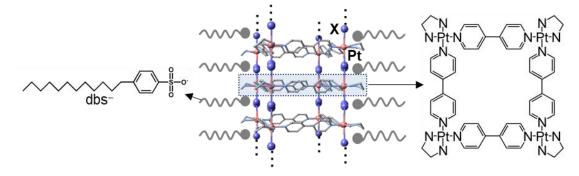
## Synthesis of MX-tube compounds soluble in organic media

(<sup>1</sup>Graduate School of Science, Kyoto University) ○Hao Liang,<sup>1</sup> Kazuya Otsubo,<sup>1</sup> Hiroshi Kitagawa<sup>1</sup>

**Keywords**: MX-tube compounds; Metal-organic frameworks; Mixed-valence compounds; Anionic surfactant

MX-tubes are an emerging kind of tubular compounds obtained from oxidative polymerization of macrocyclic complexes using halogen, which have the advantages of tunable electronic structure and channel structures.<sup>1,2</sup> Recently, a variety of MX-tube compounds were synthesized and their unique properties were also well demonstrated, such as selective sorption and high proton conduction.<sup>3,4</sup> However, these studies were limited to the aggregated solid state, leading to difficulties in direct observation of physical and chemical behavior in a single MX-tube. Besides, in light of application use of MX-tube such as thin film devices, to obtain the MX-tube compounds soluble in organic media is highly desired and challenging.

Herein, an anionic surfactant (dodecyl benzene sulfonate, dbs<sup>-</sup>) was used as the counter anion of MX-tube to improve the solubility in organic media. Two kinds of MX-tubes,  $[Pt(en)(bpy)X]_4(dbs)_8$  (en: ethylenediamine, bpy: 4,4'-bipyridine, X = I (1), Br (2)), were synthesized by mixing the precursor solution and aqueous solution containing dbs<sup>-</sup> at room temperature (**Figure 1**). The nanotubular structure of obtained compounds were confirmed by TEM observation, PXRD, Raman, and diffuse reflectance spectra. Due to the existence of anionic surfactants, 1 and 2 showed high solubility in chloroform solution, which is quite different from that of typical MX-tube compounds. Details are presented.



**Figure 1.** Schematic view of synthesized MX-tube,  $[Pt(en)(bpy)X]_4(dbs)_8(en:$  ethylenediamine, bpy: 4,4'-bipyridine, X = Br, I).

1) K. Otsubo, *CrystEngComm*, **2014**, *16*, 6277. 2) K. Otake, *Angew. Chem., Int. Ed.*, **2016**, *55*, 6448. 3) K. Otsubo, *Nature Materials*, **2011**, *10*, 291. 4) K. Otake, *Nat. Commun.*, **2020**, *11*, 843.