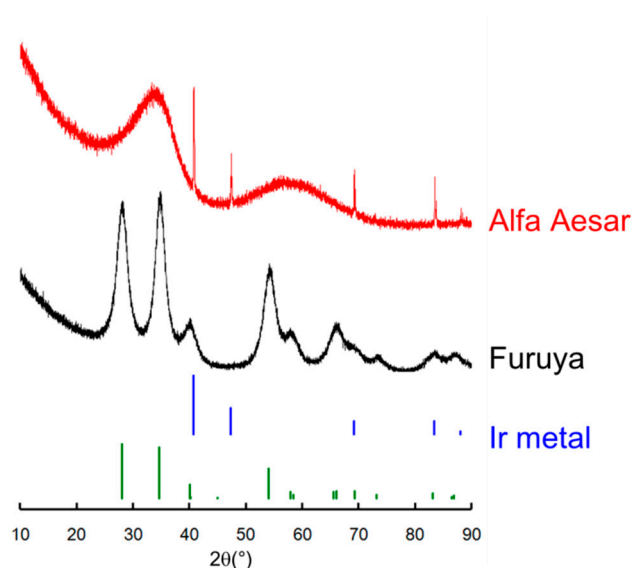
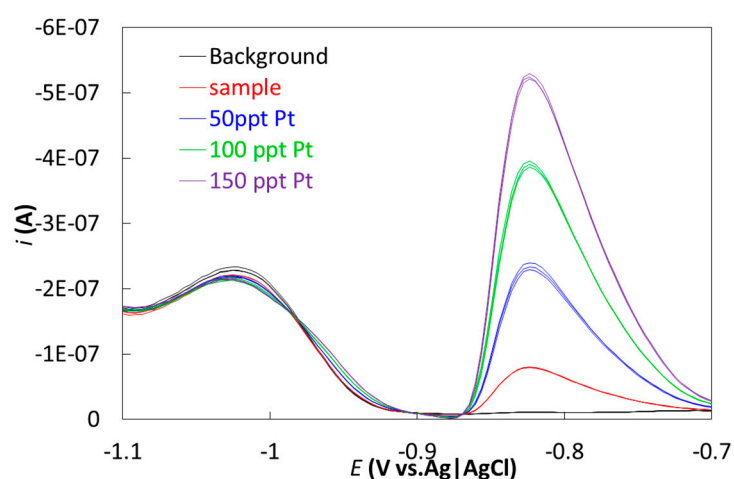
Procedure	Details
1. Electrolyte deaeration	Ar bubbling (30 min)
2. Electrode cleaning	Potential cycling 0.05–1.2 V, 500 mV s ⁻¹ , 100 cycles
3. ECSA determination	Potential cycling 0.05–1.0 V, 50 mV s ⁻¹ , 5 cycles
4. Background voltammetry	0.05–1.0 V, 10 mV s ⁻¹ , 3 cycles
5. O ₂ saturation	O ₂ bubbling (30 min)
6. ORR measurement	0.05–1.0 V, 10 mV s ⁻¹ , 400 rpm, 1 cycle

Table S4. Experimental procedures for accelerated durability test.

Procedure	Details
1. ORR test	Same condition with Table S2
2. Electrolyte change	adding 12 M HCl, 35 μL
3. ORR measurement	in the electrolyte containing 48 ppm Cl ⁻
4. Electrolyte deaeration	Ar bubbling (30 min)
5. ECSA measurement	0.05–1.0 V, 50 mV s ⁻¹ , 5 cycles
6. Potential step cycle	at 60 °C: 0.4–1.0 V each 3 s hold, 400 rpm
7. ECSA measurement	0.05–1.0 V, 50 mV s ⁻¹ , 5 cycles after 100, 200, 300, 400, 500, 600, 700, 800, 900, 1000, 2000, 3000, 4000, 5000, 6000, 7000, 8000, 9000 and 10,000 potential step cycles
8. Electrolyte sampling	(3 mL) for Pt ion concentration analysis after 0, 100, 200, 1000, 2000 and 10,000 Potential step cycles
9. ORR measurement	at 30 °C after O ₂ bubbling (30 min)
10. ECSA measurement	after Ar bubbling (30 min)
11. ORR measurement	after exchanging electrolytes to one without Cl ⁻

Table S5. Pt ion concentration determination.

Item	Detail
Apparatuses	Metrohm VA663, Metrohm IME663
Electrodes	WE: hanging mercury drop, CE: Glassy carbon, RE: Ag/AgCl/3 M KCl
Supporting Electrolyte	0.6 M sulfuric acid, 0.18 mM hydrazine and 6.6 mM formaldehyde mixture deaerated by nitrogen bubbling 10 min
Operational modes	Potential sweep mode Differential pulse mode (modulation time: 0.05 s, modulation amplitude: 0.05 V, Interval: 0.2 s, step potential: 0.004 V)
Procedures	Nitrogen bubbling: 10 s Mercury drop refresh: 5 drops Pre-condensation: at -0.7 V, stirrer 1500 rpm, 60 s Potential sweep 10 s after stirrer stop: -0.7 to -1.1 V
Signal Peak	at -0.83 V Pt: 50, 100 and 150 ppt. See Figure S2.
Calibration	Pt: 50, 100 and 150 ppt. See Figure S2.

**Figure S1.** XRD patterns of two commercial iridium oxides.**Figure S2.** Stripping voltammogram for Pt concentration determination.

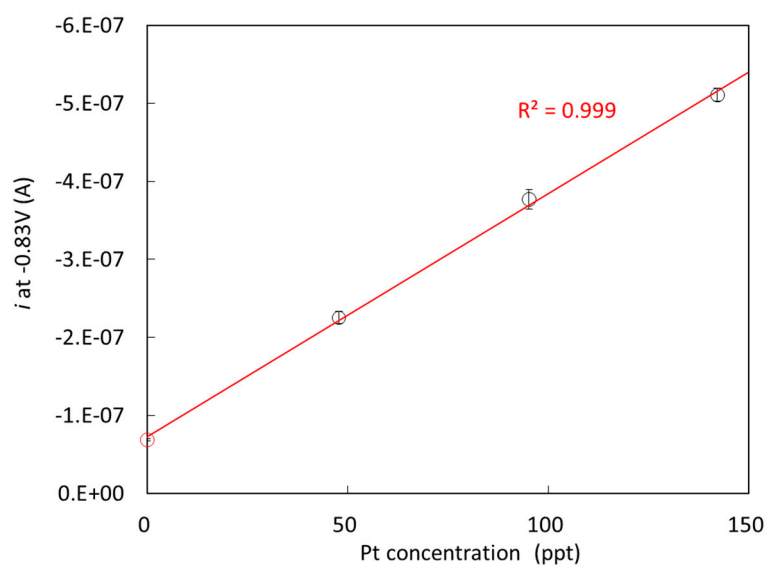


Figure S3. Calibration line for Pt concentration determination using the peak current at -0.83 V .

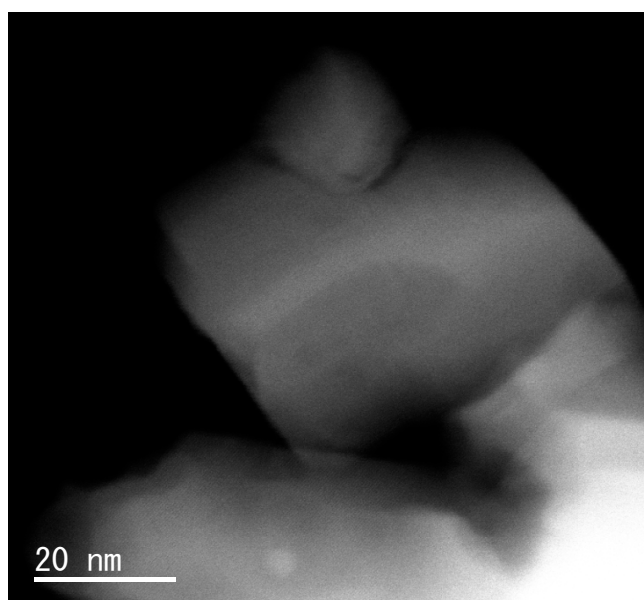


Figure S4. TEM image of Co-Pt bronze after ADT.