## CHAPTER V CONCLUSIONS AND RECOMMENDATIONS

In this work, TiO<sub>2</sub> particles prepared via sol-gel process were introduced into polypropylene by the master batch manufacturing process. The resulting composites were compared with the blends prepared with commercial nanoparticles TiO2. The effect of TiO2 contents and the properties of TiO2 from two sources of TiO<sub>2</sub> on the properties of resulting nanocomposites were studied. Results from DSC indicated that there was no change in the crystallization exotherms of TiO<sub>2</sub> from solgel process whereas composites with commercial nano-TiO2, the added TiO2 acted as a nucleating agent. At 5wt% and 10wt% of TiO2 in polypropylene were able to shift the crystallization exotherms by 6°C and 8°C, respectively. TiO<sub>2</sub> particles prepared from sol-gel process were still in micro size whereas the commercial TiO2 were much smaller, therefore it had more effect on the properties of polymer. The wide angle X-ray diffraction analysis showed that commercial TiO2 can increase the degree of crystallinity of nanocomposites by increasing weight contents of commercial TiO<sub>2</sub> which had an correspondence to the DSC results. Thermogravimetric analysis results showed that the degradation temperature of composites prepared from both sources of TiO<sub>2</sub> was increased by increasing contents of nano-TiO<sub>2</sub> from two sources, suggesting that the addition of nano-TiO<sub>2</sub> can improve the thermal stability of polymer. At 1-5wt% of TiO2 from two sources in PP, there were no adverse effects on mechanical properties of the composites, whereas at 10wt% commercial TiO<sub>2</sub> can increase much in Young's modulus due to higher in degree of crystallinity in this content.

The advantage of synthesis TiO<sub>2</sub> in sol-gel tetraol based method is the starting materials are stable in room temperature and the condition during the reaction is mild, but the obtained nanoparticles were agglomerated after calcination at high temperature. The grinding after calcination cannot control the size of particles as expected. The author would like to suggest the development of systhesis of TiO<sub>2</sub> to be of smaller particles in order to have larger surface area. The first is to try reducing the calcined temperature of TiO<sub>2</sub> to prevent the particles coming to agglomerate, but still need to make sure that it can also remove the organic substance

from the compound and after calcinations the particles should be sonicated in the solution instead of grinding. The other way is to find some coating compounds such as Hydroxypropyl cellulose and Carboxymethyl cellulose to prevent the agglomeration of TiO<sub>2</sub> particles.