

# Study of Effect of Temperature, Light intensity and Irradiation time during Decolouration Process of Methyl Green Dye

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**Abstract :** Methyl green is the oldest known synthetic groups. Most basic dyes of this series are beautiful crystalline Compounds with a reflex the colour of which is often complementary to the colour in solution. Methyl green a triarylmethane dye providing toxic effluents can be highly degraded using  $\text{TiO}_2$  catalyst. Photo catalytic degradation of methyl green dye have been studied with the help of variety of parameters which are effect of temperature, light intensity and irradiation time. The influence of temperature has been studied in the range from  $30^\circ\text{C}$  to  $55^\circ\text{C}$ . The rate constant increased with the increase in the light intensity.

**Keywords –** Methyl green, Toxic, Catalyst, Temperature, Light Intensity.

**Introduction -** It is well Known that dyes and their degradation by Products originated through oxidation, hydrolysis reactions are highly carcinogenic (1) These substances are highly toxic, Stable to natural decomposition. Decomposition of dye effluents has therefore required increasing attention (2) Earlier studies have shown that a wide range of organic substrates can be completely photo mineralized in the presence of  $\text{TiO}_2$  (3) Aqueous degradation of the commonly used textile dye methyl green has already been studied using heterogeneous catalysts such as  $\text{TiO}_2$  with artificial visible light source. The aim of this paper is to study the effect of temperature, light intensity and irradiation time during Decolouration process of Methyl green dye.

**Experimental:** Methyl green was obtained from Loba Chemie. Photo catalyst  $\text{TiO}_2$  was obtained from the S.D. Fine Company. All Solutions were prepared in doubly distilled water. Photo catalytic experiments were carried out with 50 ml of dye solution ( $3.8 \times 10^{-5} \text{ mol dm}^{-3}$ ) using 300mg of  $\text{TiO}_2$  photo catalytic under exposure to visible irradiation in specially designed double-walled slurry type batch reactor vessel made up of Pyrex glass (7.5 cm height, 6 cm diameter) surrounded by thermostatic water circulation arrangement to keep the temperature in the range of  $30 \pm 0.3^\circ\text{C}$ . Irradiation was carried out using 500 w halogen lamp surrounded by aluminum reflector to avoid irradiation loss. During photo catalytic experiments after stirring for 10 min slurry composed of dye solution and catalyst was placed in dark for  $\frac{1}{2}$  h in order to establish equilibrium between adsorption and desorption phenomenon of dye molecule on photo catalyst surface. Then slurry containing

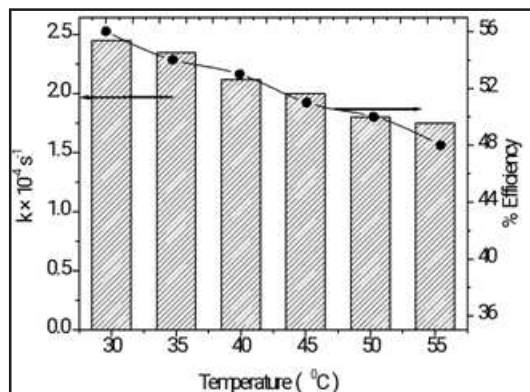
aqueous dye solution and  $\text{TiO}_2$  was stirred magnetically to ensure complete suspension of catalyst particle while exposing to visible light. At specific time intervals aliquot (3ml) was withdrawn and centrifuges for 2 min at 3500 rpm to remove  $\text{TiO}_2$  particle from aliquot to assess extent of decolourisation photo metrically. Changes in absorption spectra were recorded at 480 nm on double beam UV-Vis, spectrophotometer (Systronic Model No. 166) Intensity of visible radiation was measured by a digital luxmeter (Lutron LX 101). pH of solution was measured using a digital pH meter.

## Results and Discussion:

**Effect of temperature:** One of the advantages of photoreaction is that it is not affected or slightly affected by temperature change (4, 5). The influence of temperature has been studied in the range from  $30^\circ\text{C}$  to  $55^\circ\text{C}$ . The results are shown in Table 1 and Fig. 1. Increase in temperature led to decrease the rate of degradation. This gradual decrease in the reaction rate values could be attributed to the following reasons: the adsorption rate decreased with increasing temperature because the adsorption is a heat releasing process, increase in reaction temperature tend to increase electron-hole recombination and with increase in temperature the solubility of oxygen in water decreased (6).

**Table 1: Effect of temperature:**  $[\text{MG}] = 2.5 \times 10^{-5} \text{ mol dm}^{-3}$ , pH = 10.0  
 $\text{TiO}_2 = 100 \text{ mg/ } 10 \text{ mL}$ , Light intensity =  $20 \times 10^3 \text{ lux}$ ,  
 Temperature =  $30 \pm 0.3^\circ\text{C}$ .

Temperature ( $^{\circ}\text{C}$ )	$k \times 10^{-4} \text{ s}^{-1}$	$t_{1/2} \times 10^3 \text{ s}$
30	2.45	2.82
35	2.35	2.94
40	2.12	3.26
45	2.00	3.46
50	1.80	3.85
55	1.75	3.96



**Fig. 1: Effect of temperature**

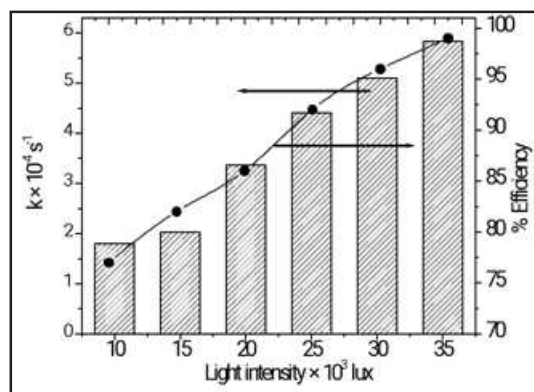
#### Effect of light intensity and irradiation time:

Photocatalytic reaction rate depends largely on the radiation absorption of the photocatalyst. Research studies revealed increase in the degradation rate, with increase in light intensity during photocatalytic degradation. The influence of light intensity on the rate of degradation has been examined at constant dye concentration ( $2.5 \times 10^{-5} \text{ mol dm}^{-3}$ ) and catalyst loading (100 mg/ 100 mL). Rate constant increased from  $1.80 \times 10^{-4} \text{ s}^{-1}$  to  $5.83 \times 10^{-4} \text{ s}^{-1}$  on increase light intensity from  $10 \times 10^3 \text{ lux}$  to  $35 \times 10^3 \text{ lux}$ .

**Table 2: Effect of light intensity:** [MG] =  $2.5 \times 10^{-5} \text{ mol dm}^{-3}$ , pH = 10.0

$\text{TiO}_2$  = 100 mg/ 100 mL, Light intensity =  $20 \times 10^3 \text{ lux}$ , Temperature =  $30 \pm 0.3 \text{ }^{\circ}\text{C}$ .

Light intensity $\times 10^3 \text{ lux}$	$k \times 10^{-4} \text{ s}^{-1}$	$t_{1/2} \times 10^3 \text{ s}$
$10 \times 10^3$	1.80	3.85
$15 \times 10^3$	2.03	3.41
$20 \times 10^3$	3.37	2.05
$25 \times 10^3$	4.41	1.57
$30 \times 10^3$	4.49	1.54
$35 \times 10^3$	5.83	1.18



**Fig. 2: Effect of light intensity**

**Conclusion :** The influence of temperature has been studied in the range from  $30^{\circ}\text{C}$  to  $55^{\circ}\text{C}$ . The rate constant was decreased with increase in temperature photo catalytic reaction rate depends largely on the radiation absorption of the photo catalyst. Rate constant increased with increase in light intensity.

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#### References:-

1. Nam S., Tratnyek P.G. 2000 Reduction of azo dyes with zero valent iron. Water Res. 1837 -1845.
2. Brown M.A., Vito SC. Crit Rev. Environ. Sci. Technol 1993 – 249 – 32.
3. Blake DM Bibliography of work on the photo catalytic removal of hazardous compounds from Water and air USA : National Renewal Energy Laboratory 2001, – 253.
4. Herrera F., Kiwi J., Lopez A., and Nadtochenko V., Environ. Sci. Technol., 1999, 3145.
5. Amiri A. S., Bolton J. R. and Caster S. R., J. Adv. Oxid. Technol., 1996, 18.
6. Jiang Y., Sun Y., Liu H., Zhu F. and Yin H., Dyes and Pigments, 2008, 77.

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