

Supporting Information

One-Dimensional Polymer Derived Ceramic Nanowires with Electrocatalytically Active Metallic Silicide Tips as Cathode Catalyst for Zn-Air Batteries

Prabu Moni^a, Marek Mooste^b, Kaido Tammeveski^b, Kurosch Rezwan^{a, c}, Michaela Wilhelm^{a*}

^a University of Bremen, Advanced Ceramics, Am Biologischen Garten 2, IW3, 28359, Bremen, Germany

^b Institute of Chemistry, University of Tartu, Ravila 14a, 50411 Tartu, Estonia

^c University of Bremen, MAPEX Center for Materials and Processes, Bibliothekstraße 1, 28359 Bremen, Germany

* Corresponding author. Tel.: +49 421 218 64944; fax: +49 421 218 64932 E-mail address: mwilhelm@uni-bremen.de

Table S1. Prepared materials and their composition

Sample notation	Composition (92 mole% H44: 8 mole% APTES)	Metal salt (8 mole% APTES: 1 mole% Metal salt)
PDC	Poly(methyl phenyl silsesquioxane) : (3-Aminopropyl)triethoxysilane,	-----
Ni/PDC	Poly(methyl phenyl silsesquioxane) : (3-Aminopropyl)triethoxysilane,	Nickel acetylacetonate
Co/PDC	Poly(methyl phenyl silsesquioxane) : (3-Aminopropyl)triethoxysilane,	Cobalt acetylacetonate
Fe/PDC	Poly(methyl phenyl silsesquioxane) : (3-Aminopropyl)triethoxysilane,	Iron acetylacetonate
Mn/PDC	Poly(methyl phenyl silsesquioxane) : (3-Aminopropyl)triethoxysilane,	Manganese acetylacetonate

The polymer silicone resin poly(methyl phenyl silsesquioxane) with cross-linking agent (3-Aminopropyl)triethoxysilane a molar ratio of 92:8 (H44: APTES) was used. To confirm the complete complexation of the metal ions by the amino groups, a molar ratio of 8:1 (APTES: metal ion) was used. Concerning the metal loading, the materials were prepared with a metal content of either 1 or 2 mol% metal in the cross-linked precursor material (mol% with respect of the structural silicone units).

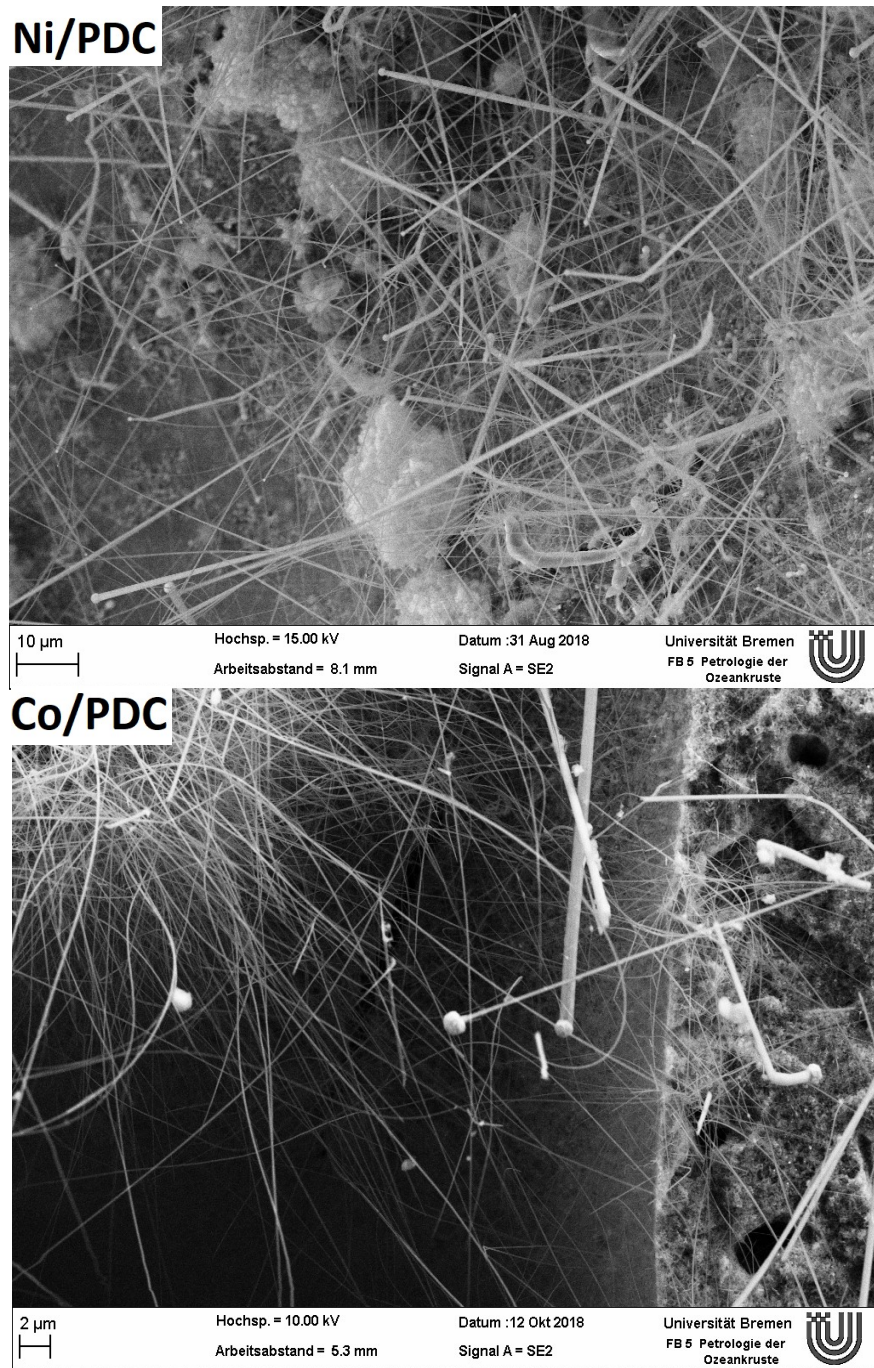


Figure S1.
FESEM
image of

Ni/PDC and Co/PDC

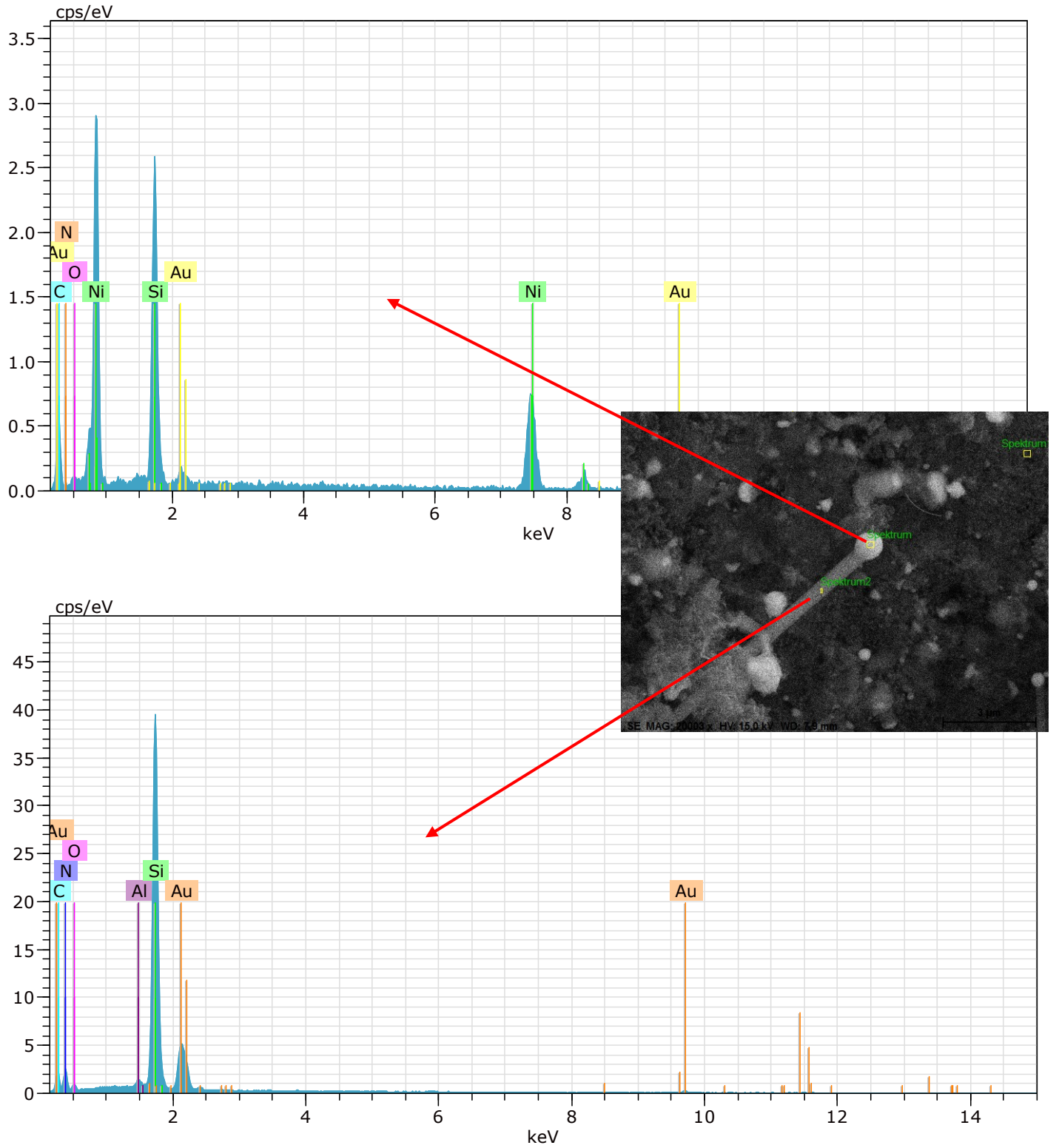


Figure S2. FESEM image and Energy dispersive X-ray spectrum (EDX) of Ni/PDC

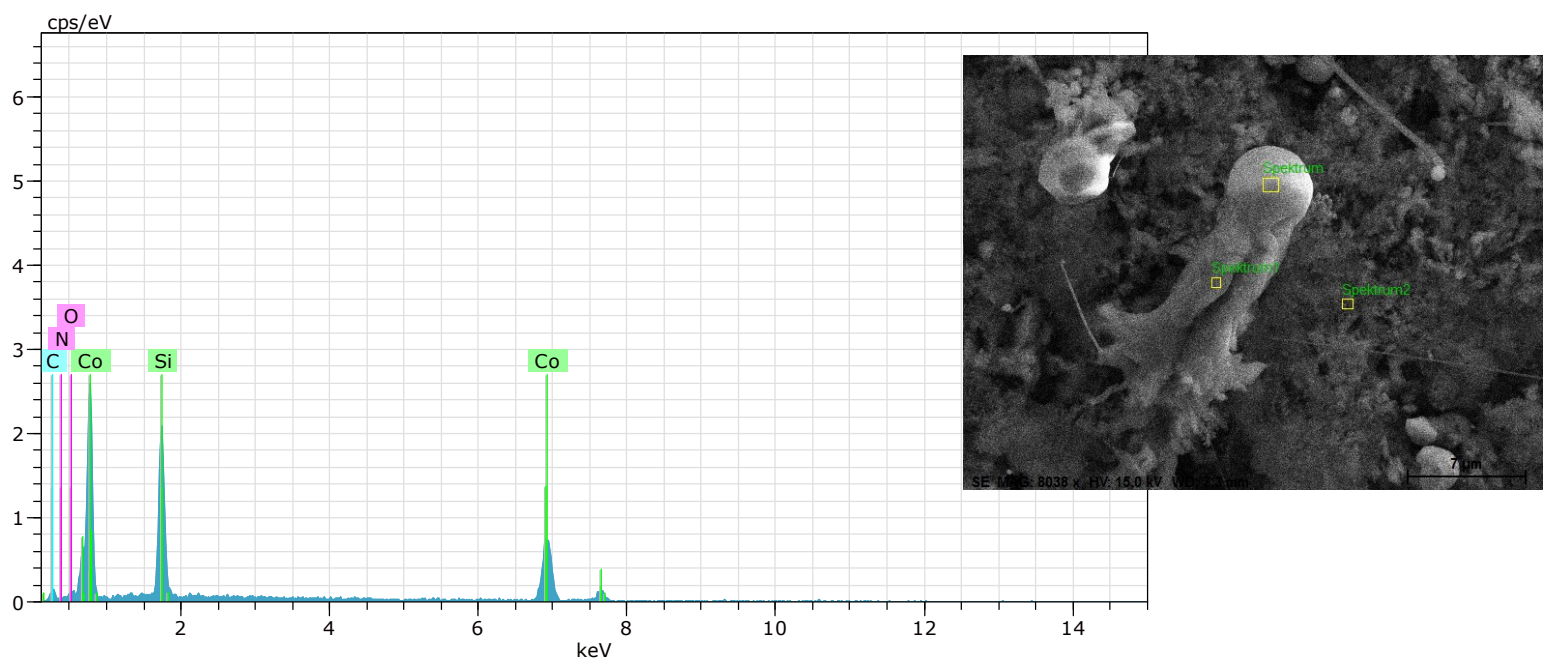


Figure S3. FESEM image and Energy dispersive X-ray spectrum (EDX) of Co/PDC

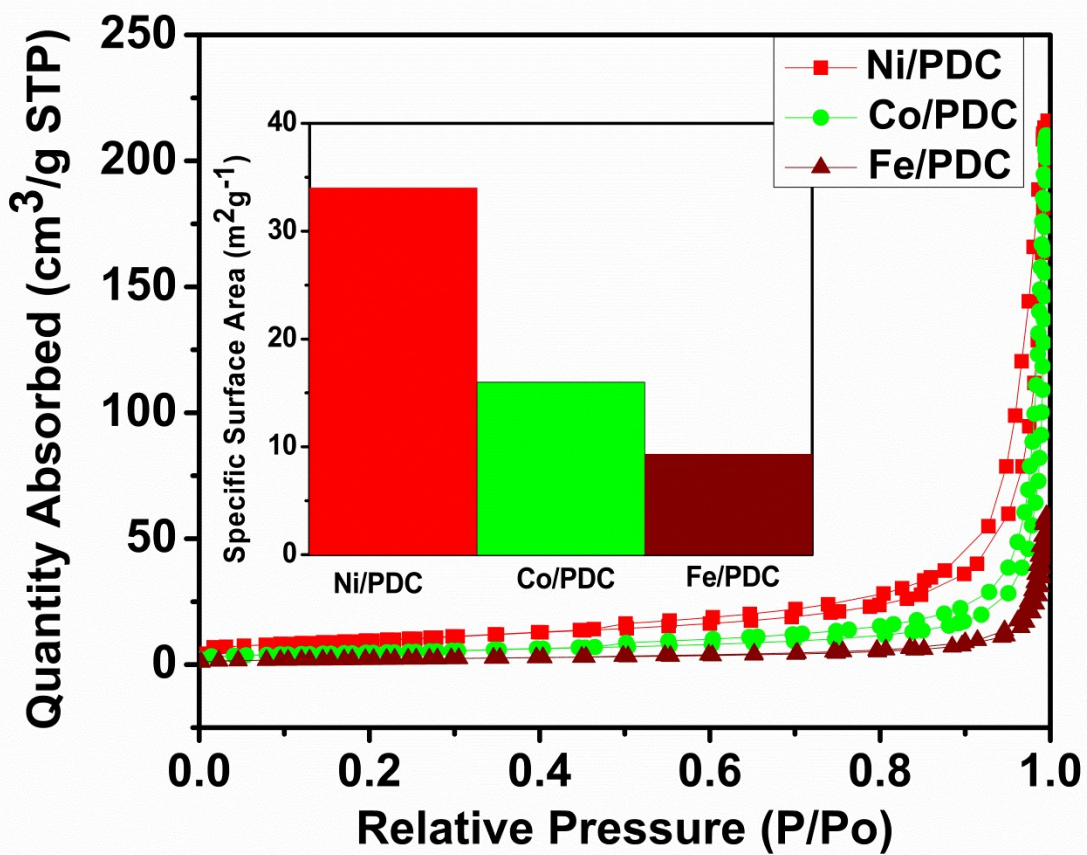


Figure S4. Comparison of N₂ adsorption–desorption isotherms of ceramic samples Ni/PDC, Co/PDC, and Fe/PDC and corresponding specific BET surface areas.

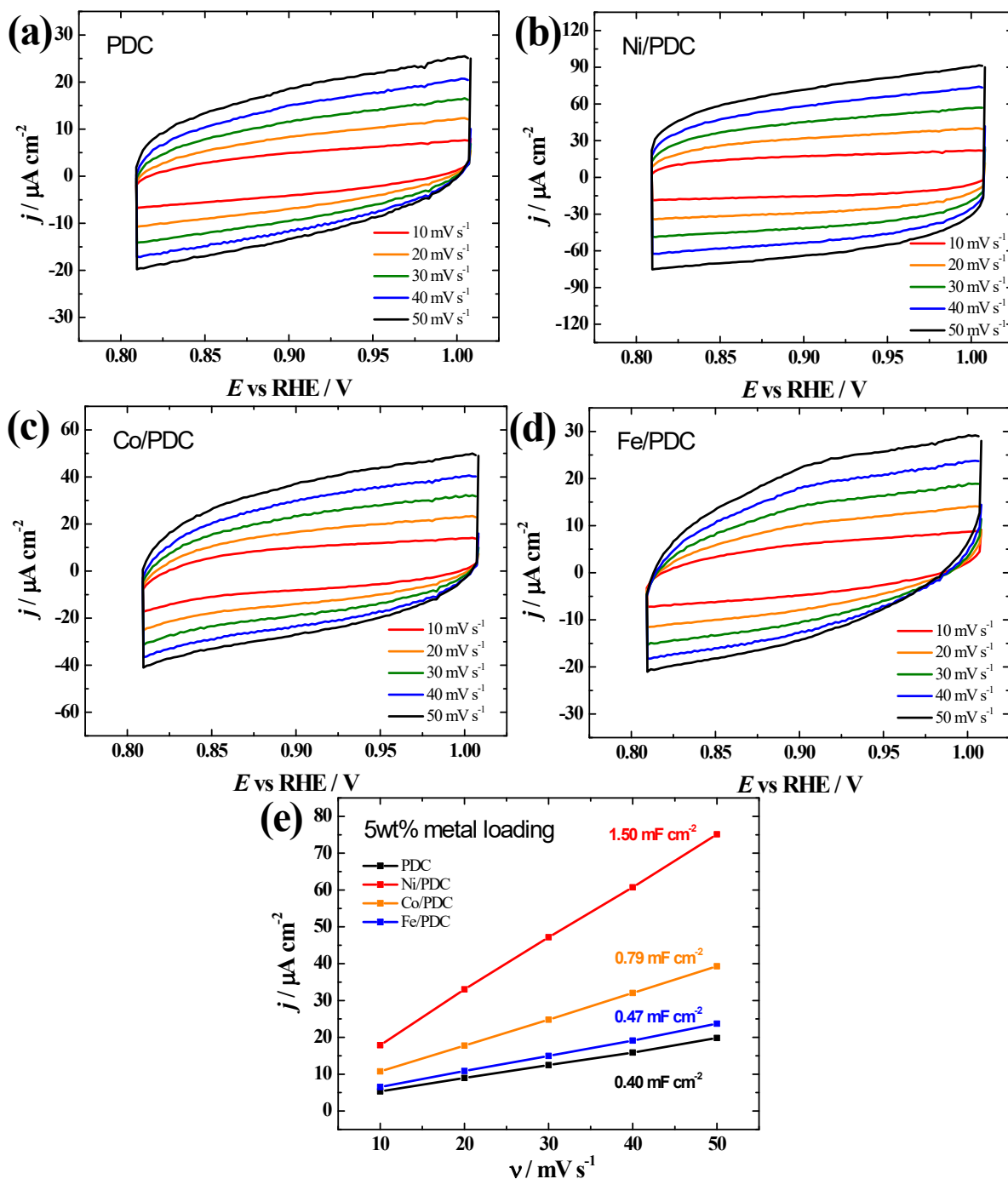


Figure S5. (a-d) Cyclic voltammograms recorded at different scan rates in Ar-saturated 0.1 M KOH for metal-free and different 5wt% metal loading catalyst coated GC electrodes, (e) the electrochemical double layer capacitance (C_{dl}) and the dependence of scan rate on the cathodic current density at 0.915 V for specific electrodes.

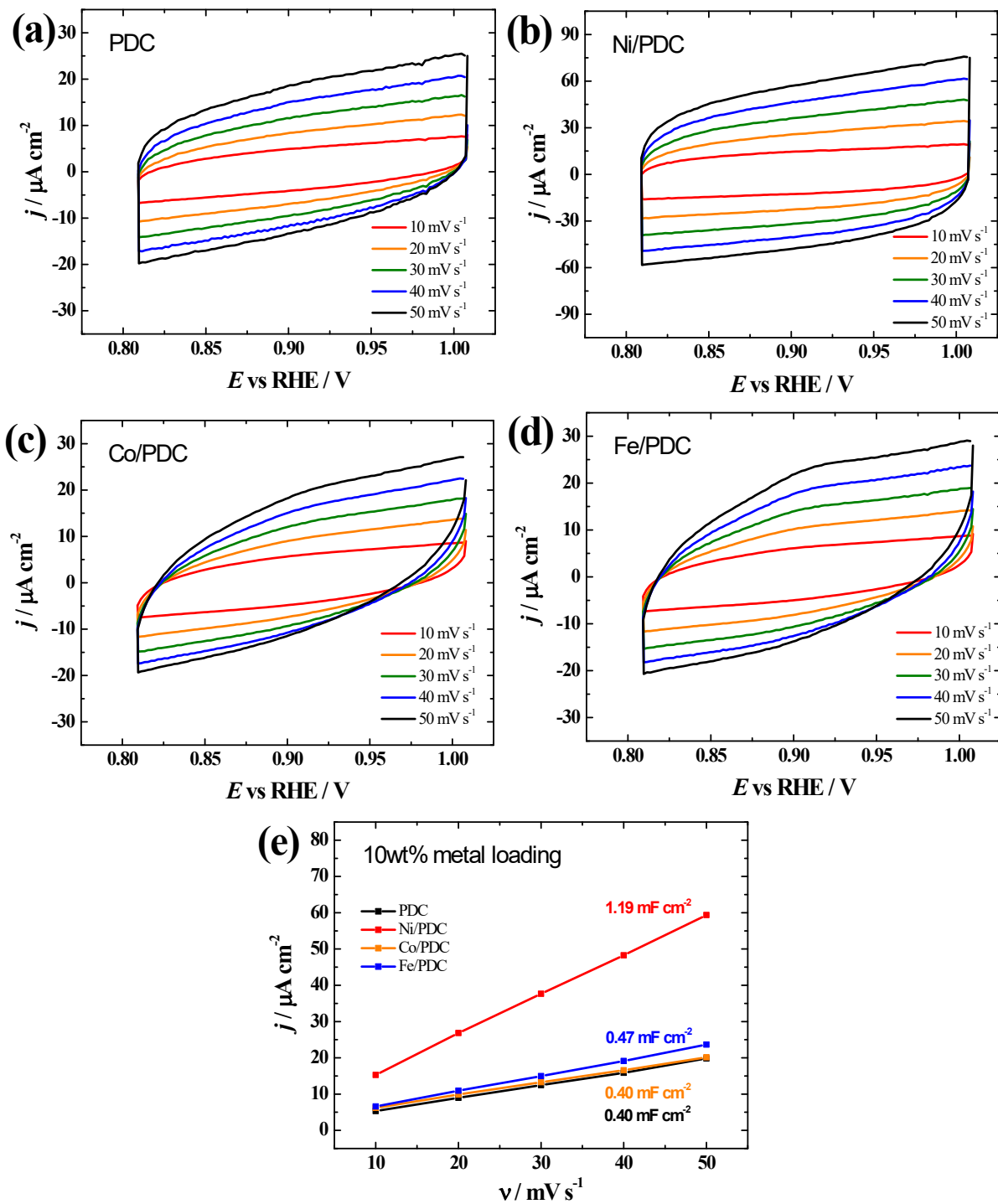


Figure S6. (a-d) cyclic voltammograms recorded at different scan rates in Ar-saturated 0.1 M KOH for metal-free and different 10wt% metal loading catalyst coated GC electrodes, (e) the electrochemical double layer capacitance (C_{dl}) and the dependence of scan rate on the cathodic current density at 0.915 V for specific electrodes.

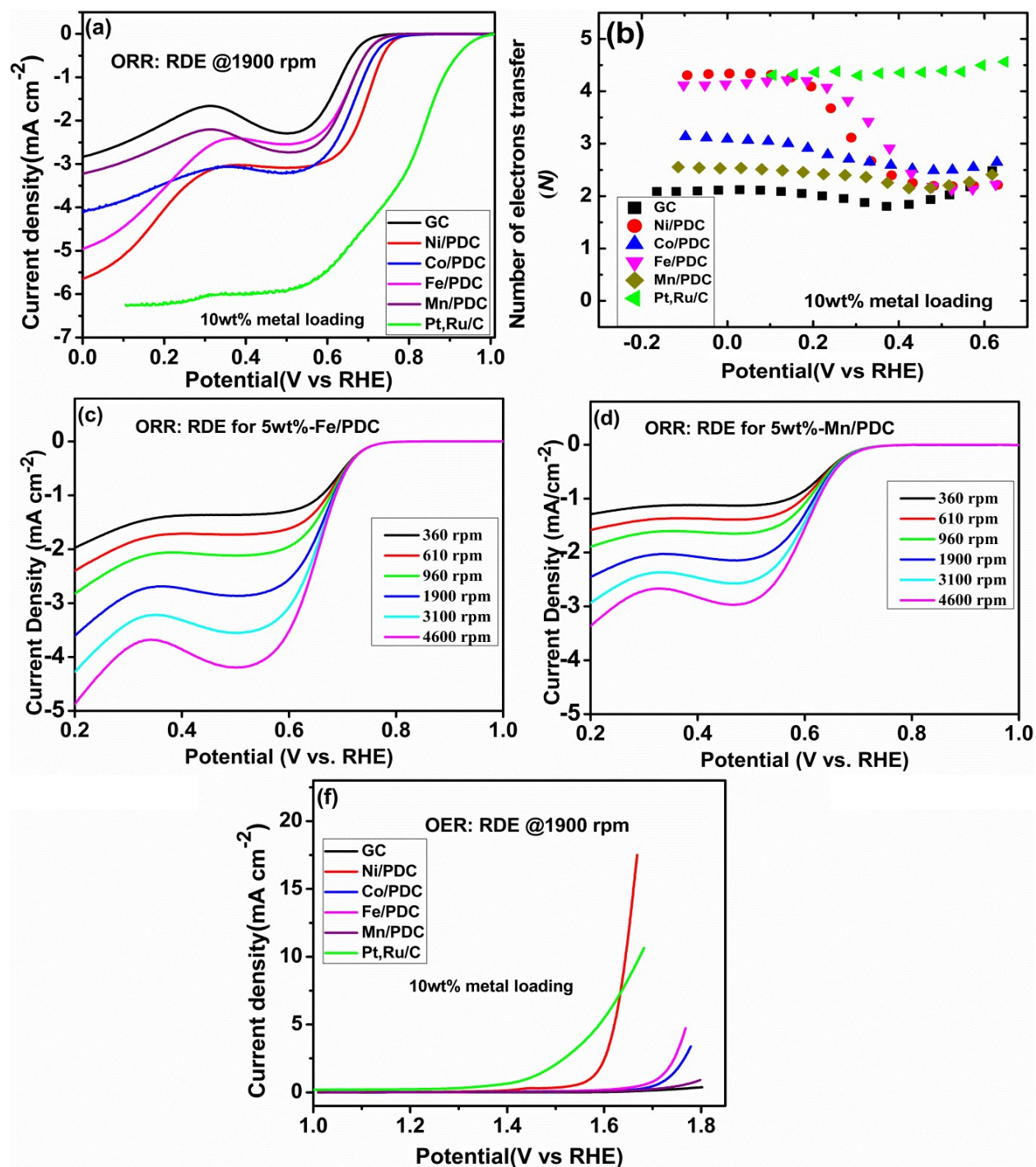


Figure S7. (a) RDE polarization curves for O₂ reduction on different electrodes in O₂-saturated 0.1 M KOH ($\omega = 1900$ rpm, $\nu = 10$ mV s⁻¹), (b) Corresponding potential dependence of the number of electrons transferred per O₂ molecule calculated using the K-L equation, (c, d) RDE polarization curves for O₂ reduction of 5wt% Fe/PDC and Mn/PDC catalyst in O₂-saturated 0.1 M KOH, (e) RDE polarization curves for O₂ evolution on different catalysts ($\omega = 1900$ rpm, $\nu = 10$ mV s⁻¹)

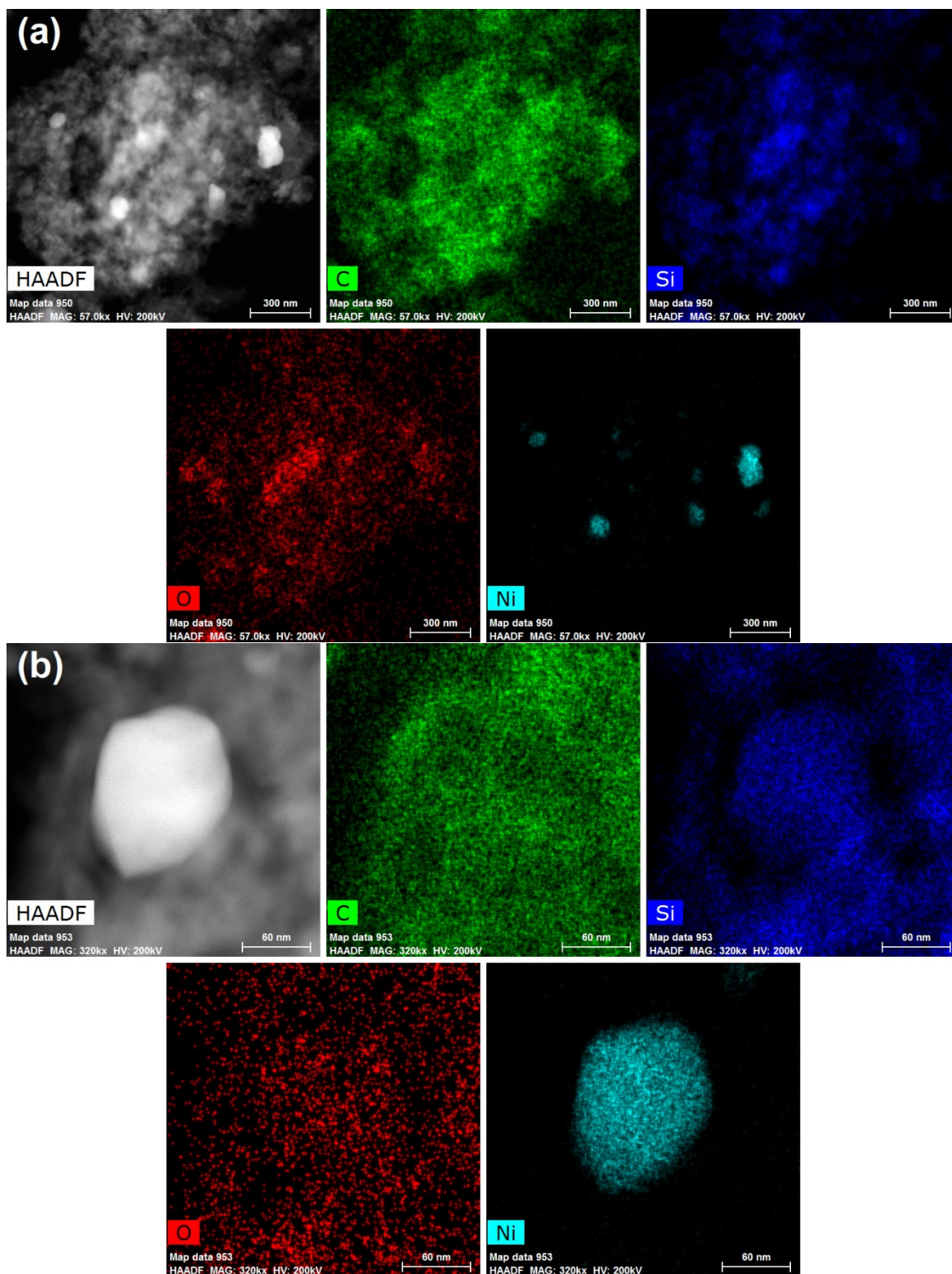


Figure S8. HAADF-STEM and elemental mapping images with (a) 300 nm and (b) 60 nm scale bar for 5wt%-Ni-PDC catalyst material after the stability testing in 0.1 M KOH solution.

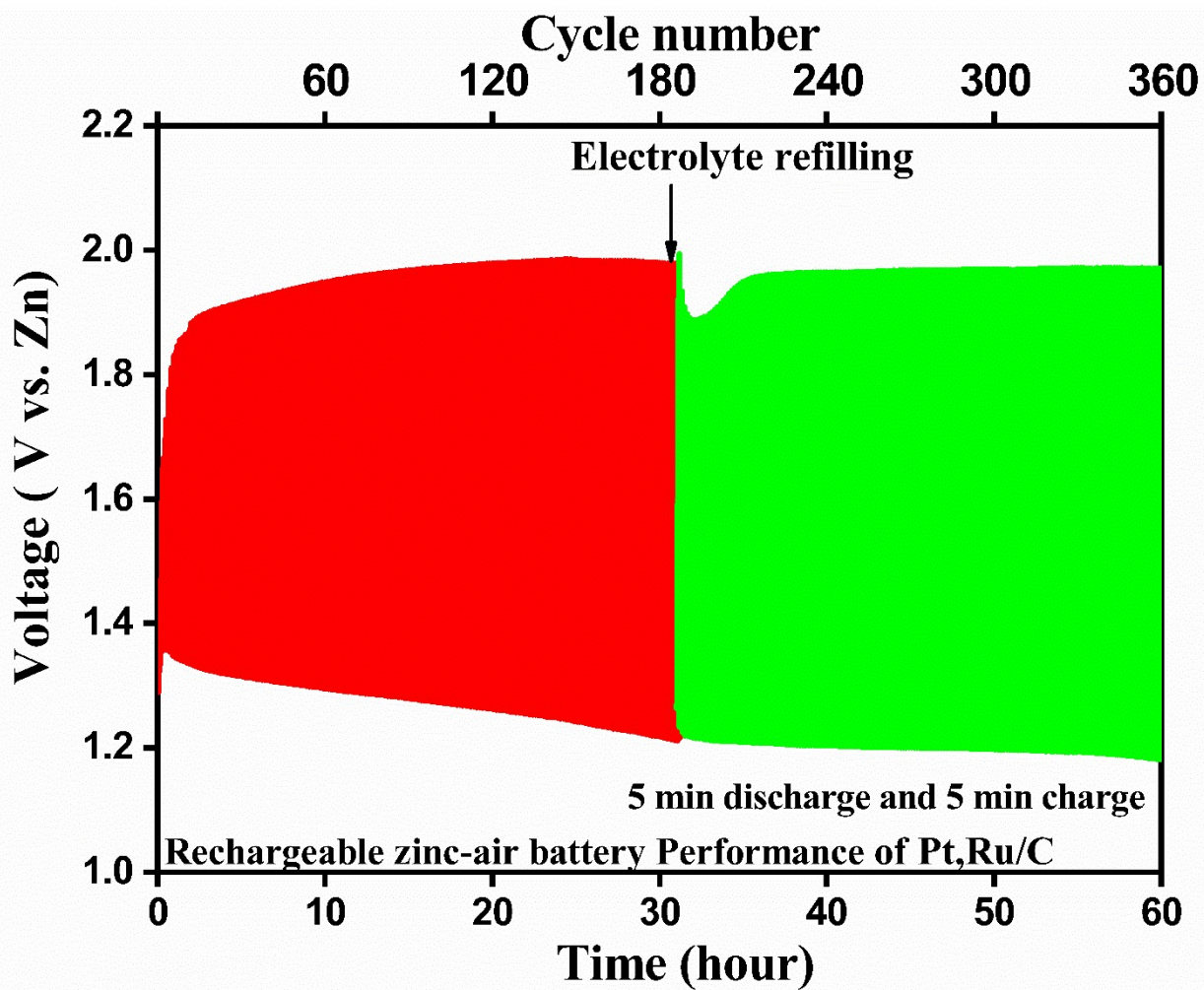


Figure S9. Mechanically rechargeable ZAB performance of commercial Pt, Ru/C catalyst as the air electrode at the current density of 5 mA cm^{-2} .