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# Evaluation of new couple Nb<sub>2</sub>O<sub>5</sub>/Sb<sub>2</sub>O<sub>3</sub> oxide for photocatalytic degradation of Orange G dye

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**Abstract:** This paper involves the study the effect of differences in calcination temperatures on the prepared couple  $Nb_2O_5/Sb_2O_3$  oxide at percentage 1:1 with 400, 500 and 600 °C of calcination temperature and it was characterized by x-ray diffraction. Degradation test of orange G dye was carried out to determine photocatalyst activity for new couple  $Nb_2O_5/Sb_2O_3$  oxide. The photocatalyst experiments were performed at mass of the catalyst (0.05-0.3) g, pH solution in the range (2 -9) and dye concentration 10 ppm were monitored as a function of time at wavelength of 475 nm with recording optical absorbance. The photocatalyst destruction of Orange G dye was found to follow first order kinetics. The results indicate that 0.15 g was the best weight of the catalyst and the best pH for degradation orange G dye was at pH of solution equal to 6.8.

Keywords : Nb<sub>2</sub>O<sub>5</sub>/Sb<sub>2</sub>O<sub>3</sub> oxide, Orange G dye, Degradation, photolysis.

#### Introduction

Niobium oxide catalyst has received attention in the recent years due to broad industrial application such as solar cell ,optoelectronic technology and catalytic activity<sup>1</sup>. It is used in photo reaction as pure powder or couple with other semiconductor for degradation of organic compounds such as  $dyes^{2-12}$ . This paper used new couple of Nb<sub>2</sub>O<sub>5</sub>/Sb<sub>2</sub>O<sub>3</sub> that detectors the photo activate by using it for degradation of orange G dye.

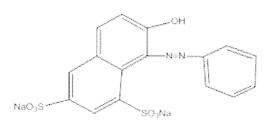
Orange G dye is a type of azo dyes and there are commonly used in many filed of industries such as textile, leather, paint, food, cosmetics and pharmaceuticals<sup>13-15</sup>. The are also the are used in biological field by giving various colors between different of the tissues to allow examination under light of microscopy. In recent years have been focused researchers attention heavily on the treatment of dye pollutants because the dye pollutants from industry are main sources for environmental contamination. It is estimated that 1-15% of the dye is lost during dyeing processes found many of new chemical treatment methods called advanced oxidation processes(AOP)<sup>16-18</sup>, one of the method is using couple semiconductors as heterogeneous photocatalyst the efficient of chemical oxidation processes for decontamination of drinking water by generation of very reactive free radical such as hydroxyl radical (OH<sup>-</sup>) which can oxidize organic compounds in to solution.

# Experimental

#### Materials

Chemical and materials were used in this work are antimony trioxide  $Sb_2O_3$  purity 98%, supplied by Fluka AG.3-Normal ,Niobium pent oxide  $Nb_2O_5$  has pure 99% supplied by Fluka AG.3-Normal ,and orange G

dye was supplied by sigma-Aldrich has purity more than 90%. NaOH and HCl were purchased from Merck (Germany), structure of orange G dye is given in Figure 1.



### Figure 1: Structure of Orange G dye molecule.

#### **Result and Discussion**

The effect of calcination temperatures on the formation of couple  $Sb_2O_3/Nb_2O_5$  at percentage (1:1) with different calcination temperatures (400,500 and 600) °C were investigated by X-ray diffraction as shown in Figure 2. These results are listed in Table 1. Shows  $2\theta$ , (d) and average partial size at different calcination temperatures.

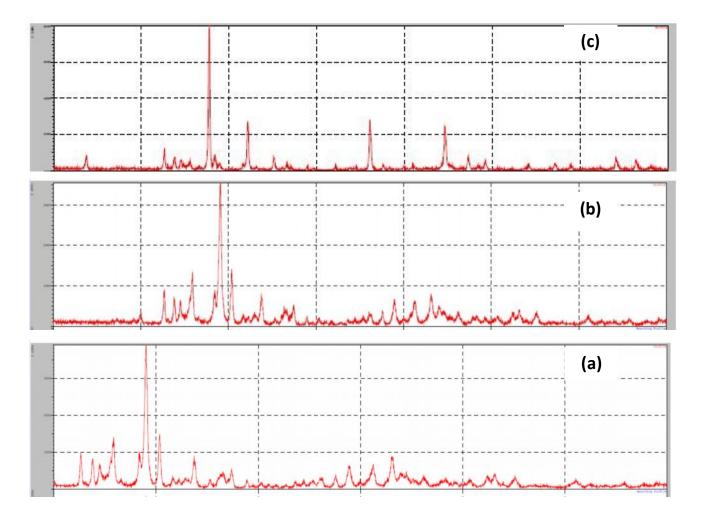


Figure 2: x-ray diffraction for couple of Sb<sub>2</sub>O<sub>3</sub>/Nb<sub>2</sub>O<sub>5</sub> (0.5:0.5) at different calcination temperatures (a. 400 °C ,b. 500 C and c. 600 °C)

| Average partial<br>size/ nm | FWHM(deg) | 2Theta(deg) | d (A)   | Temperature/ °C |
|-----------------------------|-----------|-------------|---------|-----------------|
| 45.80                       | 0.18330   | 27.7395     | 3.21339 | 400             |
| 42.20                       | 0.18810   | 46.0461     | 1.96957 | 400             |
| 45.95                       | 0.19440   | 54.5780     | 1068012 |                 |
| 26.23                       | 0.31300   | 29.0675     | 3.06953 |                 |
| 30.15                       | 0.26330   | 30.3817     | 2.93968 | 500             |
| 23.64                       | 0.34450   | 25.8517     | 3.44361 |                 |
| D =25.40                    | 0.32300   | 29.0311     | 3.07330 |                 |
| D = 32.04                   | 0.24790   | 30.3607     | 2.94167 | 600             |
| D = 23.85                   | 0.34170   | 25.8261     | 3.44696 |                 |

Table 1: The effect of different calcination temperatures of the Couple of Sb<sub>2</sub>O<sub>3</sub>/Nb<sub>2</sub>O<sub>5</sub> at (0.5:0.5) on the average partial size.

#### **Photocatalytic experiments**

#### Effect of the mass couple Nb<sub>2</sub>O<sub>5</sub>/ sb<sub>2</sub>O<sub>3</sub> on the photodegradation of orange G dye.

This factor was studied by using different masses of couple under optimum conditions including the use constant concentration of orange G dye (10ppm), temperature at 298.15 K and normal pH for dye 6.8. The rate constant of photodegradation process increased with weight increase the maximum degradation (86.71%) of the orange G at couple amount 0.15 g. After that weight the percentage of degradation was decreased The rate of photocatalytic was increase with increase the amount of couple this due to increase the active site on the surface of couple Nb<sub>2</sub>o<sub>5</sub>/sb<sub>2</sub>o<sub>3</sub>, but increase the amount of couple causes the decrease in the efficiency of photodegradation. At higher values of masses of the used couple this inhibition may be due to successive layers of molecules couple that prevent light from passing through other layer as Figure 3.<sup>11,19,20</sup>.

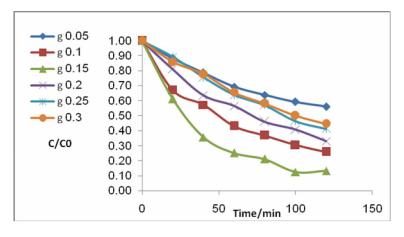


Figure 3:. Effect of the weight of couple Nb<sub>2</sub>O<sub>5</sub>/Sb<sub>2</sub>O<sub>3</sub> on photodegradation of orange G dye.

The results illustrated in Table 2.and plotted in Figure 4. which shows the pseudo first order reaction curve for various weights of couple  $Nb_2O_5/Sb_2O_3$ .

| Table 2 :Effect | of different | weights on rate con | stant. |
|-----------------|--------------|---------------------|--------|
|-----------------|--------------|---------------------|--------|

| Weight/gm | k/min   |
|-----------|---------|
| 0.05      | 0.00053 |
| 0.1       | 0.0121  |
| 0.15      | 0.0178  |
| 0.2       | 0.0094  |
| 0.25      | 0.0074  |
| 0.3       | 0.0068  |

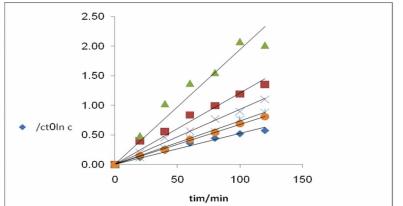


Figure 4: The change of lnC<sub>0</sub>/C at different weights with reaction time.

#### Effect of pH of reaction mixture on dye removal

It is found that many factors influence the effect of pH of reaction mixture on dye removal such as ionization state of the surface, nature of the dye, particle size and ability of molecular adsorbed adsorption on the catalyst surface<sup>21</sup>. the degradation reaction of orange G dye carried out at the pH range between 2-9 at optimum condition 10ppm dye concentration, 0.15 g catalyst amount and temperature equal to 298.15 K. the acid-base of the solution was adjusted by using HCl and NaOH prepare solutions . from measure the absorbance for different pH .It possible can observe the rate constant of different pH value from Table 3.

Table 3: The change of rate constant with different pH value.

| рН  | K*10 <sup>-3</sup> min <sup>-1</sup> |
|-----|--------------------------------------|
| 2.0 | 5.5                                  |
| 3.0 | 7.4                                  |
| 5.0 | 10.1                                 |
| 6.8 | 19.5                                 |
| 9.0 | 4.6                                  |

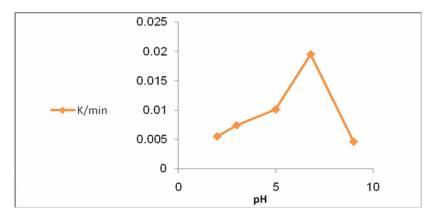


Figure 5: Effect of the pH solution on the rate constant reactions.

From Figure 5. It can be noted that rate constant was increased when the pH increase from 2 to 6.8 after that the rate constant of dye initial down on this behavior similar with literatures<sup>22-26</sup>. The reason for this result the medium of acidic pH solution causes to more  $H^+$  ions are available for the adsorption to mask the surface of the catalyst therefore preventing the photo excitation of catalyst particles , thereby reducing the generation of free radicals .

# Conclusion

From the results it was found that at the 1:1 percentage at calcination temperature 600 °C is more active than other calcination temperature. Formation the new couple  $Nb_2O_5/Sb_2O_3$  of semiconductors was investigated by x-ray diffraction patterns. The ability for photodegradation of orange G dye 86.74% in the optimum condition amount of couple 0.15 g, concentration dye 10 ppm, temperature 298.15 K and normal pH 6.8 of dye solution.

#### References

- 1. Verma A., and Singh P. K., 2013. Sol-gel derived nanostructured niobium pentoxide thin films for electrochromic applications, Indian Journal of Chemistry 52A: 593-598
- 2. Behnajady, M. A.; Modirshahla, N.; Shokri, M., 2004. Photodestruction of Acid Orange 7 (AO7) in aqueous solutions by UV/H2O2: influence of operational parameters, Chemosphere, 55: 129-134.
- 3. Algubili A. M., Alrobayi E. M., and Alkaim A. F., 2015. Photocatalytic degradation of remozal briliant blue dye by ZnO/UV process, Int. J. Chem. Sci., 13: 911-921.
- Tayade, R. J.; Natarajan, T. S.; Bajaj, H. C., 2009. Photocatalytic Degradation of Methylene Blue Dye Using Ultraviolet Light Emitting Diodes, Industrial & Engineering Chemistry Research, 48: 10262-10267.
- 5. Al-Gubury H. Y., Fairooz N. Y., Aljeboree A. M., Alqaraguly M. B., and Alkaim A. F., 2015. Photocatalytic degradation n-undecane using couple ZnO-Co2O3, Int. J. Chem. Sci., 13: 863-874.
- Gnanaprakasam, A.; Sivakumar, V. M.; Sivayogavalli, P. L.; Thirumarimurugan, M., 2015. Characterization of TiO2 and ZnO nanoparticles and their applications in photocatalytic degradation of azodyes, Ecotoxicology and Environmental Safety 121: 121-125.
- 7. Karam F. F., Kadhim M. I., and Alkaim A. F., 2015. Optimal conditions for synthesis of 1, 4naphthaquinone by photocatalytic oxidation of naphthalene in closed system reactor, Int. J. Chem. Sci., 13: 650-660.
- Pitchaimuthu, S.; Rajalakshmi, S.; Kannan, N.; Velusamy, P., 2014. Enhanced photocatalytic activity of titanium dioxide by β-cyclodextrin in decoloration of Acid Yellow 99 dye, Desalination and Water Treatment, 52: 3392-3402.
- 9. Alkaim, A. F., Dillert, R., and Bahnemann, D. W., 2015. Effect of polar and movable (OH or NH2 groups) on the photocatalytic H2 production of alkyl-alkanolamine: a comparative study, Environ. Technol., 36: 2190–2197.
- 10. El-Mekkawi, D.; Galal, H. R., 2013. Removal of a synthetic dye "Direct Fast Blue B2RL" via adsorption and photocatalytic degradation using low cost rutile and Degussa P25 titanium dioxide, Journal of Hydro-environment Research, 7: 219-226.
- 11. Sobana, N.; Krishnakumar, B.; Swaminathan, M., 2013. Synergism and effect of operational parameters on solar photocatalytic degradation of an azo dye (Direct Yellow 4) using activated carbon-loaded zinc oxide, Mat. Sci. Semicon. Proc., 16: 1046-1051.
- 12. Kandiel T. A., Robben L., Alkaim A., and Bahnemann D.,2013. Brookite versus anatase TiO<sub>2</sub> photocatalysts: phase transformations and photocatalytic activities, Photochem. Photobiol. Sci., 12: 602-609.
- 13. Thennarasu G., Kavithaa S., and Sivasamy A., 2012. Photocatalytic degradation of orange G dye under solar light using nanocrystalline semiconductor metal oxide, Environ. Sci. pollut. Res., 19: 2755-2765.
- 14. Chenini H., Djebbar K., Zendaoui S. M., Sehili T., and Zouchoune B., 2011. Removal of an Azo dye (orange G) by various methods in homogeneous phase . comparative study, Jordan Journal of chemistry, 6: 307-319.
- 15. Sun J., Qiao L., Sun S., and Wang G., 2008. Photocatalytic degradation of Orange G on nitrogen-doped TiO2 catalysts under visible light and sunlight irradiation, J. Hazard. Mater., 155: 312-319.
- 16. Tijani J., Fatoba O., Madzivire G., and Petrik L., 2014. A Review of Combined Advanced Oxidation Technologies for the Removal of Organic Pollutants from Water, Water. Air. Soil. Pollut., 225: 1-30.
- 17. Mohamed F. F., Allah P. M., Mehdi A. P., and Baseem M., 2011. Photoremoval of Malachite Green (MG) using advanced oxidation process, Res.J.Chem.Environ., 15: 65-70.
- 18. Mika E.T. Sillanpää, Tonni Agustiono Kurniawan, and Wai-hung Lo 2011. Degradation of chelating agents in aqueous solution using advanced oxidation process (AOP), Chemosphere, 83: 1443-1460.

- 19. Kaur, J.; Singhal, S., 2014. Heterogeneous photocatalytic degradation of rose bengal: Effect of operational parameters, Physica B, 450: 49-53.
- Van Doorslaer, X.; Heynderickx, P. M.; Demeestere, K.; Debevere, K.; Van Langenhove, H.; Dewulf, J., 2012. TiO<sub>2</sub> mediated heterogeneous photocatalytic degradation of moxifloxacin: Operational variables and scavenger study, Appl. Catal. B-Environ., 111-112: 150-156.
- Alkaim A. F., Aljeboree A. M., Alrazaq N. A., Baqir S. J., Hussein F. H., and Lilo A. J., 2014. Effect of pH on Adsorption and Photocatalytic Degradation Efficiency of Different Catalysts on Removal of Methylene Blue, Asian Journal of Chemistry, 26: 8445-8448.
- 22. Sohrabi, M. R.; Ghavami, M., 2008. Photocatalytic degradation of Direct Red 23 dye using UV/TiO2: Effect of operational parameters, Journal of Hazardous Materials, 153: 1235-1239.
- 23. Muruganandham, M.; Swaminathan, M., 2006. TiO2–UV photocatalytic oxidation of Reactive Yellow 14: Effect of operational parameters, Journal of Hazardous Materials, 135: 78-86.
- 24. Habibi, M. H.; Hassanzadeh, A.; Mahdavi, S., 2005. The effect of operational parameters on the photocatalytic degradation of three textile azo dyes in aqueous TiO2 suspensions, Journal of Photochemistry and Photobiology A: Chemistry, 172: 89-96.
- 25. Daneshvar, N.; Salari, D.; Khataee, A. R., 2003. Photocatalytic degradation of azo dye acid red 14 in water: investigation of the effect of operational parameters, Journal of Photochemistry and Photobiology A: Chemistry, 157: 111-116.
- 26. Palmer F. L., Eggins B. R., and Coleman H. M., 2002. The effect of operational parameters on the photocatalytic degradation of humic acid, J. Photochem. Photobiol. A, 148: 137-143.

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