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Effect of Temperature and Particle Motion against the ability of ZnO Semiconductor Photocatalyst in Humic Acid

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ABSTRACT

It has been studied to determine the effect of temperature and particles motion of zinc oxide semiconductor from commercial ZnO powder on phototransformation of humic acid, using B & L 20 Spectronics. This study was done under some optimum conditions: lighting time of 2 hours, using Hg lamp of low pressure ($\lambda = 254$ nm), humic acid of 100 ppm and pH=9.2. it was disclosed that 97.51 % of humic acid transformed into minerals product. This research can contributed to solve the environmental problems in peat water.

Keywords: Photocatalyst, Zinc Oxide, Temperature, Particle Motion, Humic Acid

INTRODUCTION

Semiconductor applications widely known, both to obtain energy from indoor lights using oxide cuprum[1], photo-splitting water for hydrogen production[2] until the compound photo-transformation. All of these for overcoming the problem of the industry and the environment is a very interesting study. Various types of compounds [primarily organic compounds] contaminants that have been successfully transformed with these methods, among others, alkanes, aliphatic alcohols, aliphatic carboxylic acids, alkenes, phenolic compounds, aromatic carboxylic acids, dyes. Simple aromatic compounds, halogenated alkanes and alkenes, surfactants and pesticides[3].

Semiconductor zinc oxide [ZnO] is an alternative option after the semiconductor TiO_2 in its application as a photocatalyst. As a basic consideration is the semiconductor ZnO resistant to corrosion, has a relatively low light energy [$E_g = 3.40$ eV], and is much cheaper than TiO_2 [3]. Research photocatalyst ZnO has been made include studies in nano material in the form of suspension of nano-ZnO[4], nanowire ZnO[5], a thin film of ZnO[6], nanocrystalline ZnO[7], nano fiber ZnO [8], and several nano composite ZnO, such as Au-ZnO[9], FGS/ZnO[10], ZnO/CuO[11], ZnO- TiO_2 [12]. In this research, ZnO used is a commercial ZnO powder, because of economic considerations for direct application on water purification peat containing humic acid, fulvic acid and humin.

This study aimed to test the ability of the ZnO semiconductor photocatalysts for transforming humic acid. How the activity and efficiency of the photocatalyst ZnO for phototransforming humic acid in the solvent water under irradiation with a low-pressure mercury vapor lamp ($\lambda = 254$ nm), with the temperature parameters and particle dynamics ZnO semiconductor photocatalysts.

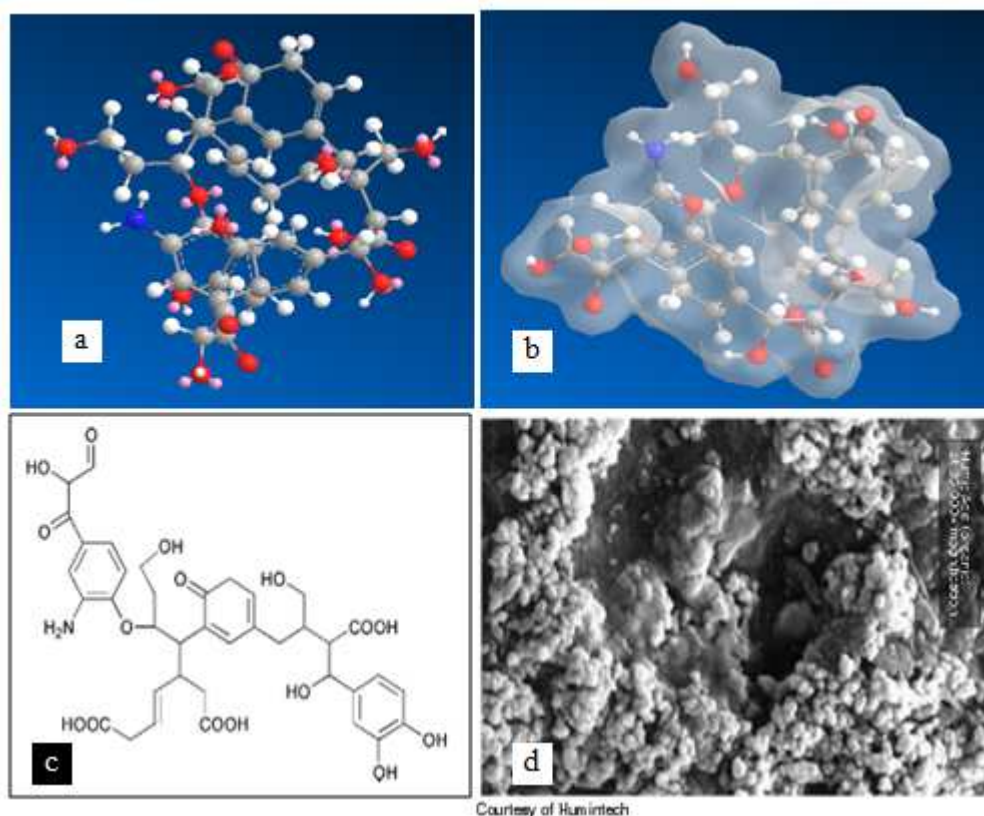


Figure 1. Humic Acid (HA) modeling (a) 3D Stealing model by ChemBio3D computerizing with software Cambridgesoft Ultra 11.0 (b) modeling surface HA of Connolly Molecular with Translucent display mode, (c) Humic Acid monomer structure according Steelink Model(13) and (d) SEM image of supramolecular humic acid with 35,000 x magnification (Humintech)

MATERIALS AND METHODS

Samples are peat water that was in the location of PT Mutiara Agam, Tiku, Pariaman. The tools used are Spectronic 20 B & L, balance of electric fan, mercury-vapor lamp low pressure ($\lambda = 254 \text{ nm}$), oven Gallenkamp, burette 25 mL, stopwatch, centrifuges, shakers, stirrers automatically, thermometer, desiccator, tube lighting, tweezers, glass plates, an electric cooker and tools commonly used glassware.

Materials used in the form of commercial ZnO powder, concentrated sulfuric acid, uranyl acetate 0:02 M, oxalic acid 12:02 M, potassium permanganate 0:02 M, HCL 0.1N NaOH 0.2 N, NaCl, fenolptalein, ethyl acetate, hexane, methanol, plate TLC and distilled water

Isolation Humic Acid

Humic Acid will be isolated by simple method. Into a glass beaker with a capacity of $30 \pm \text{L}$ filled 25 L of water peat 0.1 N HCl is added until a pH less than 2. After settling for 4 days, the precipitate separated by centrifuge. The precipitate was washed with 95% alcohol and then dried in the desiccator. Structure of Humic Acid can be modeled and showed in figure 1.

Identification and Purification Humic Acid

To test whether the isolated form brownish powder is humic acid or not done tests with a solution of NaCl. When the positive test solution precipitates a brown and grayish. Humic acid purification is done by washing the blackish brown powder with 95% alcohol many times until got 2 staining on Thin Layer Chromatography (TLC).

Determination of Effective Condition

Determination of the effective conditions include determining the distance of irradiation effectively done actinometry of uranyl oxalate, determining the effective concentration of ZnO, the determination of the effective exposure time and the determination of the effective pH.

Activity Test of ZnO Semiconductor Photocatalyst

Into the radiation tube already containing 20 mL of 100 ppm humic acid added ZnO x mg (corresponding to an effective concentration) and the pH was adjusted in accordance with the effective pH. The distance between the

tubes and mercury-vapor lamp is 5 cm. Stirrer is automatically turned on along with the mercury-vapor lamp ignites. Long exposures are defined in accordance with the effective exposure time. Temperatures are recorded each a half-hour irradiation.

RESULTS AND DISCUSSION

Activity Test of ZnO Photocatalyst

Activity test of photo-transformation will under due in reactor as showed in figure 2. By using all effective conditions (distance irradiation, ZnO concentration, exposure time and pH), then tested the activity of ZnO semiconductor photocatalysts that the data and results of calculations percent of its transformation can be seen in table 1 and figure 3. It can be observed that in 2 hours irradiation using light low-pressure, ($\lambda=254$ nm) to 20 mL of 100 ppm of humic acid at pH 9.2, was 97.51% humic acid can be transformed with 10 mg ZnO semiconductor photocatalysts. With the four treatment (treatment 2, 3, 4 and 5) the prediction of the effects of temperature, and the stirring of the photocatalyst photo-transformation process of humic acid.

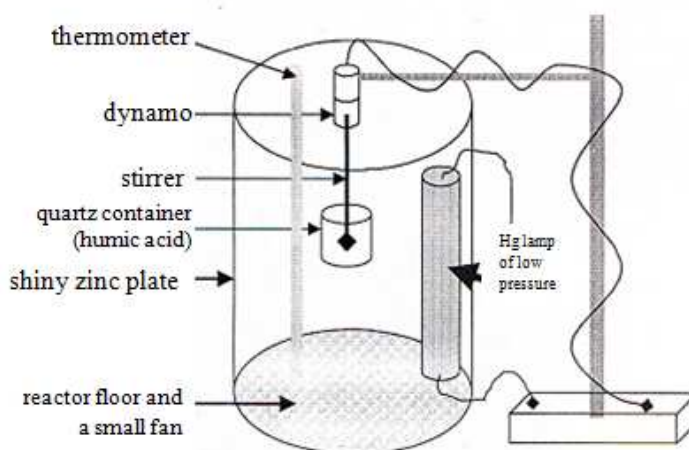


Figure 2. Schematic Photo-reactor for transformation humic acid

Modification of semiconductors by changing the environmental conditions found that at pH 12, E_g (gap energy) inexplicable existence. This is due to undergo dissolution to form ZnO, $Zn(OH)_2$ due to adsorb OH group (hydroxyl). This is due to an increase in catalytic activity due to increases in semiconductor ZnO valence band energy greater than the increase of energy in the conduction band. Such changes would reduce the energy gap (E_g) between PV and PK so that more electrons can be excited. At pH 3, two-thirds of PV energy rose while PK rose only one-third that E_g is reduced [14].

Oxygen plays an important role in the ZnO semiconductor photocatalysts (including TiO_2), then process involving contaminants photo-transformation semiconductor photocatalyst, light (can be used indoor lights) and oxygen can produce products that are environmentally friendly [15].

Table 1. Percent transformation upon irradiation of 100 ppm humic acid at pH 9.2

NO	TREATMENT	Percent Tranformation ($\lambda=730$ nm)	Percent Transformation ($\lambda=340$ nm, control)
1	ZnO + AH + hv (Stirred)	97,51	98,67
2	ZnO + AH (Dark) (Stirred)	1,33	2,04
3	ZnO + AH (54°C) (stirred)	5,38	5,43
4	ZnO + AH (indoor lights) (Stirred)	2,67	3,21
5	ZnO + AH + hv (No Stirring)	42,31	43,42
6	AH + hv (Stirred)	26,00	27,62

HA : Humic Acid 100 ppm pH 9,2

hv : photon from of low-pressure mercury vapor lamp ($\lambda = 254$ nm)

2-5 : with treatment

6 : as control

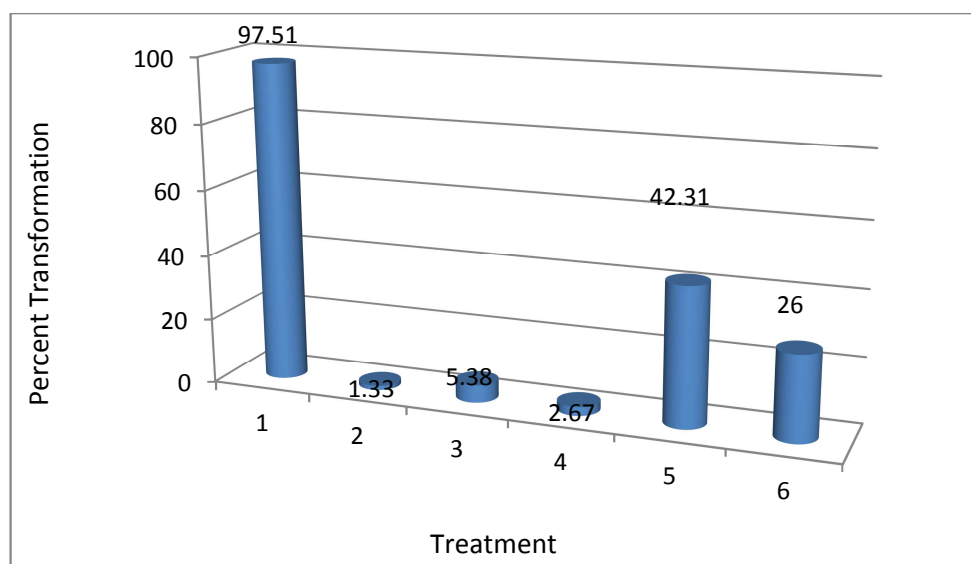


Figure 3. : Activity Test of Photocatalyst Semiconductor ZnO in various treatment

Effect of Temperature

As revealed by Cotton *et al* (1995) and Sharpe (1992) that the temperature is one of the drivers for the formation of the couple charged (h^+_{pv}/e^-_{pk}) which are vital to the working of semiconductor photocatalysts [16; 17]. The theory could explain why the treatment 2 (average temperature) 29°C there is still showing a ZnO semiconductor photocatalyst performance of 1.33% and increased in 3 treatments (average temperature of 34°C = average temperature of treatment 1) to 5.38%.

It can be observed that the result of joint work between ZnO semiconductor photocatalyst, irradiation and temperature above the Fermi level (0 K) gives the photocatalyst performance 97.51% humic acid within 2 hours. However, when compared with the treatment of temperature with radiation treatment plus a photocatalyst, the effect of irradiation plus photocatalyst (97.51% - 5.38% = 92.13%) is still much greater than the effect of temperature (34°C) plus a photocatalyst (5.38 %).

Effect of Particles Motion

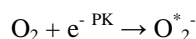
Actually, the process of transformation photo humic acid is a joint work between the photocatalyst, temperature, radiation, and stirring. Although, only a photocatalyst working alone or working radiation alone. There is interplay between the events of the photocatalyst, temperature, and radiation are mutually continuous in catalyzing the photo-transformation humic acid. However, the greatest influence is still given by the cooperation between the photocatalyst by irradiation.

When compared with the control treatment 1 and 6, it is evident that the presence of ZnO semiconductor photocatalysts, the process of photo-transformation lasts more powerful (97.51%). While the absence of ZnO process only efficiency photo-transformation of 26.00%. So the ZnO semiconductor photocatalyst help the photo-transformation process of humic acid by 71.51% (derived from the difference between 97.51% - 26.00%). The results of this study further support for what has been done Requel and Nogueira (1993), which compares labor semiconductor photocatalyst with two controls [18]. They obtained the fact that with the cooperation between the semiconductor and the radiation is able to transform 99% of methylene blue compared to 15% who did not wear a semiconductor.

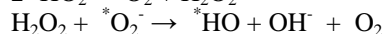
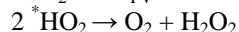
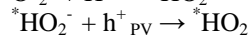
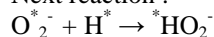
Effect of Stirring

With the stirring, the process of transformation lasts photos faster. The results of this study prove that treatment 1 (no stirring) gives percent larger transformation that is 97.51% compared 5 treatment (without stirring) is only 41.31%

The most important thing with the implementation of stirring is helping the process of dissolution of oxygen into the suspension. In accordance with Mao and Bakac report (1996) and Adamson (1990) that oxygen plays an important role in the process of photocatalysts [15; 19]. The oxygen function here is as an electron acceptor conduction band (e^-_{pk}) on the reaction:



Next reaction :



Highly reactive hydroxyl radicals will attack the substrate (in this case is the humic acid molecules). Furthermore, there will be a chain reaction that resulted transformation of humic acid.

The hydroxyl radical as the most important species to trigger the process is highly dependent photo-transformation amount of oxygen involved in the reaction. When oxygen levels are high then the more hydroxyl radicals are formed. Unsurprisingly, a process that uses oxygen dissolution stirring helps to have a higher performance than the process without stirring in a reaction semiconductor photocatalyst.

Another function is to flatten stirring so that the entire suspension portion exposed beams and enhance interaction between ZnO semiconductor photocatalyst molecules of humic acid. Further stirring can balance the adsorption-desorption processes on the surface of a semiconductor, because if the process is not balanced (higher than the adsorption desorption) then by Hoffman et al. (1995) substrate entry process will then be blocked. As a result, the efficiency will be lower semiconductor photocatalysts [3].

CONCLUSION

Based on the research process of photo-transformation 20 mL of 100 ppm of humic acid at pH 9.2 under irradiation mercury low pressure lamps ($\lambda = 254 \text{ nm}$) and with the help of 10 mg ZnO capable of transforming a 97.51% humic acid within 2 hours. Stirring helps the process of transformation of humic acid catalyzed by ZnO semiconductors amounted to 55.2%

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