

Electronic Supplementary Information

**Changes induced by transition metal oxides in Pt nanoparticles unveil
the effects of electronic properties on oxygen reduction activity**

Felipe B. Ometto¹, Emilia A. Carbonio², Érico Teixeira-Neto^{3#}, Hebe M. Villullas^{1*}

*1 São Paulo State University (UNESP), Institute of Chemistry, Rua Prof. Francisco Degni, 55 -
14800-060 Araraquara, São Paulo, Brazil.*

*2 Helmholtz-Zentrum Berlin für Materialien und Energie, BESSY-II, Albert-Einstein-Straße 15,
12489, Berlin, Germany.*

*3 Brazilian Nanotechnology National Laboratory (LNNano), Brazilian Center for Research in
Energy and Materials (CNPEM), Campinas, São Paulo, Brazil.*

*Corresponding author: E-mail address:mercedes.villullas@unesp.br

#Present address: University of Campinas (Unicamp), Institute of Chemistry. Inomat – Bloco J, Cidade Universitária. 13083-970 Campinas, São Paulo, Brazil. CP 3154.

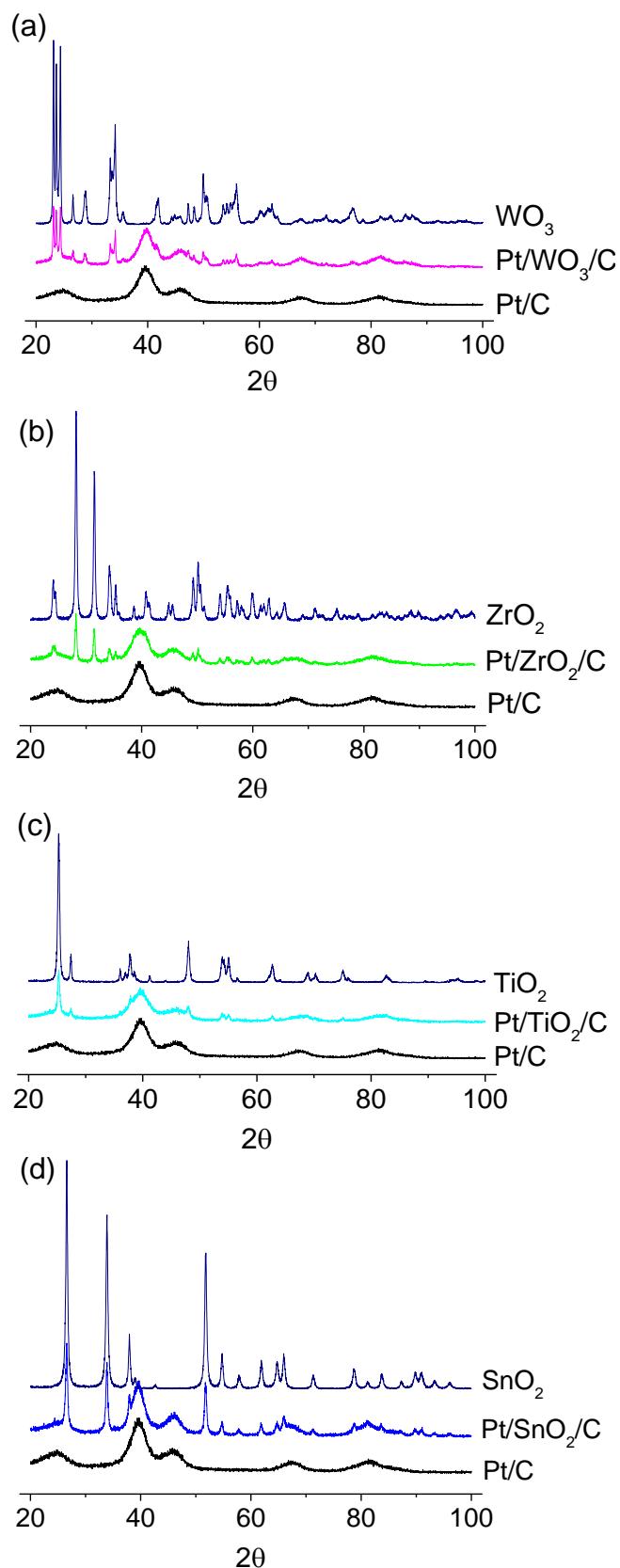


Fig. S1 Comparison of X-ray diffraction patterns of Pt/WO₃/C (a), Pt/ZrO₂/C (b), Pt/TiO₂/C (c) and Pt/SnO₂/C with those of Pt/C and the corresponding transition metal oxide.

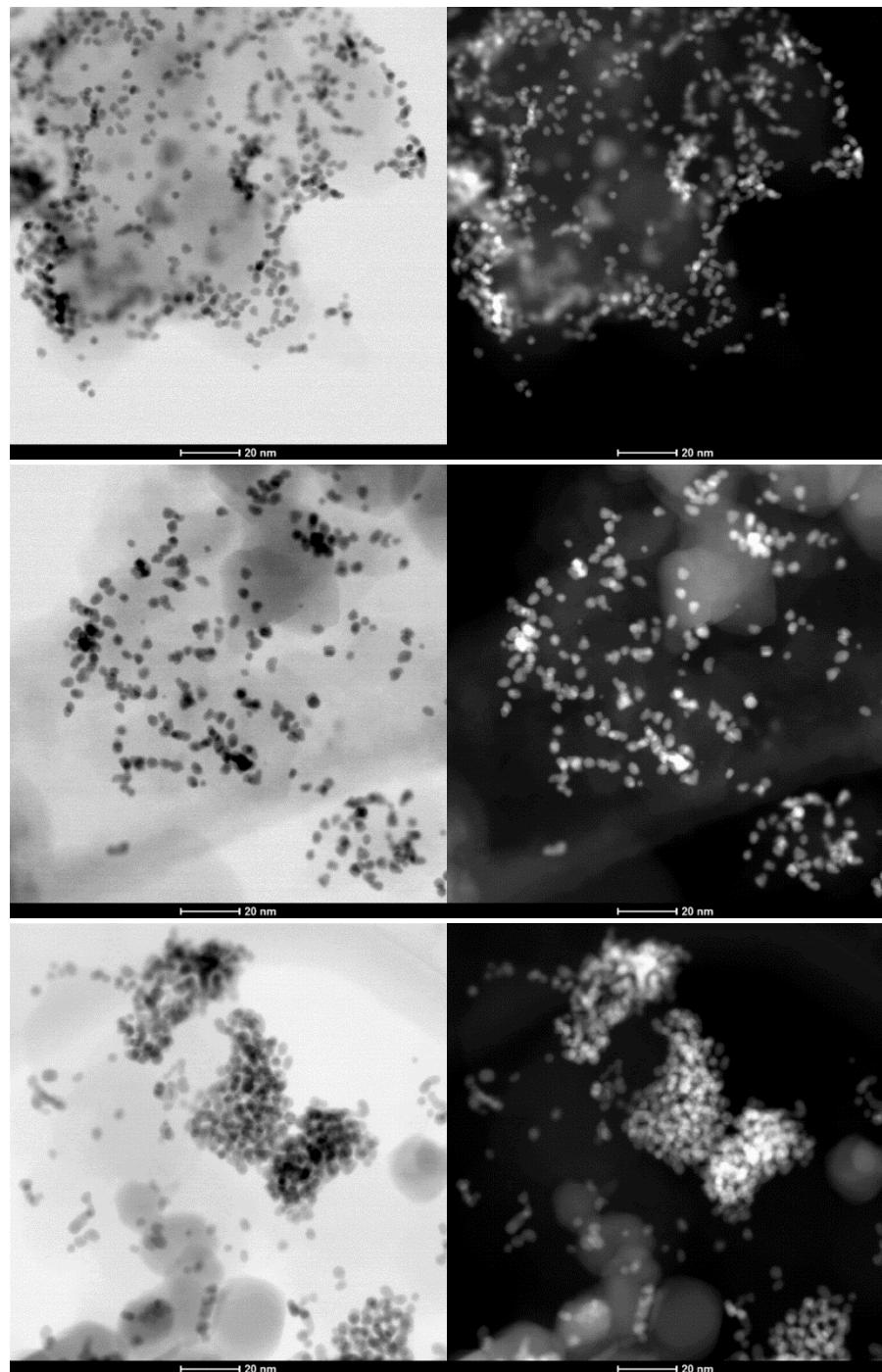


Fig. S2 HR-STEM images obtained in bright and dark field. From top to bottom: Pt/C, Pt/TiO₂/C and Pt/SnO₂/C.

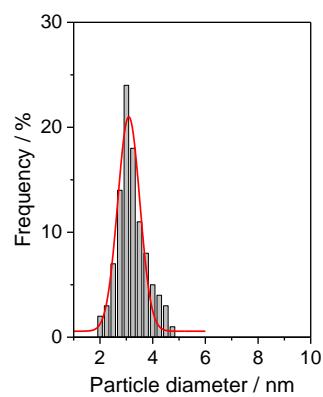


Fig. S3 Histogram of particle size distribution for Pt/C catalyst.

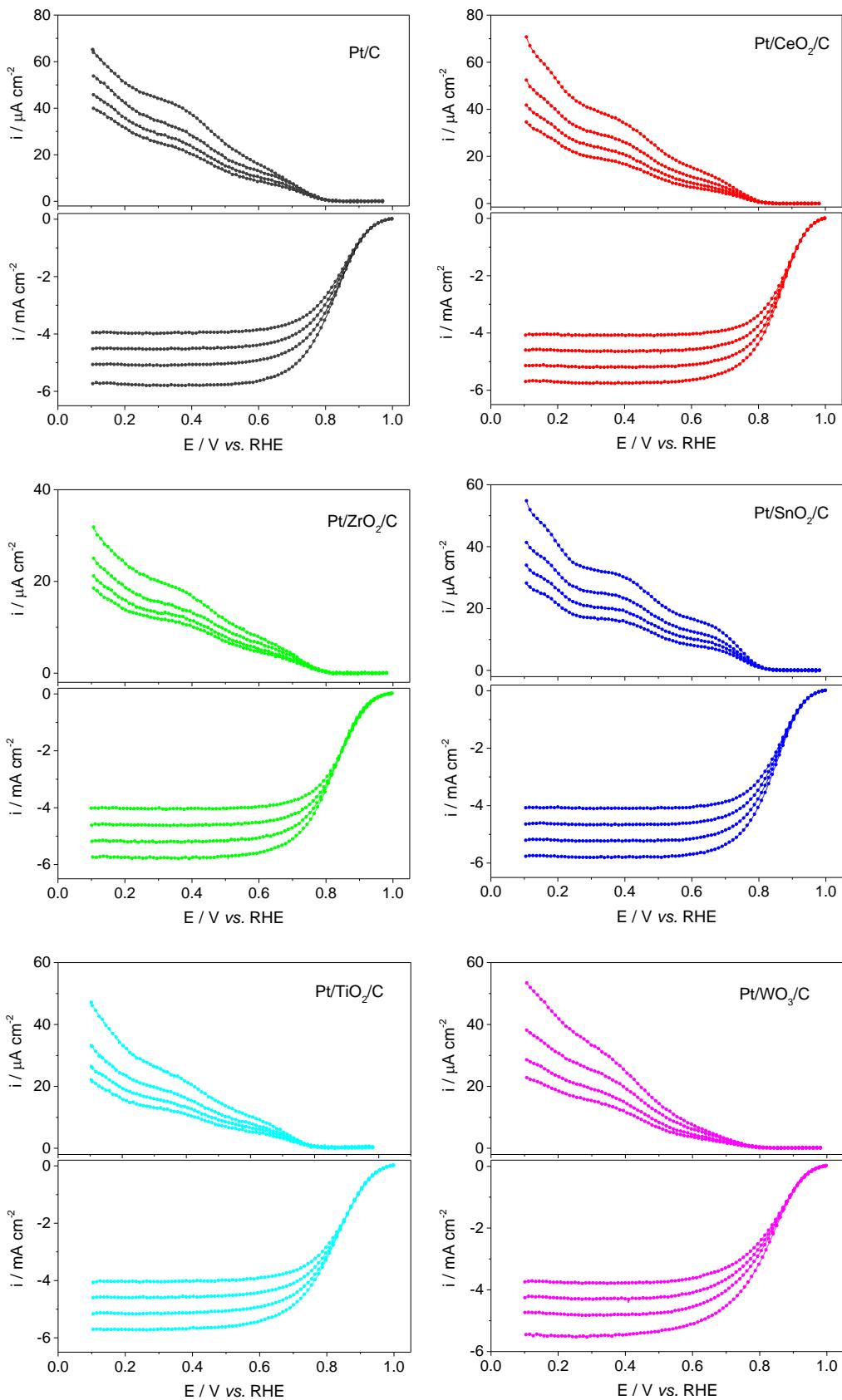


Fig. S4 Current densities of oxygen reduction and H_2O_2 oxidation measured in 0.5 M H_2SO_4 solution at rotation rates of 1225, 1600, 2025 and 2500 rpm. Currents are presented normalized by the geometric areas of the electrodes (0.247 cm^2 for disk and 0.187 cm^2 for ring). RRDE collection efficiency: 0.37.

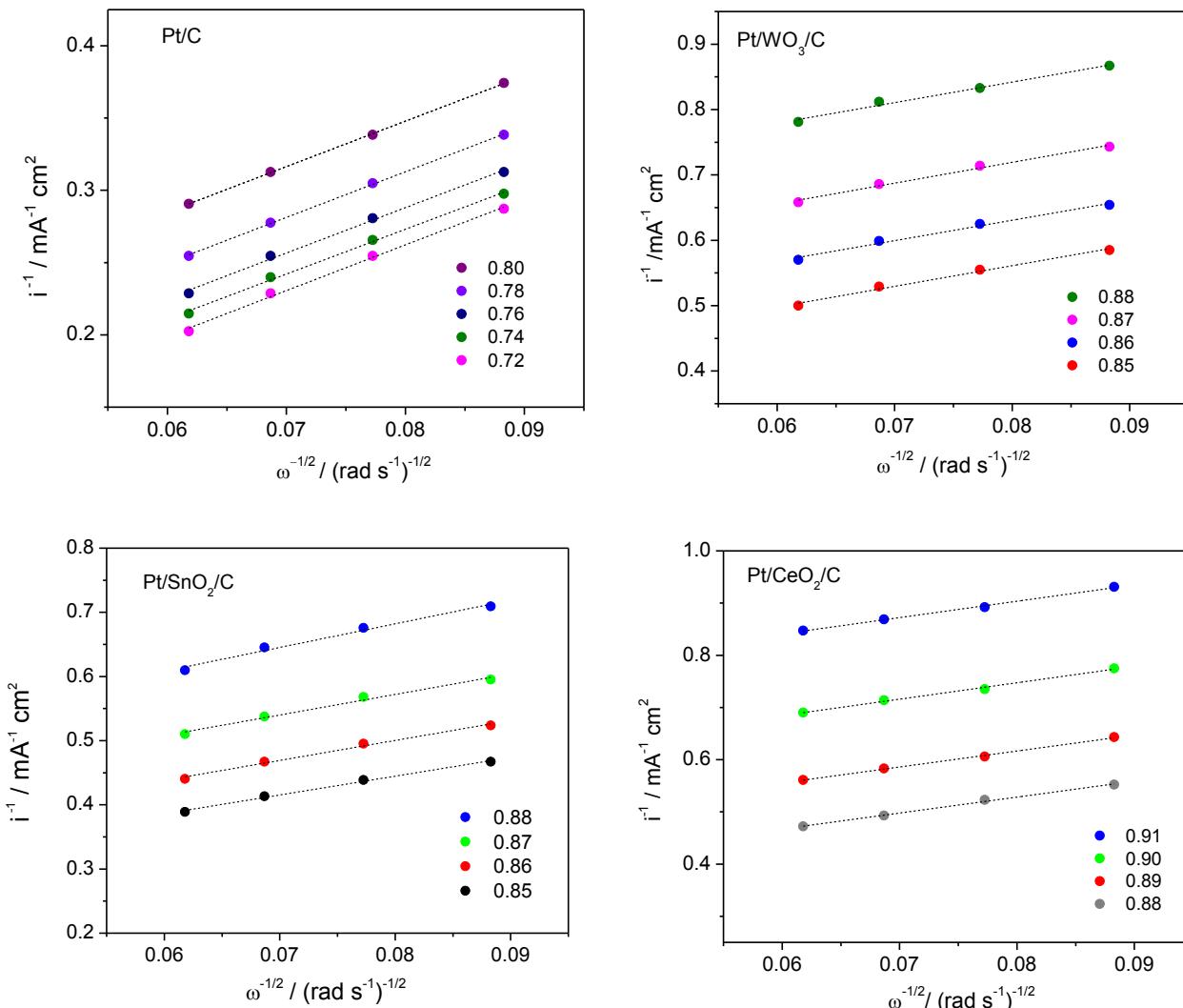


Fig. S5 Koutecky-Levich plots for Pt/C, Pt/WO₃/C, Pt/SnO₂/C and Pt/CeO₂/C catalysts constructed from disk currents at different potentials. Electrolyte: 0.5 M H₂SO₄. The number of electrons was estimated using Co= 1.1 10⁻⁶ g mol cm⁻³, D= 1.4 10⁻⁵ cm² s⁻¹ and v= 1.2 10⁻² cm² s⁻¹.

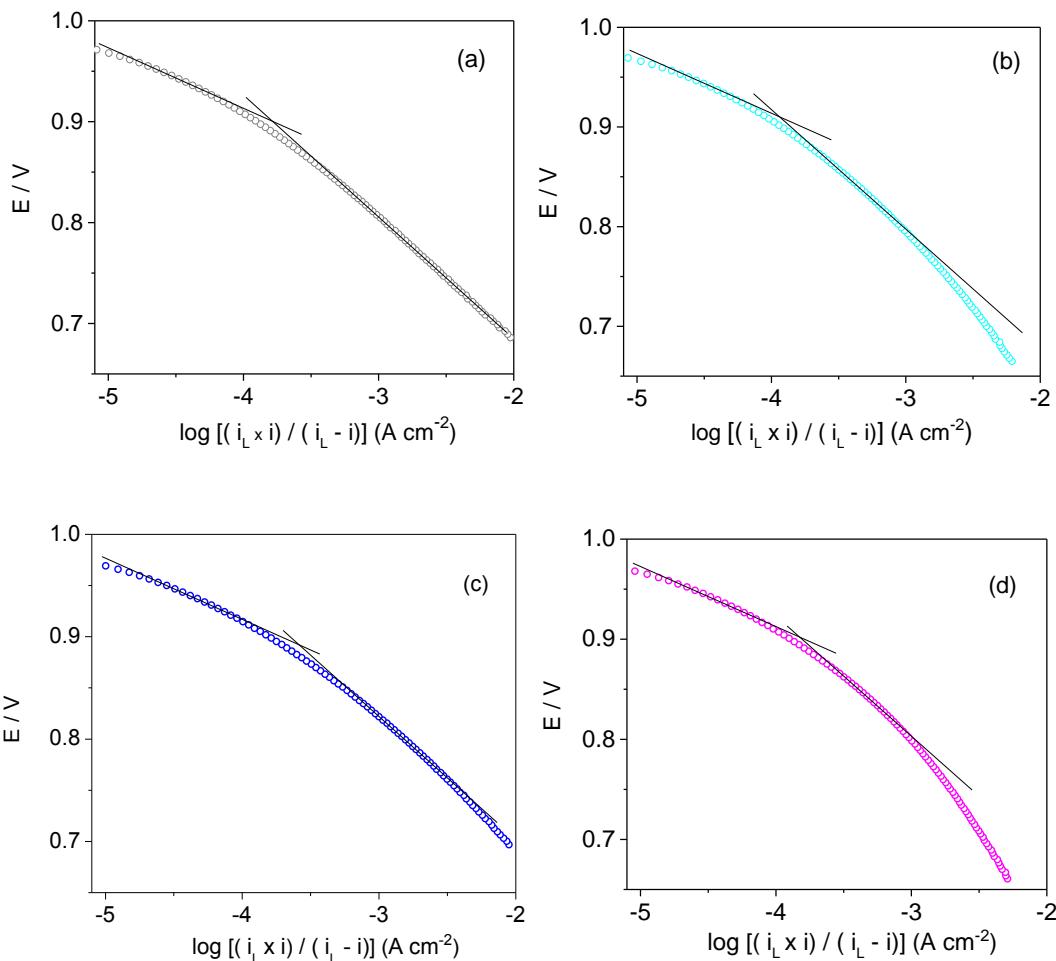


Fig. S6 Tafel plots for Pt/C (a), Pt/TiO₂/C (b), Pt/SnO₂/C (c) and Pt/WO₃/C (d). The lines correspond to -60 mV dec^{-1} and -120 mV dec^{-1} . Kinetic current densities obtained by normalization by Pt EAA.

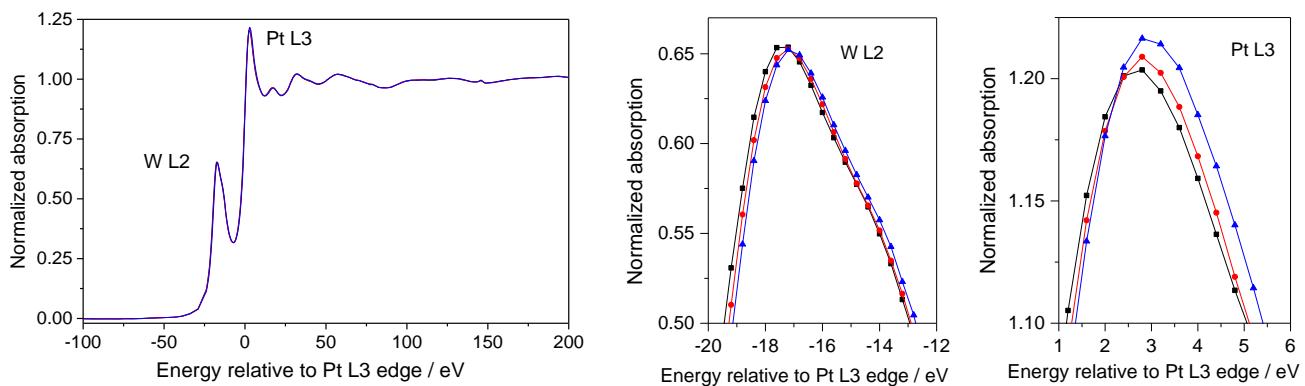


Fig. S7 Normalized *in situ* XAS spectra for Pt/WO₃/C obtained in 0.5 M H₂SO₄ and regions of maximum intensity enlarged. Applied potential: 0.55 V (black), 0.80 V (red) and 0.90 V (blue).

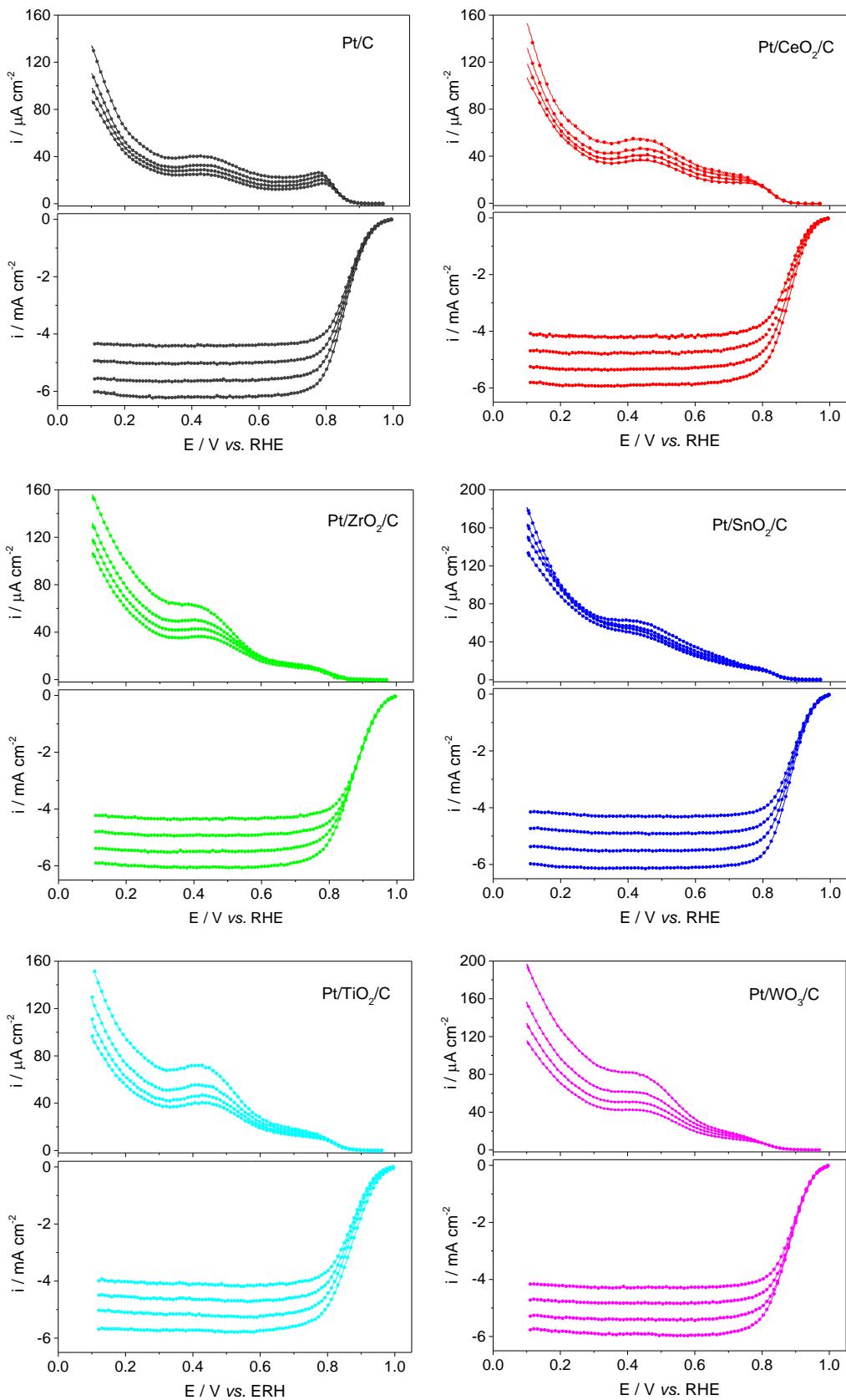


Fig. S8 Current densities of oxygen reduction and H_2O_2 oxidation measured in 0.1 M KOH solution at rotation rates of 1225, 1600, 2025 and 2500 rpm. Currents are presented normalized by the geometric areas of the electrodes (0.247 cm^2 for disk and 0.187 cm^2 for ring). RRDE collection efficiency: 0.37.

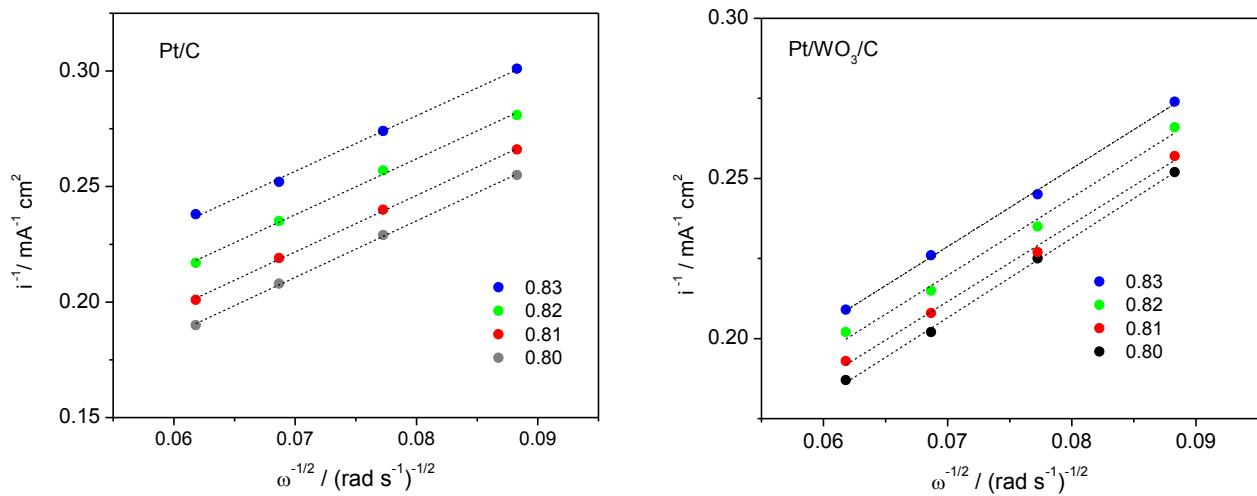


Fig. S9 Koutecky-Levich plots for Pt/C and Pt/WO₃/C catalysts constructed from disk currents at different potentials. Electrolyte: 0.1 M KOH. The number of electrons was estimated using Co= 1.15 10^{-6} g mol cm⁻³, D= 1.9 10^{-5} cm² s⁻¹ and v= 1.9 10^{-2} cm² s⁻¹.

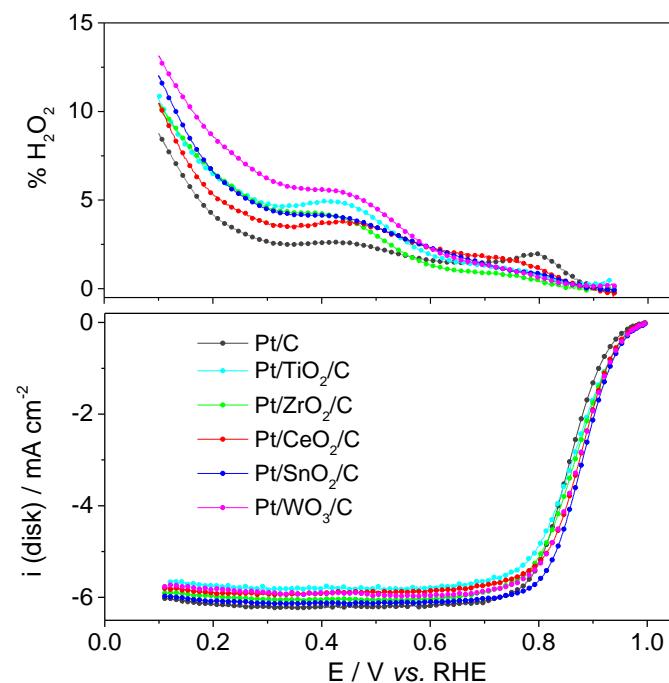


Fig. S10 ORR polarization curves measured at the disk electrode and H₂O₂ yield in 0.1 KOH solution Scan rate: 5 mV s⁻¹. Rotation rate: 2500 rpm.

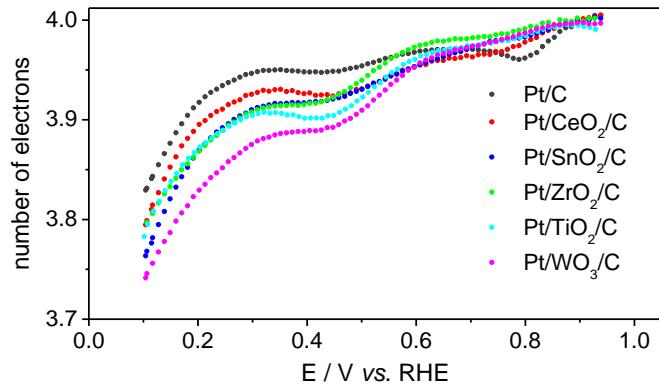


Fig. S11 Number of electrons calculated from RRDE currents measured in 0.1 M KOH solution as a function of potential.

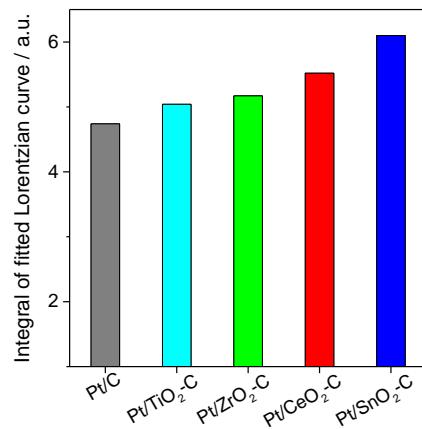


Fig. S12 Values of the integral of the Lorentzian curve adjusted to the XAS spectra measured *in situ* in 0.1 M KOH with applied potential of 0.90 V.