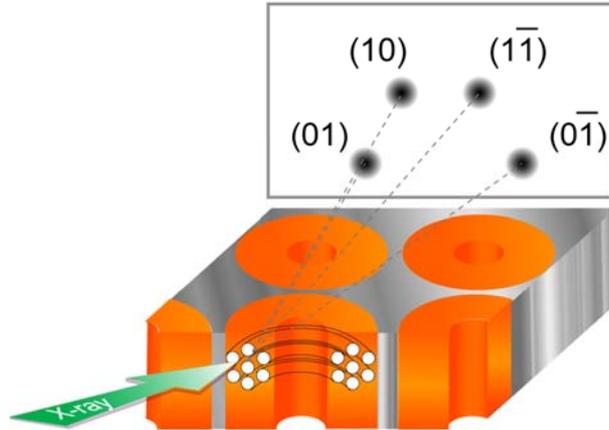


## Electronic Supplementary Information (Experimental section, Figure S1~S8)

### Experimental section

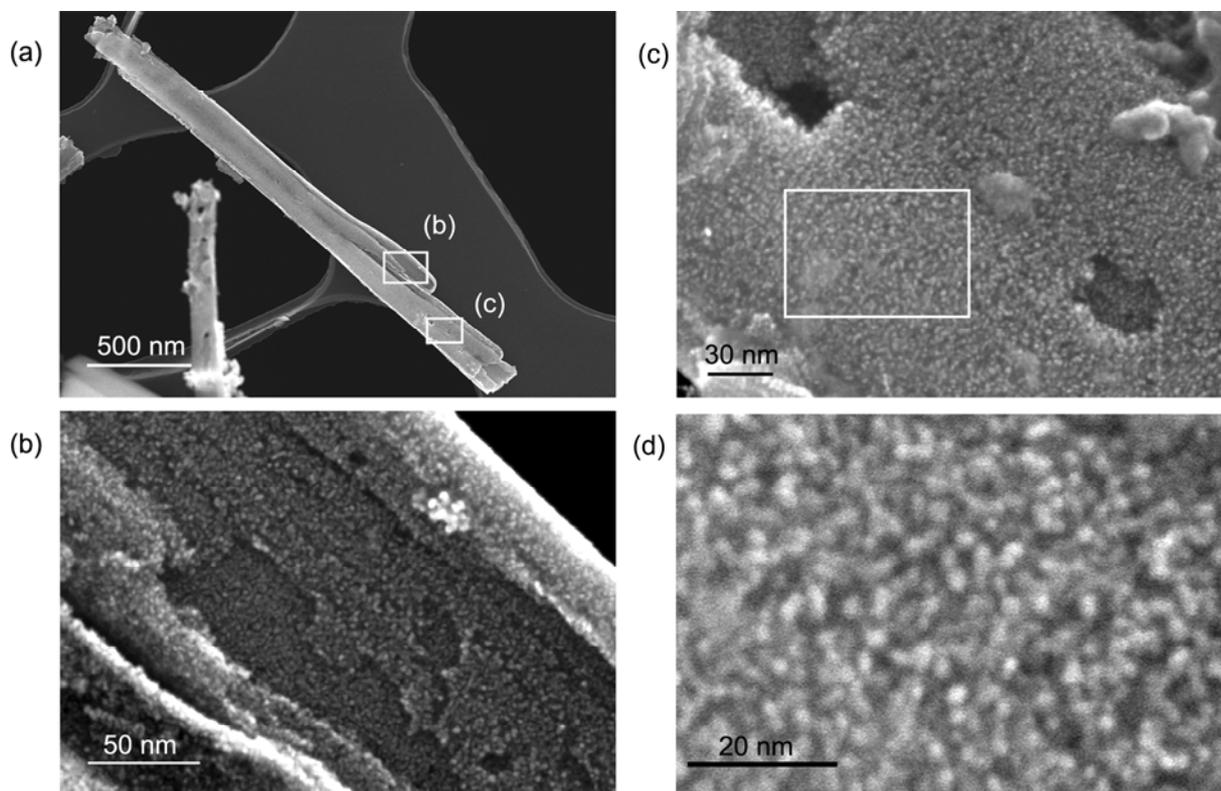
Mesoporous silica was synthesized using a precursor solution according to a previous  
5 report though C<sub>16</sub>EO<sub>8</sub> was used instead of Brij 56.<sup>20</sup> TEOS (10.4 g, Kishida Chemical Co.), ethanol  
(12 g), and dil. HCl (pH 2, 4.5 g, Kanto Chemical Co.) were mixed. After the mixture was stirred  
for 20 min, C<sub>16</sub>EO<sub>8</sub> (2.9 g) and ethanol (8.0 g) were added, and the mixture was stirred for 3 h. The  
total molar ratio was TEOS/C<sub>16</sub>EO<sub>8</sub>/HCl/ethanol/H<sub>2</sub>O=16.6:0.14:0.015:145:83.3. The precursor  
solution was dropped onto a PAAM and introduced into channels. The PAAM was dried under a  
10 reduced pressure condition. As-prepared PAAM was calcined at 400 °C for 15 h at a heating rate  
of 1 °C·min<sup>-1</sup>. The calcined PAAM was dissolved with an aqueous solution of about 10 wt%  
phosphoric acid. Then, silica nanorods were washed with water to remove phosphoric acid. When  
this precursor solution dropped onto a flat glass substrate, a 2D-hexagonally arranged mesoporous  
silica thin film was obtained.

**Figure S1**



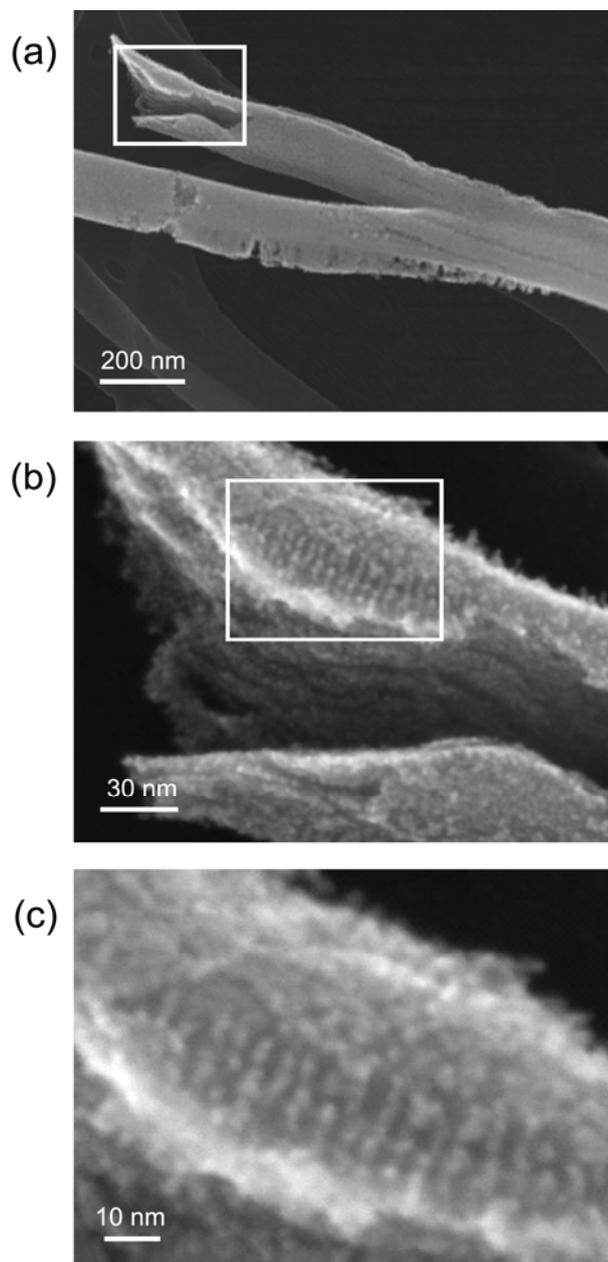
• **Figure S1** The 2D-XRD geometry and the proposed LLC mesophase structure within the PAAM after the reduced pressure process.

**Figure S2**



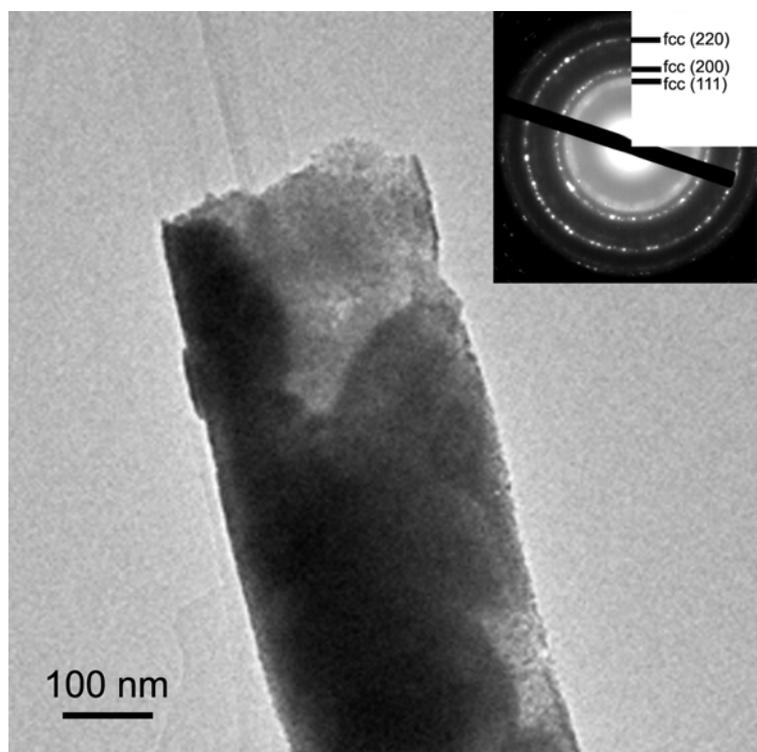
**Figure S2** SEM images of mesoporous Pt nanotubes. Figures (b) and (c) are internal and outer surfaces of the nanotubes. Figure (d) is a highly magnified image of Figure (c). Figures (b) and (c) are highly magnified images of the square areas in the Figure (a), and Figure (d) is a highly magnified image of the square area in Figure (c).

**Figure S3**



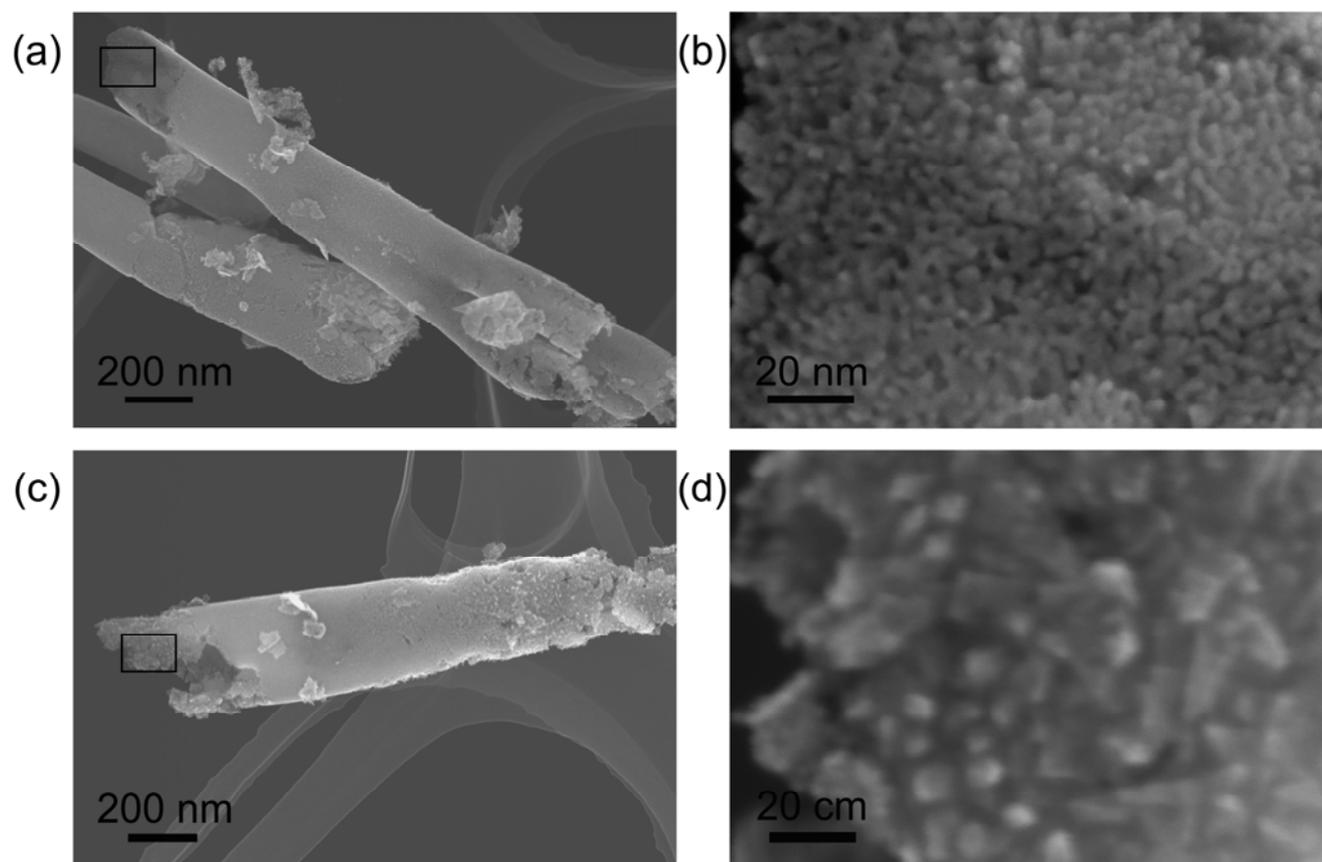
**Figure S3** HR-SEM images of mesoporous Pt nanotubes. Figures (b) and (c) are highly magnified images of the square areas in the Figures (a) and (b), respectively.

**Figure S4**



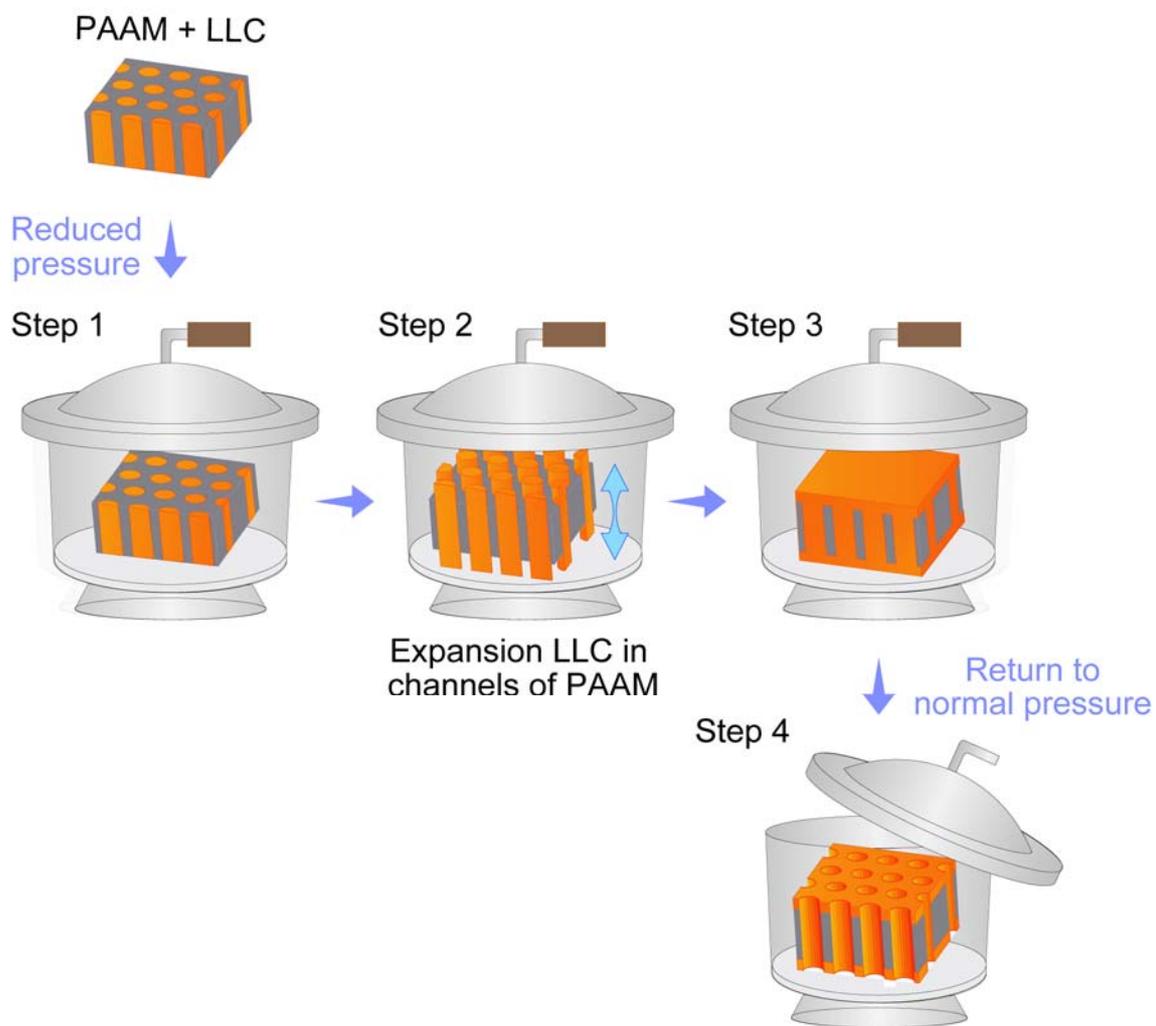
**Figure S4** TEM image of the mesoporous Pt nanotubes. Inset image is the selected ED patterns of a 100 nm region.

**Figure S5**



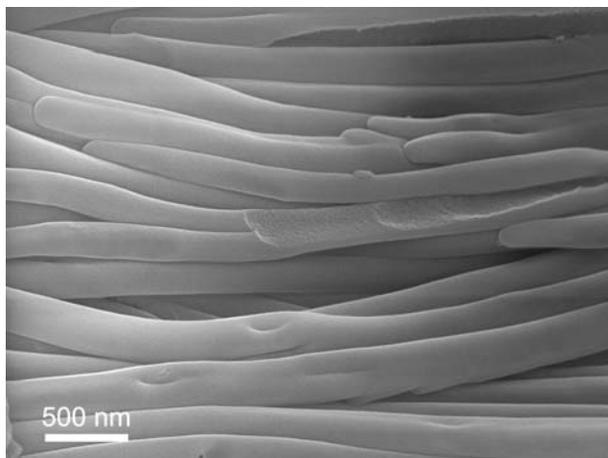
<sup>5</sup> **Figure S5** SEM images of the mesoporous Pt nanotubes treated at (a), (b) 100 °C and (c), (d) 150 °C, respectively. Figures (b) and (d) are highly magnified images of the square areas in the Figures (a) and (c), respectively.

**Figure S6**



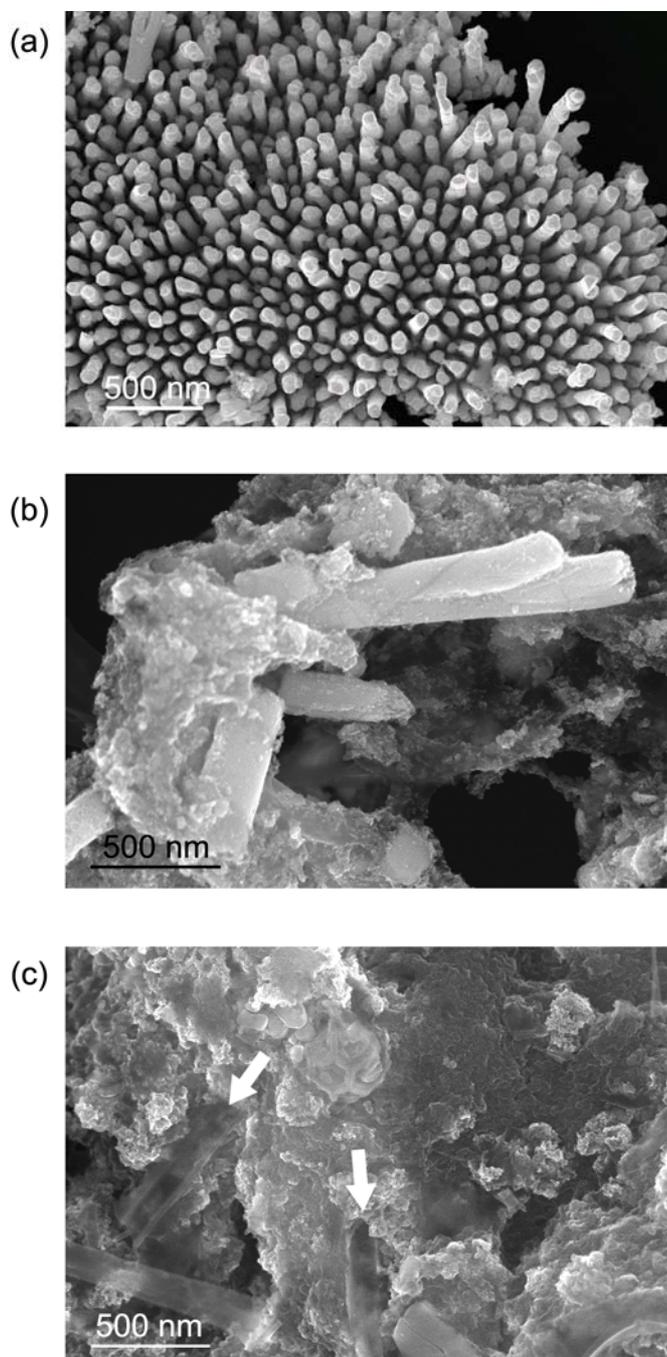
**Figure S6** Suggested formation mechanism for the mesoporous Pt nanotubes.

**Figure S7**



**Figure S7** SEM image of the mesoporous silica nanorods using PAAM.

**Figure S8**



**Figure S8** SEM images of the products prepared by changing the amount of ethanol/surfactant weight ratios: (a) 30, (b) 50, and (c) 100. In the Figure (c), mesoporous Pt nanotubes are indicated by arrows.