Role of Water and *p*-Fluorobenzoic Acid in Mn(II)T(*p*-Cl)PP Catalyzed Aerobic Oxidation of *p*-Fluorotoluene

YONG-QIN FANG*, CHUN-XU LV and MING LU

School of Chemical Engineering, Nanjing University of Science and Technology, Nanjing, Jiangsu Province, P.R. China

*Corresponding author: Fax: +86 519 86330198; Tel: +86 519 86330180; E-mail: fyq_cczu@126.com

(Received: 4 November 2011;

Accepted: 23 July 2012)

AJC-11870

A solvent-free manganoporphyrin-catalyzed oxidation of *p*-fluorotoluene was developed, using oxygen as a clean and cheap oxidant. The parameters that have relationship with conversion were investigated. The conversion of *p*-fluorotoluene was found to be depend on the stability of catalytic environment. Water and *p*-fluorobenzoic acid were the final oxidation products of *p*-fluorotoluene, would inhibit the activity of catalyst. The conversion would be increased from 7.0 to 16.5 %, by the dehydration of acetic anhydride.

Key Words: *p*-Fluorotoluene, Manganoporphyrin, Water, *p*-Fluorobenzoic acid.

INTRODUCTION

The oxidation derivatives of *p*-fluorotoluene play important roles in the field of medicine manufacture, such as paroxetine, rosuvastatin, mosapride citrate¹⁻³, etc.

Nowadays, these oxidation derivatives are synthesized by the chlorination of *p*-fluorotoluene followed by hydrolysis in industry. However, the corrosion of reactors and the serious environment threat have become the major drawbacks⁴. During the past decades, metalloporphyrins have been widely used as cytochrome P-450 models catalyzing the oxidation of various substrates in association with a wide variety of oxygen donors such as iodosylarenes⁵, sodium periodate⁶, hydrogen peroxide⁷, etc.

However, because of the inertness of saturated hydrocarbon, few reports are available for the oxidation of toluene using molecular oxygen as oxidant⁸⁻¹⁰. Furthermore, the electrophilic reactivity of high valence metalloporphyrin could be decreased due to the electron-withdrawing effect of fluorine atom.

Guo and co-workers have demonstrated a solvent-free biomimetic oxidation system to the aerobic oxidate of toluene, which used simple metalloporphyrin without any co-reductants⁸. However, the application of this method in industry is hindered due to the low conversion. Although the catalysts can be recycled by supported metalloporphyrins on different carriers^{9,10}, the conversion of toluene was limited to 10 %. Thus, the development of a efficiency procedure for such transformation remains a highly desired goal for organic chemists. Herein, we report the Mn(II)T(*p*-Cl)PP-catalyzed biomimetic oxidation of *p*-fluorotoluene using oxygen. This work may

provide a better understanding of the catalytic behavior of metalloporphyrin in oxidation reaction.

EXPERIMENTAL

Gas chromatography analysis was performed on a Fuli-9790 chromatograph equipped with a 30 m × 0.25 mm × 0.25 μm SE54 capillary column and a flame ionization detector. MS spectra were measured on a Shimadzu GCMSQP2010. *p*-Fluorotoluene was distilled and analyzed by GC before use. Mn(II)T(*p*-Cl)PP was synthesized and purified according to documented procedures¹¹. Other reagents and solvents were all analytical grade.

25 mL of *p*-fluorotoluene and a measured amount of Mn(II)T(*p*-Cl)PP were charged into a 100 mL autoclave reactor with nitrogen at 0.85 MPa. The reaction mixture was heated to 180 °C under magnetic stirring. The oxygen was injected to replace the nitrogen three times and then charged to 0.85 MPa. The loss of pressure was recorded every hour and the system was recharged to 0.85 MPa. After the loss of pressure was less than 0.01 MPa per hour, the mixture was cooled, neutralized by 1 mol/L NaOH. *p*-Fluorobenzoic acid was obtained by filtering the water phase neutralized by 1 mol/L HCl. The organic phase was identified by GC-MS and quantified by GC using chlorobenzene as the internal standard. GC analysis was carried out as follows: nitrogen was used as the carrier gas, at the flow rate of 30 mL/min; the column temperature was 120 °C; the injector temperature was 200 °C; and the detector temperature was 200 °C.

RESULTS AND DISCUSSION

The oxidation of *p*-fluorotoluene by Mn(II)T(*p*-Cl)PP is shown in **Scheme-I**. The main oxidation products were

p-fluorobenzaldehyde and *p*-fluorobenzoic acid and only trace *p*-fluorobenzyl alcohol could be detected by GC-MS. The blank experiment confirmed that *p*-fluorotoluene could not be oxidized in the absence of Mn(II)T(*p*-Cl)PP.

Scheme-I: Oxidation of *p*-fluorotoluene catalyzed by Mn(II)T(*p*-Cl)PP

Effect of temperature on oxidation reaction: The effects of reaction temperatures on the conversion of *p*-fluorotoluene and the product selectivity are shown in Table-1.

TABLE-1 EFFECT OF TEMPERATURE ON OXIDATION REACTION			
Temperature (°C)	Selectivity (%)		Conversion (%)
	<i>p</i> -Fluorobenzal- dehyde	<i>p</i> -Fluoroben- zoic acid	
150	-	-	-
160	86.8	13.2	1.7
170	54.1	45.9	3.6
180	24.2	75.8	7.0
190	16.6	83.4	5.3
<i>p</i> -Fluorotoluene, 25 mL; catalyst, 0.8 mg (4.0×10^{-5} mol/L), pressure: 0.85 MPa			

As can be seen from Table-1, the reaction could not take place below 150 °C and the conversion increased until the temperature is elevated to 180 °C. Compared with the reported optimum reaction temperature of toluene in 160 °C⁸, the optimum temperature of *p*-fluorotoluene is much higher. This is at least partly due to the electron-withdrawing effect of fluorine atom. However, because metalloporphyrin is easily oxidized by oxygen under high temperature⁹, the improvement of conversion should be resolved by other pathways.

Effect of the amount of catalyst on oxidation reaction: The effects of the amount of catalyst on the the conversion of *p*-fluorotoluene and the product selectivity are listed in Table-2.

TABLE-2 EFFECT OF THE AMOUNT OF CATALYST ON OXIDATION REACTION			
Catalyst (mol/L)	Selectivity (%)		Conversion (%)
	<i>p</i> -Fluorobenzal- dehyde	<i>p</i> -Fluorobenzoic acid	
1.0×10^{-5}	32.5	67.5	1.5
2.0×10^{-5}	30.3	69.7	2.8
4.0×10^{-5}	24.2	75.8	7.0
6.0×10^{-5}	32.9	67.1	6.2
8.0×10^{-5}	36.7	63.3	4.5
<i>p</i> -Fluorotoluene, 25 mL; temperature, 180 °C; pressure, 0.85MPa			

The results indicate that the conversion of *p*-fluorotoluene increases significantly with the increasment of catalyst loading from 1.0×10^{-5} to 4.0×10^{-5} mol/L. But the conversion decreases with the further increasment of catalyst. This phenomenon is contrary to supported metalloporphyrins¹². A probable reason

is that superfluous catalyst in homogeneous system would collide with each other and convert to inactive dimer¹³.

Effect of water on oxidation reaction: It is well known that the hydroxylation of methyl group in biological systems via a free radical pathway¹⁴. As a free radical inhibitor, water would quench the reaction. In present case, the oxidation reaction could not run by adding 3 wt % of water. To further understand of this phenomenon, we studied the oxidation of *p*-fluorotoluene in the presence of acetic anhydride, as shown in Scheme-II. The main oxidation products were *p*-fluorobenzaldehyde, *p*-fluorobenzoic acid and *p*-fluorobenzyl acetate. The curves of pressure variations are shown in Fig. 1 and the effects of the amount of acetic anhydride on the conversion of *p*-fluorotoluene and the product selectivity are shown in Table-3.

Scheme-II: Oxidation of *p*-fluorotoluene in the presence of acetic anhydride catalyzed by Mn(II)T(*p*-Cl)PP

Fig. 1. Curves of pressure variations in different oxidation systems. System 1: *p*-fluorotoluene, 25 mL; catalyst, 0.8 mg, temperature, 180 °C; pressure, 0.85 MPa, System 2: *p*-fluorotoluene, 25 mL; acetic anhydride, 4.4 g; catalyst, 0.8 mg; temperature, 180 °C; pressure, 0.85 MPa

TABLE-3 EFFECT OF THE AMOUNT OF ACETIC ANHYDRIDE ON OXIDATION REACTION				
Acetic anhydride (wt %)	Selectivity (%)			Conversion (%)
	<i>p</i> - Fluorobenz- aldehyde	<i>p</i> - Fluoroben- zoic acid	<i>p</i> - Fluorobenzyl acetate	
0	24.2	75.8	-	7.0
10	17.8	71.8	10.6	13.1
15	18.3	68.9	12.8	16.5
20	17.1	67.8	15.2	5.4
50	18.0	65.7	16.3	3.2
<i>p</i> -Fluorotoluene, 25 mL; catalyst, 0.8 mg, temperature, 180 °C; pressure, 0.85 MPa				

The results indicate that *p*-fluorobenzyl alcohol can be protected by esterification with acetic anhydride and the conversion increases significantly with the increase of the amount of acetic anhydride from 0 to 15 wt %, because of the dehydration effect of acetic anhydride. However, the conversion decreases with further increase of acetic anhydride.

As can be seen from Fig. 1, in the oxidation system with acetic anhydride, the rate of oxygen consumption decreases rapidly in the initial 5 h, which is quite different to another system. This means that, although the conversion could be increased by removing water from the reaction system, there is still some fact inhibit the oxidation reaction. Owing to strong coordination effect, acetate could be used as an axial ligand and attached to metalloporphyrin¹⁵. However, the excessive axial ligand would decrease the catalytic activity of metalloporphyrin¹⁶. Hence, it is supposed that *p*-fluorobenzoic acid also played the same role as acetate in the oxidation reaction.

Effect of *p*-fluorobenzoic acid on oxidation reaction:

The effects of the amount of *p*-fluorobenzoic acid on the conversion of *p*-fluorotoluene and the product selectivity are listed in Table-4.

TABLE-4
EFFECT OF THE AMOUNT OF *p*-FLUOROBENZOIC ACID ON OXIDATION REACTION

<i>p</i> -Fluorobenzoic acid (g)	Selectivity (%)		Conversion (%)
	<i>p</i> -Fluorobenzaldehyde	<i>p</i> -Fluorobenzoic acid	
0.5	31.7	68.3	5.7
1.0	32.4	67.6	3.6
1.5	34.1	65.9	1.2
2.0	39.6	60.4	0.7
2.5	41.3	58.7	0.5

p-Fluorotoluene, 25 mL; catalyst, 0.8 mg, temperature, 180 °C; pressure, 0.85 MPa

As can be seen from Table-4, the conversion decreases with increase of the amount of *p*-fluorobenzoic acid. The oxidation reaction is nearly terminated when the amount of *p*-fluorobenzoic acid is up to 0.08 g/mL. This observation supported the aforementioned supposition.

Conclusion

This work studies the application of metalloporphyrin for the oxidation of *p*-fluorotoluene without any solvent or co-reductants. Water and *p*-fluorobenzoic acid, which are the final oxidation products of *p*-fluorotoluene would inhibit the catalytic activity of metalloporphyrine. Hence, not only the improvement of the structure of metalloporphyrin but also stability of catalytic system should be considered.

ACKNOWLEDGEMENTS

The authors acknowledged the assistance from Nanjing University of Science and Technology.

REFERENCES

1. W.R. Scott, T.S. Neville, L. Zhao and N. Ran, WO Patent 2009005647 (2009).
2. P. Benkic, D. Bevk, R. Lenarsic, S. Zupancic, A. Vajs and D. Jakse, WO Patent 2010081861 (2010).
3. M.S. Hashmi, M.K. Madhra and M.I. Inamdar, WO Patent 2011107903 (2011).
4. G. Billeb and P. Burg, US Patent 5347054 (1994).
5. T. Nakano, S. Kawabata, T. Sugihara, N. Agatsuma, H. Kakuda and Y. Mori, *Bull. Chem. Soc. Jpn.*, **76**, 2353 (2003).
6. M. Moghadam, S. Tangestaninejad, V. Mirkhani, I. Mohammadpoor-Baltork, N. Sirjanian and S. Parand, *Bioorg. Med. Chem.*, **17**, 3394 (2009).
7. P. Tagliatesta, D. Giovannetti, A. Leoni, M.G.P.M.S. Neves and J.A.S. Cavaleiro, *J. Mol. Catal. A: Chem.*, **252**, 96 (2006).
8. C.C. Guo, Q. Liu, X.T. Wang and H.Y. Hu, *Appl. Catal.*, **282**, 55 (2005).
9. G. Huang, F. Xiang, T.M. Li, Y.X. Jiang and Y.A. Guo, *Catal. Commun.*, **12**, 886 (2011).
10. G. Huang, J. Luo, C.C. Cai, Y.A. Guo and G.W. Luo, *Catal. Commun.*, **9**, 1882 (2008).
11. A.D. Adler, F.R. Longo, J.D. Finarelli, J. Goldmacher, J. Assour and L. Korsakoff, *J. Org. Chem.*, **32**, 476 (1967).
12. H. Kargar, M. Moghadam, V. Mirkhani, I. Mohammadpoor-Baltork, S. Tangestaninejad and M. Naghipour, *Polyhedron*, **30**, 1463 (2011).
13. J.E. Lyons, P.E. Ellis and H.K. Myers, *J. Catal.*, **155**, 59 (1995).
14. B. Meunier, S.P. Visser and S. Shaik, *Chem. Rev.*, **104**, 3947 (2004).
15. J. Poltowicz and J. Haber, *J. Mol. Catal. A: Chem.*, **220**, 43 (2004).
16. D. Mohajer and L. Sadeghian, *J. Mol. Catal. A: Chem.*, **272**, 191 (2007).