Supporting Information for:

Advanced electron microscopy characterization for pore structure of mesoporous materials; a study of FDU-16 and FDU-18

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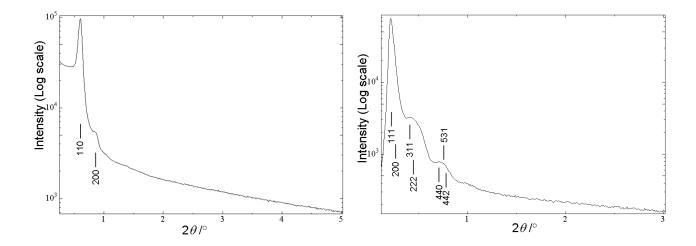


Figure S1. Powder X-ray diffraction (XRD) patterns of ordered mesoporous carbons FDU-16 (left) and FDU-18 (right). The observed reflections are indexed based on *Im-3m* for FDU-16 and *Fm-3m* for FDU-18, respectively. The cell parameters calculated from the highest peak positions are 13.5 nm for FDU-16 and 45.5 nm for FDU-18.

The XRD measurements were performed at BL02B2 in a synchrotron radiation facility SPring-8 (Japan) using a wavelength = 0.100 nm. The powder specimen was mounted in Lindemann glass capillary (diameter = 0.4 mm). The XRD profiles were recorded using an imaging plate set into a Debye-Scherrer type detector.

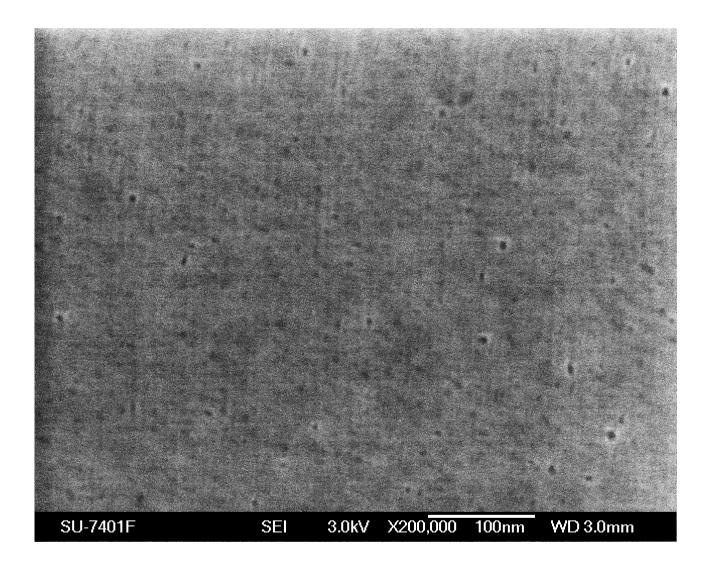


Figure S2. High resolution scanning electron microscopy (HRSEM) image of FDU-16 sample polished by a cross-section polisher (CP). This figure is an enlarged version of Fig. 5A.

A JEOL JSM-7401F was used for this HRSEM imaging. The sample powders were cross-sectioned by an Ar ion (JEOL SM-09010) with an accelerating voltage of 4 kV for 10 h and an emission of ~ 0.050 mA. Afterward, the powders were mounted on the bottom side of a silica wafer with the Ar-beam first irradiating the Si-wafer.