

Chongdian Si¹, Jing Zhou², Yuhong Sun¹,
Guangjun Liu¹, Hongtao Gao¹

**KINETICS OF PHOTODEGRADATION OF ALIZARIN
GREEN IN AN ACOUSTIC FLUIDIZED
BED USING TiO₂ CATALYST**

¹ Jining University, Qufu, China;

² Department of Energy and Material Engineering,
Shandong Career Development College, Jining, China
chongdiansi@126.com

The kinetics of photodegradation of alizarin green (AG) in an acoustic fluidized bed was investigated using Degussa P25 catalyst and N-doped TiO₂ catalyst, respectively. The effects of initial concentration of AG, catalyst amount, initial pH value and liquid flow rate on the photodegradation rates were the main goal of present study. The degradation rate increased with increasing amount of photocatalyst, pH value and liquid flow rate, reached a maximum and then decreased. The degradation rate decreased with increasing initial concentration. According to the Langmuir-Hinshelwood model, the disappearance of the AG followed pseudo-first-order kinetics. Experimental results showed that AG can be degraded effectively by Degussa P25 catalyst and N-doped TiO₂ catalyst in an acoustic fluidized bed reactor. N-doped TiO₂ catalyst has high and significant photocatalytic activity and in comparison with Degussa P25 catalyst was photocatalyst.

Keywords: TiO₂ catalyst, photodegradation, fluidized bed, acoustic.

Introduction

Advanced oxidation processes (AOP) are recognized as a high and significant alternative to transform and decontaminate the dye-containing waste water [1]. These processes named AOP all rely on the generation of highly reactive ·OH radical as the main oxidative species for the potential destruction and conversion of organic pollutants into harmless substances [2, 3]. Photocatalysis is one typical example of AOP for the degradation of dye-containing and other organic pollutants in water [4, 5].

© Chongdian Si, Jing Zhou, Yuhong Sun, Guangjun Liu, Hongtao Gao, 2016

References

- [1] Cheng H., Chou S., Chen S., Yu C. // *J. Environ. Sci.*, 2014, 26, P. 1307 – 1312.
- [2] Azimi Y., Allen D.G., Farnood R.R. // *Water Res.*, 2014, 54, P. 179 – 187.
- [3] Mohapatra D.P., Brar S.K., Tyagi R.D., Picard P., Surampalli R.Y. // *Sci. Total Environ.*, 2014, 470/471, P. 58 – 75.
- [4] Kochkodan V.M., Rolya E.A. Goncharuk V.V. // *J. Water Chem. and Technol.*, 2009, 31, P. 227 – 237.
- [5] Liu P., Zhang H., Feng Y., Yang F., Zhang J. // *Chem. Eng. J.*, 2014, 240, P. 211 – 220.
- [6] Wooseok N., Woo K., Han G.Y. // *J. Ind. Eng. Chem.*, 2009, 15, P. 348 – 353.
- [7] Su C.C., Chen C.M., Anotai J., Lu M.C. // *Chem. Eng. J.*, 2013, 222, P. 128 – 135
- [8] Roberto L.P., Brandi R.J., Cassano A.E. // *Chem. Eng. Sci.*, 2010, 65, P. 1345 – 1353.
- [9] Wang R.C., Fan K.S., Chang J.S. // *J. Taiwan Inst. Chem. Eng.*, 2009, 40, P. 533 – 540.
- [10] Dong S.S., Zhou D., Bi X.T. // *Particuology*, 2010, 8, P. 60 – 66.
- [11] Si C.D., Zhou J., Guo Q.J. // *Int. J. Chem. React. Eng.*, 2012, 10, P. 1 – 13.
- [12] David R.E., Theodore J.H. // *Chem. Eng. J.*, 2013, 243, P. 411 – 420.
- [13] Gao H.T., Si C.D., Zhou J., Liu G.J. // *J. Taiwan Inst. Chem. Eng.*, 2011, 42, P. 108 – 113.
- [14] Xiong Z.D., Xu A.H., Li H.Y., Ruan Xi.C., Xia D.S., Zeng Q.F. // *Ind. Eng. Chem. Res.*, 2013, 2, P. 362 – 369.
- [15] Satoshi K., Li N., Ito K., Katsumata H., Suzuki T., Ohta K. // *Chem. Eng. J.*, 2009, 148, P. 50 – 56.
- [16] Pierre F.B., Bouzaza A., Wolbert D. // *Environ. Sci. Technol.*, 2007, 41, P. 2908 – 2914.
- [17] Hao X.G., Li H.H., Zhang Z.L. // *Chem. Eng. Res. Des.*, 2009, 87, P. 1604 – 1611.

Recieved 14.04.2014