Facile Synthesis of AgCl/BiYO$_3$ Composite for Efficient Photodegradation of RO16 under UV and Visible Light Irradiation
(Sintesis Mudah Komposit AgCl/BiYO$_3$ untuk Fotodegradasi Cekap RO16 di bawah Sinaran UV dan Cahaya Boleh Lihat)

URAIWAN SIRIMAHACHAI*, HUSNA HAROME & SUMPUN WONGNAWA

ABSTRACT
AgCl/BiYO$_3$ composite was successfully synthesized via the aqueous precipitation method followed by calcination. The varied amount of AgCl (10, 20 and 30%) was mixed into BiYO$_3$ via sonochemical-assisted method. The structures and morphologies of the as-prepared AgCl/BiYO$_3$ composite were characterized by x-ray diffraction (XRD), scanning electron microscopy (SEM) and UV-vis diffused reflectance spectroscopy (UV-vis DRS). The optical absorption spectrum of AgCl/BiYO$_3$ composite showed strong absorption in visible region. The photocatalytic activity of AgCl/BiYO$_3$ composite was evaluated by the photodegradation of reactive orange16 (RO16), which was selected to represent the dye pollutants, under UV and visible light irradiation. The results indicated that 20% AgCl/BiYO$_3$ photocatalyst was the most capable photocatalyst in this series in the degradation of RO16 under both UV and visible light illumination within 1 h. Moreover, the mechanism of photocatalytic degradation of AgCl/BiYO$_3$ was elucidated using three types of free radical scavengers. The significant enhancement was attributed to the formation of AgCl/BiYO$_3$ heterojunction resulting in the low electron-hole pair recombination rate.

Keywords: BiYO$_3$; photocatalyst; visible light

INTRODUCTION
Heterogeneous photocatalytic oxidation has received overwhelming interest as a potential method in the degradation of environmental pollutant. It is a process in which a photoexcitable solid catalyst is illuminated by light with the energy greater than the band gap of the semiconductor and electron-hole pairs are generated within the semiconductor. If electron and hole do not recombine, each can migrate to the surface of the solid catalyst and participate in the degradation (oxidation and reduction) of pollutants (Chong et al. 2010; Tariq et al. 2007). Semiconductor photocatalyst can be modified to expand their photoresponse to the visible region for pollutant decomposition such as by doping with metal/nonmetal (Chen et al. 2010; Hu et al. 2010; Zhou et al. 2011) or coupling with other small band gap semiconductor to form composite photocatalyst (Cao et al. 2011).

In order to develop novel photocatalysts for solar photocatalytic degradation of pollutants such as organic dyes, the Bi-based photocatalyst is one of the very interesting materials (Cai et al. 2015; Cheng et al. 2015; Xu et al. 2014; Xu et al. 2014). The efficient sunlight active photocatalyst, α-Bi$_2$O$_3$ nanorod, was synthesized and characterized which showed an excellent performance in the photocatalytic degradation of rhodamine B and 2,4,6-trichlorophenol (Kansal et al. 2015). However, another Bi-containing photocatalyst, BiYO$_3$, which having the perovskite structure has not much been studied in the photocatalytic process because of its poor photocatalytic performance (Qin et al. 2009). Wu et al.

Kata kunci: BiYO$_3$; cahaya boleh nampak; fotomangkin
(2015) synthesized a nanorod structure of BiYO₃ under hydrothermal condition which could photocatalytically degrade 68.5% of tetracycline within 3 h under visible light irradiation. In another work, the effect of templates on the formation of BiYO₃ nanostructure was studied and followed by their photocatalytic reduction of CO₂ under visible-light irradiation (Qin et al. 2016). Until recently, the enhancement of photocatalytic activity of semiconductor has become a key strategy in the design of composites, therefore, it is interesting to develop the photocatalytic activity of BiYO₃ by using the heterojunction system.

In this paper, AgCl/BiYO₃ composite, with different mole percents of AgCl fabricated onto BiYO₃, were prepared with the aim for low cost of production and ease of synthesis. The morphology and optical property were characterized. In addition, the photocatalytic activity and mechanism of AgCl/BiYO₃ composite toward RO16 dye under UV and visible light irradiation were also investigated.

MATERIALS AND METHODS
Bismuth (III) nitrate (Bi(NO₃)₃, Sigma-Aldrich), yttrium (III) nitrate (Y(NO₃)₃, Sigma-Aldrich), ammonia solution (NH₃, Labscan), silver (I) nitrate (AgNO₃, VWR, Belgium) and potassium chloride (KCl, Merck) were used with no further purification. First, the Bi(NO₃)₃ solution was added dropwise to Y(NO₃)₃ solution with stirring. The mixture was adjusted to pH = 8 by using 0.5 M NH₃ solution to yield a white precipitate. The precipitate was dried at 90°C. The obtained powder was ground and calcined at 750°C for 3 h under normal atmosphere (Qin et al. 2009). The final product was yellow powders.

The ultrasound-assisted preparation of AgCl/BiYO₃ composite with various amounts of AgCl loading was as follows: the desired amount of the as-prepared BiYO₃ and KCl were added to 50 mL of deionized water. The powders were mixed in a 100 mL glass beaker for 30 min and then sonicated for 30 min. Subsequently, the desired amount of AgNO₃ solution was rapidly injected to the mixture above. Finally, the mixture was further sonicated for 1 h. The product was collected by washing with deionized water and dried at 80°C for 12 h (Xue et al. 2015). The products were assigned as 10, 20 and 30% AgCl/BiYO₃.

The power x-ray diffraction (XRD) patterns of the samples were measured by X’Pert MPD diffractometer (Philips, Netherlands) using CuKα (λ = 0.154 nm) at a scanning rate (2θ) of 0.05° s⁻¹ and a 2θ range of 5-90° at 40 kV and 30 mA. The morphology of the prepared samples was analyzed using electron microscope (SEM, Quanta-400, FEI). UV-vis diffused reflectance spectra of the synthesized samples were determined by UV-2450 UV-Visible spectrophotometer.

The photocatalytic activity of AgCl/BiYO₃ was evaluated by the degradation of reactive orange16 (RO16), a cationic dye, under UV and visible light. The UV and visible light sources were 3 bulbs of 18 Watt black light lamp (Sylvania, USA) and 3 bulbs of 18 Watt fluorescent lamp (Sylvania, USA), respectively. In a typical experiment, 150 mL of the reaction suspension containing 150 mg catalyst and RO16 (1x10⁻⁵ M) was placed in a tightly closed reaction chamber. Prior to illumination, the suspension was magnetically stirred in the dark for 30 min to establish an adsorption-desorption equilibrium between the photocatalyst and dye. Then it was illuminated for a certain period of time with constant stirring. About 5 mL of the suspension was collected at each specific irradiation time interval, centrifuged and the supernatant was collected to analyze the absorption spectrum using UV-2600 spectrophotometer (Shimadzu, Japan). The dye concentration was determined from the wavelength at maximum absorption, 494 nm, with deionized water as a reference.

The setup of experiment with scavengers was similar to the photocatalytic degradation experiment above. A specified amount of scavengers was introduced into the RO16 solution prior to addition of the catalyst. The scavengers used in this experiment were: tert-butanol (TBA), ammonium oxalate (AO), and 1,4-benzoquinone (BQ).

RESULTS AND DISCUSSION
The XRD patterns of the as-synthesized AgCl/BiYO₃ with different AgCl content and BiYO₃ are shown in Figure 1. All of diffraction peaks can be indexed to the cubic phase of BiYO₃, which is in good agreement with the standard card (JCPDF No.00-055-0702) while AgCl (JCPDF No.01-071-5209) is cubic structure. A noticeable increase in intensity of AgCl diffraction peaks (2θ = 28.4) was observed with increasing AgCl content in AgCl/BiYO₃, whereas that of BiYO₃ (2θ = 27.8) was observed with decreasing AgCl content in AgCl/BiYO₃.

![Figure 1. XRD patterns of BiYO₃, 10%, 20%, and 30% AgCl/BiYO₃.](image-url)
The UV-Vis diffused reflectance spectra (DRS) of the photocatalysts are displayed in Figure 2(a). It can be seen that BiYO₃ and AgCl/BiYO₃ are responsive to visible light as both samples exhibit absorption in the visible light range (λ > 400 nm). The absorption edge of BiYO₃ appears at longer wavelength than that of AgCl/BiYO₃. This may come from the nature of AgCl which has its own absorption edge at 423 nm (Cao et al. 2011). Hence, the onset absorption of AgCl/BiYO₃ composite was blue shifted with the increasing of AgCl contents.

The optical bandgap energy ($E_g$) can be determined by using the Kubelka-Munk relation as shown in (1) (Pan et al. 2013; Yan et al. 2013);

$$\alpha(h\nu) = A(h\nu - E_g)^{1/2},$$

where $\alpha$ is the absorption coefficient; $\nu$ is the radiation frequency; $h$ is the Planck’s constant; $A$ is a constant; and $E_g$ is the band gap. The $E_g$ value can be obtained from Figure 2(b) by using the intersection point where the tangent of the curve intersects with the energy ($x$) axis. All the $E_g$ values obtained by this method are listed in Table 1.

The conduction band (CB) and valence band (VB) potentials can be calculated by the following empirical equations (Wang et al. 2008):

$$E_{VB} = X - E_e + 0.5E_g$$

(2)

$$E_{CB} = E_{VB} - E_g$$

(3)

where $E_{VB}$ is the valence band edge potential; $E_{CB}$ is the conduction band edge potential; $X$ is the electronegativity of the semiconductor (which is the geometric mean of the electronegativity of the constituent atoms); and $E_e$ is the energy of free electrons on the hydrogen scale (∼4.5 eV). By using (2) and (3), the $E_{CB}$ and $E_{VB}$ of BiYO₃ were calculated to be -0.04 and 2.58 eV, respectively. These values together with the $E_{CB}$ and $E_{VB}$ of AgCl (0.11 and 3.04 eV) (Cao et al. 2011) will be used to determine the photocatalytic mechanism diagram in this work.

| Compound          | $E_g$ (eV) |
|-------------------|-----------|
| BiYO₃             | 2.62      |
| 10%AgCl/BiYO₃     | 2.92      |
| 20%AgCl/BiYO₃     | 3.04      |
| 30%AgCl/BiYO₃     | 3.04      |
| AgCl              | 2.93      |

*Cao et al. 2011.

The morphology of the as-synthesized BiYO₃ and AgCl/BiYO₃ were characterized with a SEM at 50,000× magnification (Figure 3). BiYO₃ particles exhibited anomalous shape with diameters in the range of 50-60 nm and densely aggregated together (Figure 3(a)) while the size of AgCl/BiYO₃ was larger with the increasing amount of AgCl contents as illustrated in Figure 3(b)-3(e). Since AgCl could be decomposed by the high energy electron beam, the morphology at the higher resolution was not investigated. We also used a TEM to clarify the morphology of AgCl on BiYO₃ but failed to obtain any good results because the AgCl particles were not stable under electron beam. The rapid decomposition of AgCl under electron beam was observed as the AgCl particles shrunk and combined with BiYO₃ particles. The EDX spectrum of 20%AgCl/BiYO₃ composite with elemental mapping (Figure 3(e)) was acquired and used to confirm the existence of AgCl dispersed evenly on BiYO₃.

The photocatalytic decompositions of RO16 by BiYO₃, AgCl/BiYO₃, and P25 under UV and visible light illumination are shown in Figure 4. Under UV light, the photocatalytic activity of AgCl/BiYO₃ was almost equal to that in P25. In contrast, the photocatalytic performance of AgCl/BiYO₃ composite was greater than that of P25 when the catalyst was irradiated with visible light. This means that the AgCl/BiYO₃ is a better candidate visible light active material than P25. With this result, the mechanism of photocatalytic activity of AgCl/BiYO₃ under visible light irradiation was investigated and the data are displayed in Table 2. The rate constants of all samples in this work corresponded with the Langmuir-Hinshelwood (L-H)
kinetic model as given by the following equation (Dong et al. 2007; Gaya 2014; Wu et al. 2006).

$$k_{\text{app}}t$$

where $k_{\text{app}}$ is the apparent pseudo-first-order rate constant $(\text{min}^{-1})$; $C_0$ is initial concentration of RO16; and $C_t$ is RO16 concentration at time $t$. The results showed that only 10% and 7% degradations of RO16 over BiYO$_3$ were obtained under UV and visible light irradiation for 6 h, respectively. In contrast, the photocatalytic activities of AgCl/BiYO$_3$ with different AgCl contents were greatly enhanced under the same conditions. As the mole percentage of AgCl increased to 20%, the highest photocatalytic degradation was achieved at 100% photodecomposition within 2 h ($k_{\text{app}} = 0.0214 \text{ min}^{-1}$). Furthermore, the photocatalytic decomposition of RO16 was slightly decreased when the molar percentage of AgCl reached 30% ($k_{\text{app}} = 0.0211 \text{ min}^{-1}$) indicating the optimal AgCl content in AgCl/BiYO$_3$ composite was < 30%. Hence, 20% of AgCl fabricated on BiYO$_3$ was taken as the optimal value.

Moreover, it was clearly shown that the photocatalytic activity of AgCl/BiYO$_3$ photocatalyst was much higher than that of BiYO$_3$ itself. This photocatalytic enhancement of AgCl/BiYO$_3$ may result from the reduced recombination rate of the electron-hole pairs which, in turn, is attributable to the formation of AgCl and BiYO$_3$ heterojunction in AgCl/BiYO$_3$ composite.

In the photocatalytic oxidation process, electron-hole pairs were directly produced in the photocatalyst bulk after illumination. A series of photo-induced reactive species including hole ($h^+$), hydroxyl radical ($^\cdot$OH) and superoxide anion radical ($^\cdot$O$_2^-$) were suspected to be involved in the photocatalytic reaction. To examine the role of reactive species in photocatalytic degradation process, a series of scavengers were used to detect the relevant reactive species produced from photocatalytic process (Xu et al. 2013). Ammonium oxalate (AO), 1,4-benzoquinone (BQ) and tert-butanol (TBA) were quenchers to be used for hole ($h^+$), superoxide anion radical ($^\cdot$O$_2^-$), and hydroxyl radical ($^\cdot$OH), respectively.

Figure 5 shows the photocatalytic decompositions of RO16 with scavengers of which the experiment with low $k_{\text{app}}$ indicated the more important the role of the reactive species played in the reaction. For instance, the $k_{\text{app}}$ values for photodegradation of RO16 before and after adding TBA did not significantly change indicating the $^\cdot$OH radical was not the reactive species involved in this photocatalytic process. On the other hand, the addition of AO and BQ greatly reduced the $k_{\text{app}}$ indicating the hole...
and \(^{\bullet}O_2^-\) radicals played an important role to quench the photocatalytic decomposition of RO16 under UV and visible light irradiation. When AgCl and BiYO\(_3\) were irradiated under UV (or visible light), the electron was excited into the conduction band and the hole was left in the valence band, as in (5). The photogenerated electron in the valence band then reacted directly with adsorbed \(O_2\) on the surface producing the \(^{\bullet}O_2^-\) radical as in the following equation (6) (Gaya 2014; Lin et al. 2012):

\[
\text{hv} + \text{AgCl} \rightarrow h^+ + e^- \quad (5)
\]

\[
e^- + O_2 \rightarrow ^{\bullet}O_2^- \quad (6)
\]

Therefore, both hole and \(^{\bullet}O_2^-\) species were the dominating reactive species that finally degraded RO16 in this work. The possible mechanism for the RO16 decomposition over AgCl/BiYO\(_3\) composite could be proposed in Figure 6. The conduction band (0.11 eV) and valence band (3.04 eV) of AgCl lie below those of BiYO\(_3\) (\(E_{\text{CB}} = 2.58 \text{ eV}, E_{\text{VB}} = -0.04 \text{ eV}\), respectively. After irradiation, the photogenerated electrons migrated from the conduction band of BiYO\(_3\) to that of AgCl. Meanwhile, the photogenerated holes moved conversely from valence band of AgCl to the lower valence band of BiYO\(_3\). Through this process, it effectively reduced the recombination of the electron-hole pairs, which is believed to be the key factor to improve the photocatalytic performance of AgCl/BiYO\(_3\) (Lin et al. 2012). Furthermore, the accumulated electrons on the AgCl immediately reacted with the adsorbed \(O_2\) on the surface of composite to produce \(^{\bullet}O_2^-\). It was also noted that the holes, which accumulated on the BiYO\(_3\), could react directly with RO16. These two above mechanisms finally led to the degradation of RO16. However, in the case of BiYO\(_3\), it could be stimulated to produce electron-hole pairs by both UV and visible light and decomposed RO16 to ca. 20% within 6 h of irradiation, as shown in Figure 4, but the bandgap energy of BiYO\(_3\) (2.62 eV) was very narrow that the recombination of the electron-hole pairs became significant. Therefore, based on the above discussion, it can be proved that the decomposition of RO16 could be attributed to the reaction with hole and \(^{\bullet}O_2^-\) species during the photocatalytic reaction.

| Photocatalyst | % dye degradation | \(k_{\text{app}}\) (min\(^{-1}\)) | \(R^2\) |
|--------------|------------------|-------------------|--------|
| BiYO\(_3\)   | 19               | 0.0006            | 0.8247 |
| 10%AgCl/BiYO\(_3\) | 100            | 0.0190            | 0.9524 |
| 20%AgCl/BiYO\(_3\) | 100            | 0.0214            | 0.9410 |
| 30%AgCl/BiYO\(_3\) | 100            | 0.0211            | 0.9564 |
CONCLUSION

In this work, AgCl/BiYO\textsubscript{3} composite was successfully synthesized. Both BiYO\textsubscript{3} and AgCl show good visible-light absorption, but BiYO\textsubscript{3} itself displays low photocatalytic activity. The fabrication of AgCl onto BiYO\textsubscript{3} enhanced the photodegradation efficiency of R016 to 100 % within 1 and 2 h under UV and visible light, respectively, especially for 20%AgCl/BiYO\textsubscript{3} with $k_{\text{app}}$ value of 0.0214 mm\textsuperscript{-1} h\textsuperscript{-1}. The enhancement of photocatalytic activity of AgCl/BiYO\textsubscript{3} can be attributed to the effective electron-holepairs separation at AgCl/BiYO\textsubscript{3} heterojunction interfaces. The evaluation of photocatalytic mechanism illustrated that the hole and $\bullet O_2^-$ species were the dominant reactive species for photodecomposition of R016 under visible light irradiation. Results from this work show that the AgCl/BiYO\textsubscript{3} is a capable supported-AgCl composite in photocatalytic degradation of R016 under both UV and visible light illumination.

ACKNOWLEDGMENTS

This work was supported by the government budget of Prince of Songkla University (No. SCI570504S) and the Department of Chemistry, Faculty of Science, Prince of Songkla University.

REFERENCES

Cao, J., Xu, B., Luo, B., Lin, H. & Chen, S. 2011. Preparation, characterization and visible-light photocatalytic activity of Ag/AgCl/TiO\textsubscript{2}. Applied Surface Science 257: 7083-7089.

Cai, P., Zhou, S.M., Ma, D.K., Liu, S.N., Chen, W. & Huang, S.M. 2015. Fe\textsubscript{3}O\textsubscript{4}-modified porous BiVO\textsubscript{4}nanoplates with enhanced photocatalytic activity. Nano-Micro Letter 7: 183-193.

Chen, C., Ma, W. & Zhao, J. 2010. Semiconductor-mediated photodegradation of pollutants under visible-light irradiation. Chemical Society Reviews 39: 4206-4219.

Cheng, F.H., Huang, B.B., Dai, Y., Qi, Y.X. & Zhang, Y.X. 2010. One-step synthesis of the nanostructured AgCl/BiOI composites with highly enhanced visible-light photocatalytic performance. Langmuir 26: 6618-6624.

Cheng, Y., Wang, H., Zhu, Y., Liao, F., Li, Z. & Li, J 2015. One-step hydrothermal synthesis of BiVO\textsubscript{3}-Bi\textsubscript{2}O\textsubscript{3} p-n heterojunction composites and their enhanced photocatalysis properties. Journal of Material Science: Material Electron 26: 1268-1274.

Chong, M.N., Jin, B., Chow, C.W.K. & Saint, C. 2010. Recent developments in photocatalytic water treatment technology: A review. Water Research 44: 2997-3027.

Dong, X., Ding, W., Zhang, X. & Liang, X. 2007. Mechanism and kinetics model for degradation of synthetic dyes by UV-Vis/H\textsubscript{2}O\textsubscript{2}/ferrioxalate complexes. Dye and Pigments 74: 470-476.

Gaya, I.U. 2014. Heterogeneous Photocatalysis using Inorganic Semiconductor Solids. Dortrecht: Springer Netherlands. pp. 23-26 and 43-71.

Hu, X., Li, G. & Yu, J.C. 2010. Design, fabrication, and modification of nanostructured semiconductor materials for environmental and energy applications. Langmuir 26: 3031-3039.

Lin, H.L., Cao, J., Luo, B.D., Xu, B.Y. & Chen, S.F. 2012. Visible-light photocatalytic activity and mechanism of novel AgBr/ BiOBr prepared by deposition-prepiction. Chinese Science Bulletin 57: 2901-2907.

Pan, L., Hu, B., Zhu, X., Chen, X., Shang, J., Tan, H., Xue, W., Zhu, Y., Liu, G. & Li, W.R. 2013. Role of oxadiazole moiety in different D-A polyazoles and related resistive switching properties. Journal of Material Chemistry C 1: 4556-4564.

Qin, Z., Liu, Z., Liu, Y. & Yang, K. 2009. Synthesis of BiYO\textsubscript{3} for degradation of organic compounds under visible-light irradiation. Catalysis Communication 10: 1604-1608.

Qin, Z., Tian, H., Su, T., Ji, H. & Guo, Z. 2016. Soft template induced hydrothermal BiYO\textsubscript{3} catalysts for enhanced formic acid formation from the photocatalytic reduction of carbon dioxide. RSC Advances 6: 52665-62673.

Sood, S., Umar, A., Mehta, S.K. & Kansal, S.K. 2015. $\alpha$-Bi$_2$O$_3$nanorod: An efficient sunlight active photocatalyst for degradation of Rhodamine B and 2,4,6-trichlorophenol. Ceramics International 41: 3355-3364.

Tariq, M.A., Faisal, M., Muneer, M. & Bahnemann, D. 2007. Photochemical reactions of a few selected pesticide derivatives and other priority organic pollutants in aqueous suspensions of titanium dioxide. Journal of Molecular Catalysis A: Chemical 265: 231-236.

Wang, D.W., Huang, Q.F., Lin, P.X. & Yang, H.J. 2008. Visible-light-responsive photocatalyst: BiOBr-(1-x)BiOI. Catalysis Communication 9: 8-12.

Wu, C., Chang, H. & Chen, J. 2006. Basic dye decomposition kinetics in a photocatalytic slurry reactor. Journal of Hazardous Materials B 137: 336-343.

Wu, M., Xu, D., Lou, B., Shen, H., Wang, C. & Shi, W. 2015. Synthesis of BiYO\textsubscript{3} nanorods with visible-light photocatalytic activity for the degradation of tetracycline. Materials Letters 161: 45-48.

Xu, C.Y., Hu, S.P., Zhang, B.Y., Pei, Y. & Zhen, L. 2014. Solvothermal synthesis of Bi$_2$O$_3$CO$_3$nanoplates for efficient photodegradation of RhB and phenol under simulated solar light irradiation. Bulletin of Korean Chemical Society 35: 2935-2940.

Xu, H., Yan, J., Xu, Y., Song, Y., Li, H., Xia, J., Huang, C. & Wan, H. 2013. Novel visible-light-driven AgX/graphite-like C$_n$N$_x$ (X=Br, I) hybrid materials with synergistic photocatalytic activity. Applied Catalysis B: Environmental 129: 182-193.

Xu, L., Lu, H., Wei, B., Zhang, M., Gao, H. & Sun, W. 2014. Enhanced photosensitization process induced by the p-n junction of Bi$_2$O$_3$CO$_3$/BiOCl heterojunctions on the
degradation of rhodamine B. *Applied Surface Science* 303: 360-366.

Xue, B., Sun, T., Wu, J., Mao, F. & Yang, W. 2015. AgCl/TiO$_2$ nanocomposites: Ultrasound-assisted preparation, visible-light induced photocatalytic degradation of methyl orange and antibacterial activity. *Ultrasonics Sonochemistry* 22: 1-6.

Yan, J., Wang, C., Xu, H., Xu, Y., She, X., Chen, J. & Song, Y. 2013. AgI/Ag$_3$PO$_4$ heterojunction composites with enhanced photocatalytic activity under visible light irradiation. *Applied Surface Science* 287: 178-186.

Zhou, B., Zhao, X., Liu, H., Qu, J. & Huang, C.P. 2011. Synthesis of visible-light sensitive M-BiVO$_4$ (M = Ag, Co, and Ni) for the photocatalytic degradation of organic pollutants. *Separation and Purification Technology* 77: 275-282.

Department of Chemistry and Center of Excellence for Innovation in Chemistry
Faculty of Science
Prince of Songkla University
Kanjanawanich Rd., Kohong
Hatyai, Songkhla, 90110
Thailand

*Corresponding author; email: uraiwan.s@psu.ac.th

Received: 31 August 2016
Accepted: 17 January 2017