

# Photocatalytic degradation of reactive green dye by using Zinc oxide

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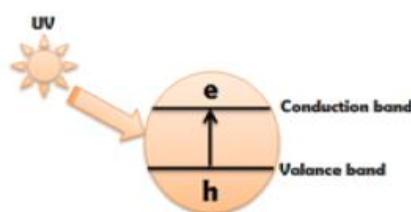
## ABSTRACT

Photocatalytic degradation of reactive green using zinc oxide have been included in this work, which is achieved by the irradiation of suspended solution consists of different concentration of reactive blue with 0.12gm/100cm<sup>3</sup> of mixed metal oxide semiconductor zinc oxide by using solar light. In order to study the effect of metal oxide zinc oxide in photocatalytic degradation of reactive blue, several experiments were achieved. Such as the effect of dye concentration, the effect of mass loading and effect of solution temperature.

**KEY WORDS:** Photocatalytic; Reactive green dye, Kinetic.

## 1. INTRODUCTION

In last decay, industry revolution in most of the word cause environmental contaminates. Huge types of dye uses led to pollute the surface water and groundwater., there are different methods used for water treatment in our previous work (Mashkour, 2011; Mohamed, 2011; Aljebori and Alshirifi, 2012; Alkaim, 2012, 2013, 2015; Kandiel, 2013; Alsalman, 2013; Jasim, 2013; Al-Robayi, 2014; Aljeboree, 2014; Hadi, 2014; Jabbar, 2014; Al-Gubury, 2015; Al-Gubury, 2015; Al-Gubury, 2015; Algubili, 2015; Ali, 2015; Karam, 2015; Mohammed B Alqaragully, 2015; Abdulrazzak, 2016; Al-Terehi, 2016; Aljeboree, 2016; Fairouz, 2016; Kamil, 2016). Advanced oxide processes (AOP) method can be used to wastewater treatment. It has been ability to convert the pollutants into the harmless materials (Hussein, 2008; Hussein, 2013; Jafar, 2013; Kami, 2014; Al-Khafaji, 2015). In (Al-gubury, 2016) hydroxyl radicals ( $\cdot\text{OH}$ ), this reactive radical capable of mineralizing organic pollutants(Saleh, 1974; Alkhateeb, 2005; Hussein, 2007; Hussein 2010; Matloob, 2011; Mohamed, 2011; Al-gubury, 2016). Zinc oxide semiconductor as a catalyst has the ability to remove from the wastewater a number of pollutants such as pesticides(Hussein, 2010). If the semiconductor zinc oxide irradiated by UV lamp, the electrons will promote from valance band to conduction band, producing electrons in conduction band undergo photo reduction will leave a positive hole in valance band inter photo oxidation processes.



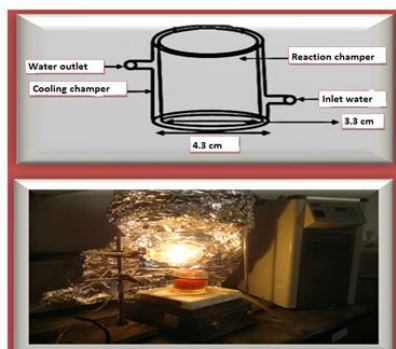
**Figure.1.General mechanism of the photo catalysis on zinc oxide particle**

## 2. MATERIALS AND METHODS

**A-Chemicals:** 1-Zinc oxide (ZnO): purity (99%), particle size (100) mesh, supplied by Fluka AG.

2 – Reactive green supplied by sigma – Aldrich.

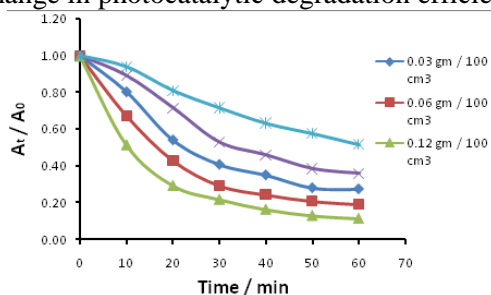
**B - Photo reactor and Procedure:** 1-Experiments were carried out in glass photochemical reactor as shown in Fig. 2-In the outside thimble; running water was passed through the thimble to cool the reaction solution. Owing to the continues cooling done by controlled water bath, in the inside the reaction vessel will consist of 100 ml of sample under irradiation.



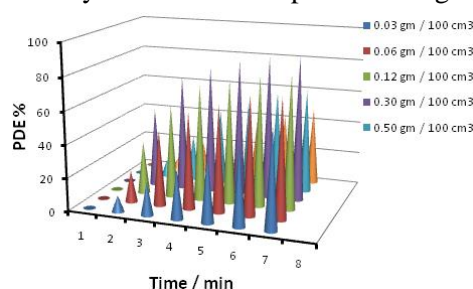
**Figure.2.Main parts of the photocatalytic cell used in Photocatalytic degradation of reactive green**

### 3. RESULTS AND DISCUSSION

**The Effect of zinc oxide mass on photo catalytic degradation of reactive green:** The effect of mass of zinc oxide on Photocatalytic degradation of reactive green, was studied using 40 ppm of reactive blue, flow rate of air 10ml/min, room temperature 298 K, Figure 3 shows different loaded mass of zinc oxide have been tested by the photo catalytic degradation processes of reactive green dye, as the masses zinc oxide increases until reach to the mass zinc oxide 0.12 gm/100cm<sup>3</sup> photocatalytic degradation of reactive green gradually increases then photocatalytic efficiency decreased. This may be attributed to the light absorption will be limited only to the first layers of reactive green and the other layers of solution do not receive light photons. Moreover at higher zinc oxide loading a light scattering will attained, this lead to decrease the photon intensity (Krishnakumara, 2011). The mass of zinc oxide 0.12gm/100cm<sup>3</sup> gives the optimum photocatalytic degradation efficiency which is equal to 88.68 %. The results of the change in photocatalytic degradation efficiency (P.D.E%) with catalyst concentration plotted in Figure 4 .

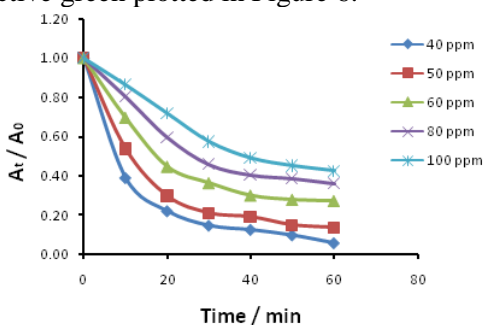


**Figure.3.The effect masses of zinc oxide on Photocatalytic degradation of reactive green**

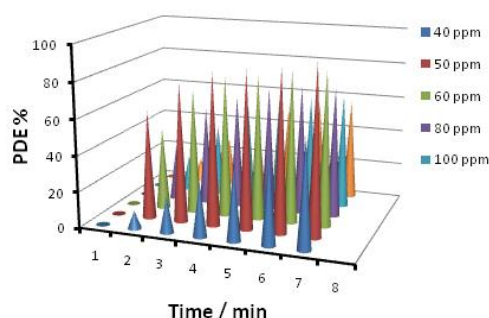


**Figure.4.The change of Photocatalytic Degradation Efficiency with irradiation time of different types of catalyst**

**The Effect of initial reactive green concentration on photo catalytic degradation processes :** A series of experiments have been done , the effect of change initial (40 – 100 ppm) on photocatalytic degradation process of reactive green was studied using 0.12gm / 100 cm<sup>3</sup>, the light intensity equal to 8.22 mW/cm<sup>2</sup>, and temperature equal to 298 K . The results are plotted in Figure 5. It has been observed that the rate of photocatalytic degradation gradually decreases with the increasing of initial reactive green concentration. This behaviour could be explained, the concentration 40ppm was the optimum concentration to cover the largest area of the zinc oxide particles, therefore absorbed maximum exciting photons to generate higher concentration of the activated titanium dioxide semiconductor. Another reason for this behaviour is the strong absorption of light by the reactive green in the sample which contain high concentration that 40ppm, reactive green on 0.12 gm /100cm<sup>3</sup> of zinc oxide. The excess of reactive green prevent the penetration of light through the successive layers of reactive green on the zinc oxide surface is weak to generate the required excited state of the reactive green on zinc oxide(Miao, 2014). The concentration of reactive green 40ppm gives the optimum photocatalytic degradation efficiency which is equal to 94.14%. The results of the change in percentage photocatalytic degradation efficiency (P.D.E %) with concentration of reactive green plotted in Figure 6.

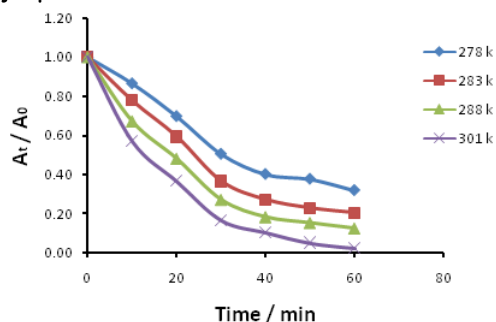


**Figure.5.The change of ( At / A0 )with irradiation time at concentration of reactive green**

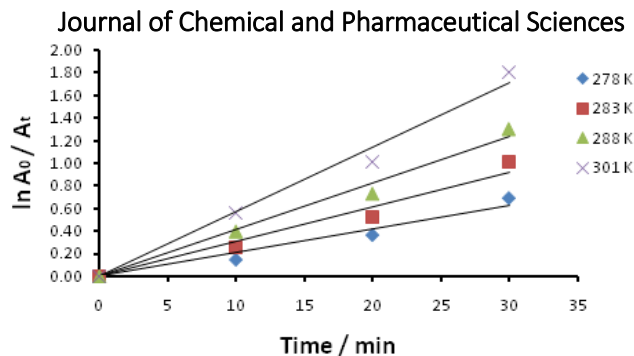


**Figure.6.The change of percentage Photocatalytic Degradation Efficiency (PDE %) with irradiation time of different types of concentration of reactive green.**

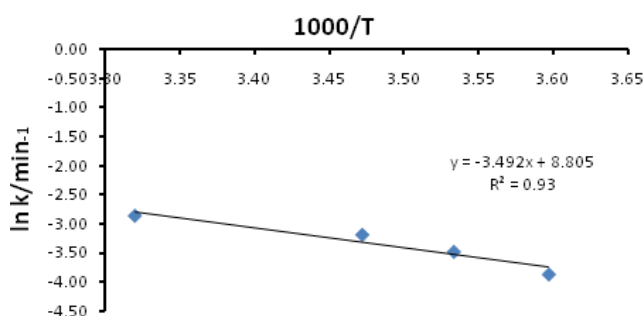
**Effect of temperature on photocatalytic degradation of reactive green:** A series of experiments were carried out to study the effect of temperature on Photocatalytic degradation rate of reactive green using zinc oxide suspension at different temperature ranging from (278 – 301 k)(Ahmed, 2011). Figure 7, shows the effect of temperature on the photo catalytic degradation rate of reactive green at fixed initial concentration 40 ppm and 0.12 gm/100ml of zinc oxide as catalyst . Figure 7 indicate that the photo catalytic degradation rate of reactive green increases with increase of temperature because increase temperature cause to increase generate free radicals and this lead to decrease recombination process. From the Figure 9 the amount of activation energy 29.04 kJ / mole using Arrhenius equation.



**Figure.7.** The change of  $(A_t / A_0)$  with irradiation time at different temperature for photocatalytic degradation rate of reactive green B using zinc oxide



**Figure.8.** The change of  $\ln(A_0 / A_t)$  with irradiation time at different temperature



**Figure.9.** The change of  $\ln K$  against  $1000/T$

#### 4. CONCLUSION

The compound has been not degraded in case of absent of catalyst. The compound has been successfully degraded when used the catalyst with the light. The optimum condition for the Photocatalytic degradation of reactive green equal 0.12gm/100ml mass of zinc oxide and 40ppm concentration of reactive green and 8.22 mW/cm<sup>2</sup>. The activation energy = 29.04 J/mole. The percentage efficiency of degradation reactive green equal 94.14%.

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