## Supporting Information for

## Core-Shell Catalysts Consisting of Nanoporous Cores for Oxygen Reduction Reaction

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Figure S1. HAADF-STEM images of dealloyed Pd-Ni/C (d-PdNi/C) with a nanoporous structure.



Figure S2. A single d-PdNi particle in sample 3: HAADF-STEM image (A) and 2D EELS mapping (B).



Figure S3. Fourier transformed EXAFS of the data (red) and first-shell fit (dotted blue) of (a) Ni K and (b) Pd K edges from the as-synthesized sample.



The number of repeated cell unites N > N'

Figure S4. Illustrations of porous and solid particles. The intensity (reverse of broadness) of a XRD peak depends on the number of repeated cell units (N) in one direction (a) in the crystal (p.6 in prism.mit.edu/xray/CrystalSizeAnalysis.ppt):

$$I = I_e F^2 \frac{\sin^2(\pi/\lambda)(s-s_o) \bullet N_1 a_1}{\sin^2(\pi/\lambda)(s-s_o) \bullet a_1} \frac{\sin^2(\pi/\lambda)(s-s_o) \bullet N_2 a_2}{\sin^2(\pi/\lambda)(s-s_o) \bullet a_2} \frac{\sin^2(\pi/\lambda)(s-s_o) \bullet N_3 a_3}{\sin^2(\pi/\lambda)(s-s_o) \bullet a_3}$$

The larger the number N, the higher the intensity (narrower of the XRD peak). For a porous particle, the number of the repeated unit cells (N') is smaller than that a solid particle with a similar overall particle size, resulting in a smaller crystallite size using Scherrer equation.



Figure S5. TEM image of sample 3 after first stage of dealloying in 1 M HNO<sub>3</sub> at 60°C for 1 hr. Porous structure was seen in some of the particles.