

Characterization of Flame-generated TiO₂ Deposited Activated Carbon Fibers (ACF)

Jae Hong Park ^{1,a}, Jun Ho Ji^{2,b}, Jeong Hoon Byeon ^{1,c}, Gyo Woo Lee ^{3,d}, and Jungho Hwang ^{1,e}

¹Department of Mechanical Engineering, Yonsei University, Seoul, 120-749, Korea

²Division of Digital Appliance Network, Samsung Electronics Co., Ltd., Suwon, 442-742, Korea

³Division of Mechanical and Aerospace System Engineering, Chonbuk National University, Jeonju, 561-756, Korea

^acheap@yonsei.ac.kr, ^bjunho.ji@samsung.com, ^cpostjb@yonsei.ac.kr, ^dgwlee@chonbuk.ac.kr, ^ehwangjh@yonsei.ac.kr (corresponding author)

Keywords: TTIP, TiO₂, Activated carbon fibers, Flame synthesis

Abstract. Even though activated carbon fibers (ACF) have been attractive due to high specific surface area and uniform micropore structure, there are only a few reports about the photocatalyst immobilization on ACF or their photodegradation behavior for removal of organic pollutants. In this study, ACF were selected as adsorptive support for photocatalyst immobilization. As photocatalysts, TiO₂ nanoparticles were synthesized by using a N₂-diluted and oxygen-enriched co-flow hydrogen diffusion flame. The visible flame length of 150mm was obtained by direct photographs. Flame temperature was measured by rapid insertion measurement technique with a R-type (Pt/Pt-13%Rd) thermocouple which was in wire shape (127μm diameter). TiO₂ particles were sampled by using a high temperature particle sampler and then were coated on ACF filters. The sampling was carried out at 70mm and 110mm above the burner. The structure of TiO₂ particles was analyzed by XRD. TiO₂ coated ACF filters were characterized by FESEM-EDX and BET analyses. TiO₂ particles on ACF filters were found to be agglomerated particles and the size of primary particles was approximately 50nm. The structure of TiO₂ particle was anatase-phase crystalline structure. The specific surface area of TiO₂ coated ACF filter was enhanced to be 1700~1860m²/g, depending on the sampling location, and the pores were micropores, regardless of the sampling location.

1. Introduction

Activated carbon fibers (ACF) are a type of highly microporous carbon materials [1], which are commercially available in the form of fiber tows, cloths (fabrics), papers, mats and felts [2]. ACF have a larger micropore volume and a more uniform micropore size distribution than granular activated carbons (GAC), and thus are considered to have a larger adsorption capacity and greater rates of adsorption and desorption. However, it is still interesting to enhance the pollutant reduction

activity of ACF, for example, by using a functional metal coating technology [3]. In this paper, we propose a coating method on ACF with flame synthesized TiO₂ nano particles.

2. Methods

Fig. 1 shows the experimental set up. As photocatalyst, TiO₂ nanoparticles were synthesized by using a N₂-diluted and oxygen-enriched co-flow hydrogen diffusion flame. TiO₂ particles were sampled by using a high temperature particle sampler and then were coated on an ACF filter which was located in the sampling duct. The sampling was carried out at 70mm and 110mm above the burner.

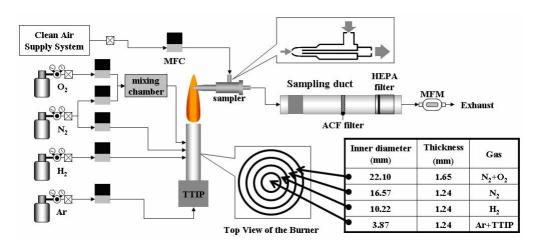
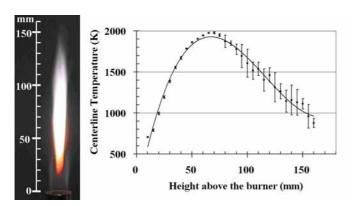
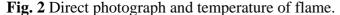


Fig. 1 Experimental setup.

The visible flame length of 150mm was obtained by direct photographs in Fig. 2. Flame temperatures were measured by rapid insertion measurement technique with an R-type (Pt/Pt-13%Rd) thermocouple which was in wire shape (127 μ m diameter) and the results are shown in Fig. 2. The structure of TiO₂ particles was anatase-phase crystalline structure (2 θ =25.5°) which was analyzed by XRD, as shown in Fig. 3.





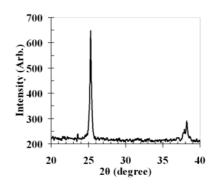


Fig. 3 XRD analysis of generated particle.

3. Results

TiO₂ coated ACF filters were characterized by field emission scanning electron microscopy (FESEM, JSM-6500F, JEOL, Japan), energy dispersive X-ray spectroscopy (EDX, JED-2300, JEOL, Japan) and BET analyses. TiO₂ particles on ACF filters were found to be agglomerated particles and the size of primary particles was approximately 50nm, as shown in Fig. 4. The concentration of TiO₂ particles which were sampled at 70mm was higher than that of particles sampled at 110mm, since the temperature at the location of 70mm was higher than the temperature at the location of 110mm.

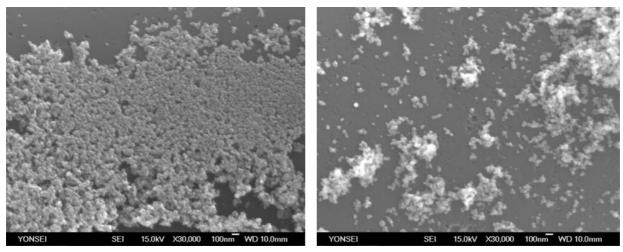


Fig. 4 FESEM images of TiO₂ particles on ACF filters at the sampling location of 70mm (left) and 110mm (right).

Elements of TiO₂ coated ACF filters were tabulated in table 1. TiO₂ coated filter of sampling location of 70mm had more Ti and O atoms than TiO₂ filter of sampling location of 110mm.

	ACF filter (sampling@70mm)		ACF filter (sampling@110mm)	
Element	mass%	atom%	mass%	atom%
С	89.61	93.79	91.64	94.86
О	6.66	5.23	5.75	4.47
Ti	3.73	0.98	2.61	0.68
Total	100	100	100	100

Table 1. Characteristics of ACF filters by FESEM-EDX

Specific surface area (m^2/g) and specific micropore area (m^2/g) of ACF filters were obtained by N_2 adsorption at 77 K by using a Micromeritics ASAP 2010. Fig. 5 shows that specific surface area of TiO_2 coated ACF filter was enhanced to be $1700 \sim 1860 m^2/g$, depending on the sampling location. The specific surface area of ACF filter having TiO_2 particles which were sampled at 70mm was higher than that of ACF filter with TiO_2 sampled at 110mm. It was found that the pores were micropores, regardless of the sampling location.

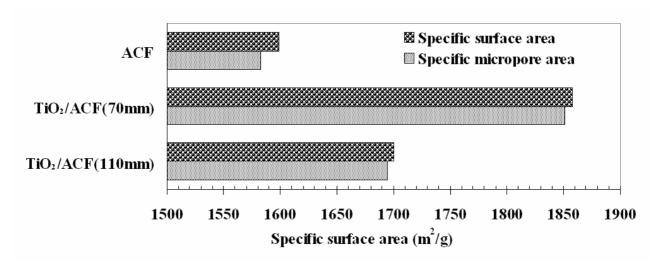


Fig. 5 Specific surface area and micro specific surface area of TiO₂/ACF filters.

4. Conclusion

In this study, ACF were selected as adsorptive support for TiO_2 nanoparticles immobilization. TiO_2 particles were synthesized by using a N_2 -diluted and oxygen-enriched co-flow hydrogen diffusion flame. TiO_2 particles were sampled by using a high temperature particle sampler and then were coated on ACF filters. The sampling was carried out at 70mm and 110mm above the burner. The structure of TiO_2 particles was analyzed by XRD. TiO_2 coated ACF filters were characterized by FESEM-EDX and BET analyses. TiO_2 particles on ACF filters were found to be agglomerated particles and the size of primary particles was approximately 50nm. The structure of TiO_2 particle was enhanced to be $1700\sim1860\text{m}^2/\text{g}$, depending on the sampling location, and the pores were micropores, regardless of the sampling location.

Acknowledgement

The authors acknowledge the supports from Samsung Electronics Co. Ltd. project for "Development of small size nano aerosol generator for HAPs removal" under grant 2005-8-0405.

References

- [1] D. Das, V. Gaur and N. Verma: Carbon Vol. 42 (2004), p. 2949
- [2] Y. Matatov-Meytal and M. Sheintuch: Applied Catalysis A General Vol. 231 (2002), P. 1
- [3] C.H. Ao and S.C. Lee: Applied Catalysis B: Environmental Vol. 44 (2003), p. 191