## **CHAPTER 4**

## CONCLUSIONS

## 4. 1 Synthesis of tungsten oxide by a hydrothermal method using ammonium metatungstate hydrate as a tungsten source

In this work, orthorhombic tungsten oxide (o-WO<sub>3</sub>) was synthesized by the 200 °C, 24 h hydrothermal reaction of ammonium metatungstate solutions containing different volumes of 1M HCl and hexadecyltrimethylammonium bromide (CTAB) cationic surfactant. In the HCl-free solution, the product was an amorphous phase. When 2.50 ml 1M HCl was added to the solution, both orthorhombic WO<sub>3</sub>·0.33H<sub>2</sub>O and WO<sub>3</sub> phases were detected. These products became pure orthorhombic WO<sub>3</sub> in the 5.00 ml and 7.50 ml 1M HCl-added precursor solutions. In 7.50 ml 1M HCl-added solution, the product was o-WO<sub>3</sub> microflowers, with microsquare layers growing out of their cores. These analyses showed that their phases and morphologies were controlled by the acidity of the solutions. FTIR and Raman vibrations of W=O, O-W-O, and W-O-W stretching modes were detected, and showed typical crystalline WO<sub>3</sub>. Their optical properties showed a maximum absorption at 275 nm in the UV region and a maximum emission peak at 375 nm.

## 4. 2 Synthesis of tungsten oxide by a hydrothermal method using sodium tungstate dihydrate as a tungsten source

Tungsten oxide (WO<sub>3</sub>) nanostructures with different morphologies were synthesized by hydrothermal reactions of sodium tungstate solutions containing different volumes of 3M HCl and ammonium sulphate as a capping reagent. A various factors influencing on the final morphologies of WO<sub>3</sub> such as acidity, reaction temperature and time were studied. The XRD patterns revealed pure phase hexagonal WO<sub>3</sub> (h-WO<sub>3</sub>) by comparing with its JSPDS database. Their morphologies were observed by SEM that shows nanoparticles, nanorods and nanowires, controlled by various factors. Hexagonal WO<sub>3</sub> nanowires with a diameter of 20-30 nm and lengths of up to several micrometers were successfully synthesized by a hydrothermal reaction at 200 °C for 48 h in a solution containing 5.00 ml 3M HCl-added precursor solution. FTIR and Raman vibrations of W=O, O-W-O, and W-O-W stretching modes were detected, and showed typical crystalline WO<sub>3</sub>. Their optical properties showed a maximum absorption at 275 nm in the UV region.

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