


 Cite this: *RSC Adv.*, 2018, 8, 14806

Preparation of metallic monolithic Pt/FeCrAl fiber catalyst by suspension spraying for VOCs combustion

 Hao Li,^a Yue Wang,^a Xiao Chen,^b Shuo Liu,^{ac} Ying Zhou,^{*a} Qiulian Zhu,^a Yinfei Chen^a and Hanfeng Lu^{ID}^{*a}

We report a facile and general strategy for the preparation of metallic monolithic catalysts. Our strategy involved subjecting the surfaces of FeCrAl fibers to thermal treatment and the spraying of Pt nanoparticles suspension liquid. The catalyst exhibited high catalytic activity and good stability in the combustion of volatile organic compounds to CO₂ and H₂O at mild temperature. The exceptional activity of the catalyst can be attributed to the well-adhered alumina coating that formed on the surfaces of the FeCrAl fibers after thermal treatment and the highly dispersed Pt nanoparticles on the surface of the alumina coating.

 Received 27th February 2018
Accepted 5th April 2018

DOI: 10.1039/c8ra01720d

rsc.li/rsc-advances

1. Introduction

As one of the major components of air pollution, volatile organic compounds (VOCs) are both precursors of ozone and photochemical smog and substances that are toxic to human health.^{1–4} One of the most effective and economically feasible VOC removal technologies is catalytic combustion because catalytic oxidation leads to degradation of VOCs to CO₂ and H₂O and it is suitable for operation with dilute VOC effluent streams (<1% VOCs).^{5–8} Monolithic catalysts are widely used in catalytic combustion due to their good attrition resistance and very low pressure drop even under high flow rate. Monoliths are commonly composed of ceramics or metallic materials.^{9,10} Metallic materials exhibit higher thermal conductivity and mechanical resistance, better electrical conductivity, and lower manufacturing costs than ceramic materials.^{11,12} Given these advantages, the application of metallic substrates as catalyst supports is becoming increasingly common.^{13–15} Monolithic catalysts are prepared through electrophoretic deposition,¹⁶ anodisation,¹⁷ or steam-only oxidation.¹⁸ Lu *et al.* reported a facile, versatile, and cost-effective one-step route for the fabrication of a composite system of hierarchically structured ALOOH 2D-nanosheets on a paper-like 3D network based on Al-fibers. Their method is aided by the steam-only oxidation of Al. The Al-fiber@ns-ALOOH@Pd catalyst presents high intrinsic activity for the oxidative coupling of CO to DMO.¹⁹

However, several important issues in the fabrication of metallic monolithic catalysts for VOC combustion remain unresolved. These issues include (1) the process complexity and poor adhesion of conventional wash-coating methods^{20–23} and (2) the limitation of active component loading due to the shape of the carrier. The preparation flow of the metallic monolithic catalyst is cumbersome: the active species is loaded through impregnation or through a slurry, which is prepared from a powder catalyst and directly coated on the surface of a metallic substrate; then, the monolithic catalyst is obtained through activation treatment. Existing methods for the fabrication of metallic monolithic catalysts require complex and enormous effort to load the active species on the catalyst supports. Nevertheless, active noble metal nanoparticles (NPs) prepared in liquid form can be randomly loaded through a simple spraying method. Moreover, the deactivated catalyst will not require replacement because it can be reactivated through respraying. This method is very economic and convenient, and it can save raw material costs.

Herein, we synthesized Pt NPs dispersion suspension *via* solvent-thermal method.^{24–26} A well-adhered alumina coating formed on the surface of FeCrAl fibers after thermal treatment at 950 °C in O₂ atmosphere and active NPs were loaded on FeCrAl fibers through simple spraying. Pt NPs were highly dispersed on the surface of the metallic substrate with a monomolecular membrane structure and were firmly bonded on the metallic substrate after calcination (Fig. 1). We evaluated the catalytic performance of the as-synthesized catalyst in the catalytic combustion of toluene, which was used as a model VOC. Our catalyst can exhibit activity toward other VOCs and may enable the significant advancement of VOC removal technologies.^{27,28}

^aInnovation Team of Air Pollution Control, Institute of Catalytic Reaction Engineering, College of Chemical Engineering, Zhejiang University of Technology, Hangzhou 310014, China. E-mail: wjfx@zjut.edu.cn; luhf@zjut.edu.cn; Tel: +86 571 88320767

^bCollege of Environment, Zhejiang University of Technology, China

^cHangzhou Runxing Technology Co., Ltd, Hangzhou, China





Fig. 1 The scheme for preparing monolithic Pt/FeCrAl fiber catalyst by suspension spraying.

2. Experimental section

2.1 Catalyst preparation

Dinitrodiammineplatinum ammoniacal (DPA, 34.6 mg) was mixed with 3 mL oleyl amine (OA) and 2 mL oleic acid (OC) in a three-necked flask equipped with a condenser and a stir bar, followed by heating and holding at 200 °C for 3 h under flowing N_2 atmosphere. The particles thus formed were purified by precipitation in ethanol, centrifuged (6000 rpm, 5 min), washed with ethanol, and re-dispersed in 10 mL *n*-hexane.

The FeCrAl fiber (~ 1.3 mm in thickness) consisting of 7% Al and 85 vol% voids was utilized as the substrate, which was taken from Kunshan Fangdou Net-structured Material Co. Ltd. (China). The FeCrAl fiber was machined to 6 circular chips (15 mm in diameter, about 2.0 g). Then, FeCrAl fiber was immersed in acetone, 10 wt% NaOH, and 10 wt% HNO_3 with ultrasonic treatment for 30 min, in sequence. Subsequently, FeCrAl fiber was calcined in O_2 atmosphere at 950 °C for 3 h. A ready-made Pt nanoparticles dispersion liquid was supported on FeCrAl fiber by spraying; then, the monolithic catalyst was dried at 110 °C for 2 h and calcined at 500 °C for 5 h. The loading of Pt was 0.1 wt% (it refers to theoretical value), the monolithic catalyst is denoted as 0.1Pt/FeCrAl fiber.

2.2 Catalyst characterization

Transmission electron microscopy (TEM) images were taken on a Tecnai G2 F30 S-Twin (Philips-FEI, Netherlands) transmission electron microscope at acceleration voltage of 200 kV. The samples were dispersed in ethanol assisted by an ultrasonic technique. Scanning electron microscopy (SEM) and energy-dispersive X-ray spectroscopy (EDS) analysis were performed on a Hitachi-SU8010 SEM with acceleration voltage of 5 kV for imaging and 15 kV for EDS collection. The X-ray diffraction (XRD) patterns were recorded on an X'pert PRO diffractometer (PAN-alytical, Netherlands) using Cu K_α radiation at a generator voltage of 40 kV and tube current of 40 mA. The samples were scanned in the 2θ range of 10–80° with scanning speed of 0.02° s^{-1} . Diffraction peaks were compared with the standard Joint Committee on Powder Diffraction Standards (JCPDS) database reported by the International Centre for Diffraction Data (ICDD). The adhesion of samples was evaluated using ultrasound tests by immersing the samples in water and applying ultrasound for 1 h. After the samples were dried at 110 °C for 2 h and calcined at 500 °C for 2 h, the weight loss was measured.

2.3 Catalytic activity measurement

The activities of catalysts were tested for VOC catalytic combustion in a fixed-bed stainless steel tubular reactor (internal diameter 16 mm and length 350 mm). In each test run, circular chips of the metallic monolithic catalyst were packed layer-by-layer into the stainless steel tubular reactor, while a K-type thermocouple was placed in the middle of the reactor. For this experiment, we used 2.0 g of catalyst with a total flow rate of the feed stream at 347 $mL\ min^{-1}$, providing a space velocity of 10 000 $mL\ h^{-1}\ g^{-1}$. The feed stream containing 2500 ppm VOCs was generated by bubbling air through a saturator containing pure VOCs chilled in an ice-water isothermal bath, and then diluted with another air stream by the mass flow controller. The concentration of VOCs at the outlet of the reactor was monitored with a gas chromatograph (Kexiao, GC1620) equipped with a flame ionization detector (FID). The concentration of the oxidative product (CO_2) was monitored with a mass spectrograph (MKS Cirrus 2). The removal of VOCs was calculated on the basis of the VOCs consumption. The removal efficiency of VOCs was calculated using the following equation:

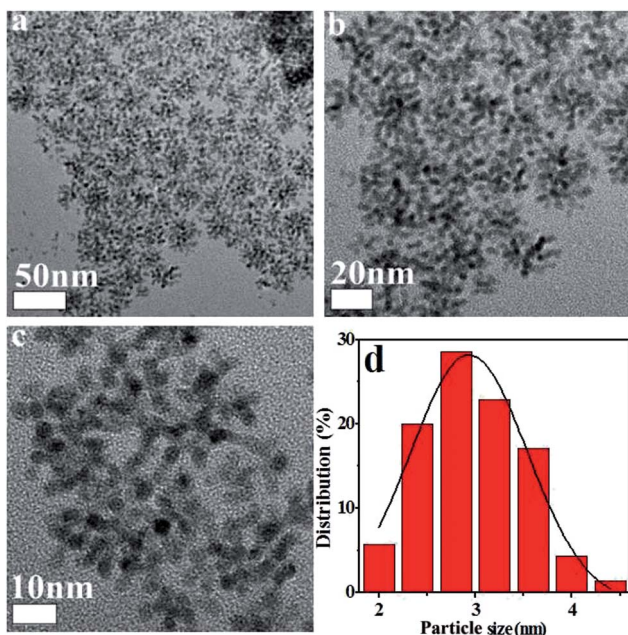


Fig. 2 TEM images of Pt NPs (a–c) and corresponding size histograms (d).

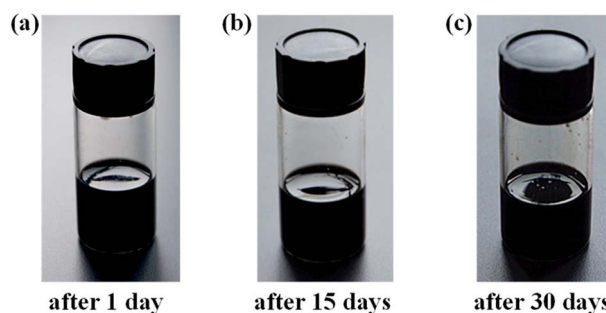


Fig. 3 Photographs of the as-prepared Pt NPs dispersion liquid after 1 day (a), 15 days (b), and 30 days (c).



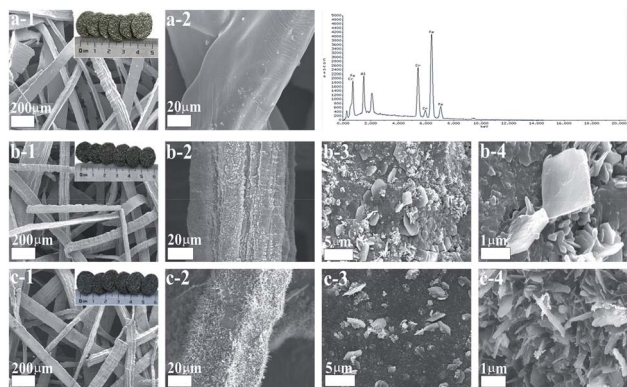


Fig. 4 SEM images (EDS results) of FeCrAl fiber (a), calcined FeCrAl fiber (b), 0.1Pt/FeCrAl fiber (c).

$$\text{VOCs conversion (\%)} = \frac{[\text{VOCs}]_{\text{in}} - [\text{VOCs}]_{\text{out}}}{[\text{VOCs}]_{\text{in}}} \times 100\%$$

3. Results and discussion

3.1 TEM characterization

Fig. 2 shows TEM images of the Pt NPs, which appear spherical in shape with low aggregation and narrow size distribution.^{29,30} The average sizes of Pt NPs were in the range of 2–4 nm. The monodispersed structure of Pt NPs promoted their dispersal in liquid. Simple spraying was used to randomly load the active species in the VOCs purification system. In addition, the Pt NPs dispersion liquid was extremely stable. It could be stored for 10–30 days without delamination and sol destruction. The storage behavior of the Pt NPs dispersion liquid indicated that it could be continuously used for long periods without changing (Fig. 3). Furthermore, this method is generally applicable and could be easily applied with the dip-coating method. Notably, the low Pt content required for catalyst loading decreased the cost.

3.2 SEM characterization

To promote anchoring of the coating to the metallic substrate and to increase the exposed area of the catalyst, FeCrAl fiber was thermally treated at 950 °C for 3 h in O₂ atmosphere. To confirm that the applied thermal treatments enabled the formation of a uniform rough surface on the FeCrAl fiber, the morphology and composition of the fibers were studied. Fig. 4 and Table 1 show SEM micrographs and EDS analyses of the surfaces of the untreated and treated FeCrAl fibers. SEM micrographs of the untreated FeCrAl fiber are shown in Fig. 4a. The untreated FeCrAl fiber exhibited slightly cracked smooth surfaces. Additionally, the mass percentages of the elements (Fe, Cr and Al) reported by EDS analysis were similar to those reported by the manufacturers. Fig. 4b shows SEM micrographs of the treated (calcined at 950 °C for 3 h in O₂ atmosphere) FeCrAl fiber, which showed the growth of uniformly roughened surfaces.²² Given that the bulk aluminum had a high chemical potential and was likely to be oxidized by oxygen, it easily migrated from the bulk to the surface and reacted with oxygen to form alumina

Table 1 Atomic composition of the sample determined by X-EDS

Sample	Element	Weight/%	Atomic/%
FeCrAl fiber	Fe	71.27	65.74
	Cr	22.41	22.20
	Al	6.32	12.06
Calcined FeCrAl fiber	Fe	30.75	15.93
	Cr	11.53	6.42
	Al	36.30	38.92
	O	21.42	38.74
0.1Pt/FeCrAl fiber	Fe	35.79	23.36
	Cr	4.67	3.44
	Al	29.15	24.36
	O	16.71	41.35
	Pt	13.68	7.49

whiskers.^{31–33} In addition, the EDS analysis results of the thermally treated fibers indicated significant changes to the mass percentages of each element. Clearly, the percentages of Al and O increased, while that of Fe decreased because of the formation of the alumina layer by thermal treatment.^{34,35} The well-adhered alumina coating promoted firm bonding of the active species to the metallic substrate. SEM micrographs of the monolithic catalyst are presented in Fig. 4c. The images show that the Pt catalyst formed a homogeneous surface on the whiskers or pores. The addition of Pt to the FeCrAl fiber was supported by the percentages reported by EDS analysis, which detected Pt content of approximately 13%. This finding indicated that thermal treatment increased the exposed area of the FeCrAl fiber. This effect ensured catalytic activity by promoting the even dispersion of NPs. Moreover, ultrasound tests were conducted to confirm adherence between the wash-coated and active phases. The total weight losses of the samples were only 0.9 and 0.6 wt%. These results indicated excellent adhesion between the wash-coated and active phases.

3.3 XRD characterization

The crystal phases of samples were identified by XRD. Fig. 5 shows the diffraction patterns of FeCrAl fiber before and after the treatments. XRD patterns of untreated FeCrAl fiber confirmed the presence of Fe–Cr alloys of ferrite type (PDF 54-0331). However, no diffraction peaks of Al were detected because the content of Al is very low. After calcination treatment at 950 °C, the formation of the crystalline phase Fe₂AlCr (PDF 54-0387) was found. This indicated that the percentage of Al increased after calcination; as confirmed from EDS analysis. Moreover, no diffraction peaks of alumina were detected, indicating high dispersion of the alumina layer. It should be noted that no Pt XRD peaks were observed for the 0.1Pt/FeCrAl fiber catalyst, indicating the feature of highly dispersed Pt. Indeed, the TEM image clearly shows that Pt is highly dispersed with average particle size of only 2–4 nm.

3.4 Activity test

The catalytic activities of the samples in toluene combustion were tested under the base condition of toluene inlet



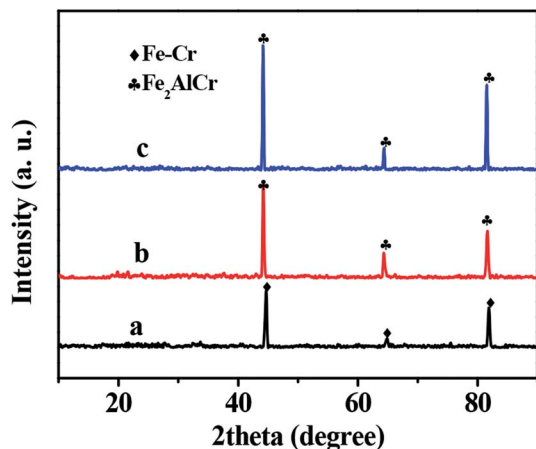


Fig. 5 XRD patterns of FeCrAl fiber (a), calcined FeCrAl fiber (b), 0.1Pt/FeCrAl fiber (c).

concentration of 2500 ppm and WHSV of $10\,000\text{ mL h}^{-1}\text{ g}^{-1}$; the results are presented in Fig. 6a. In the blank experiment (without Pt), no distinct toluene conversion was detected below $300\text{ }^{\circ}\text{C}$ and gas chromatography (GC) detected the presence of other organic compounds in the gaseous product. These results indicated that the direct thermal oxidation of toluene under test conditions and in the absence of active species is negligible.³⁶ In contrast, the 0.1Pt/FeCrAl fiber catalyst achieved complete toluene conversion ($>99\%$) at temperatures exceeding $280\text{ }^{\circ}\text{C}$. This result, in addition to SEM and TEM observations, indicated that catalytic activity improved with the formation of a well-adhered alumina coating on the surfaces of the FeCrAl fiber through thermal treatment and that 2–4 nm Pt NPs were highly dispersed on the surfaces of the alumina coating. In addition, mass spectroscopy (MS) results showed that the catalyst had nearly 100% selectivity for CO_2 at a temperature of $280\text{ }^{\circ}\text{C}$ (Fig. 6b). The ignition curves could be represented either by following the decrease in the toluene signal or by following the increase in CO_2 signal.¹¹ Furthermore, on-line MS did not detect CO during the reaction. This result indicated that our catalyst has high selectivity for CO_2 .

3.5 Stability test

To examine stability of the catalyst, we performed on-stream toluene combustion at $280\text{ }^{\circ}\text{C}$ and WHSV of $10\,000\text{ mL h}^{-1}\text{ g}^{-1}$. The results are shown in Fig. 7. Clearly, the conversion rate of toluene was maintained at 100% and catalytic activity did not decrease during 600 min of on-stream reaction. This result suggested that the catalyst has excellent stability and can be applied in the industrial combustion of VOCs.

3.6 Applicability test

VOCs include alkanes, alcohols, ketones, esters and nitrogen-containing compounds.^{7,37} To confirm that the catalyst can be applied in the catalytic combustion of other VOCs, we tested the activities of the catalyst in the combustion of ethyl acetate, acetone, *n*-hexane, isopropyl alcohol (IPA) and *N,N*-

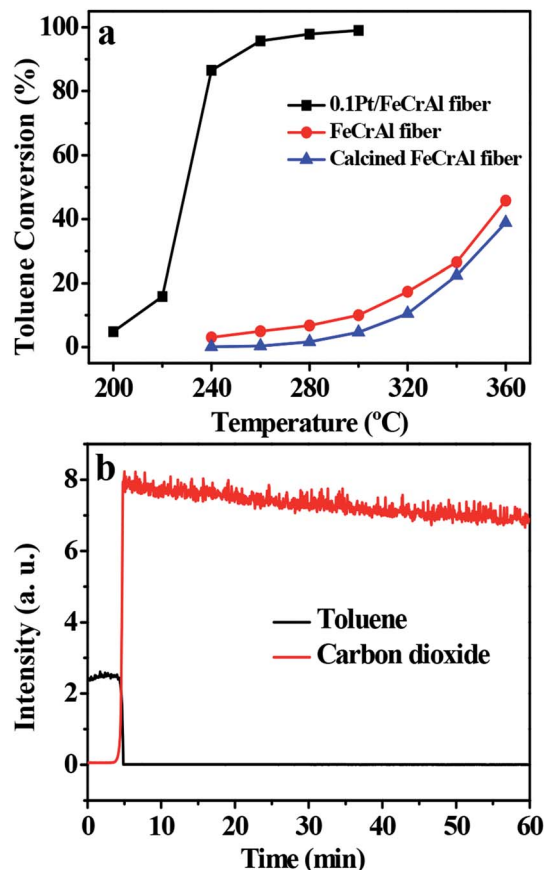


Fig. 6 Light-off curves of toluene catalytic combustion (a). MS curves of toluene catalytic combustion (temperature = $280\text{ }^{\circ}\text{C}$) (b). Other conditions: toluene concentration 2500 ppm and WHSV $10\,000\text{ mL h}^{-1}\text{ g}^{-1}$.

dimethylformamide (DMF). The catalytic performance of 0.1Pt/FeCrAl fiber in VOC combustion is shown in Fig. 8. The temperatures for VOC conversion over 0.1Pt/FeCrAl fiber catalyst for 10% (T_{10}), 50% (T_{50}) and 90% (T_{90}) of total conversion are summarized in Table 2. We conclude that the catalytic

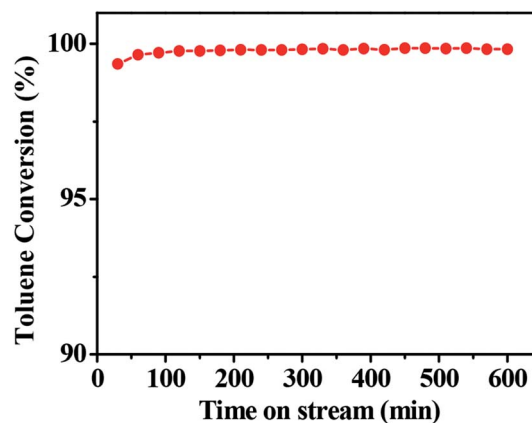


Fig. 7 The stability of toluene combustion over 0.1Pt/FeCrAl fiber ($T = 280\text{ }^{\circ}\text{C}$). Other conditions: toluene concentration 2500 ppm and WHSV $10\,000\text{ mL h}^{-1}\text{ g}^{-1}$.



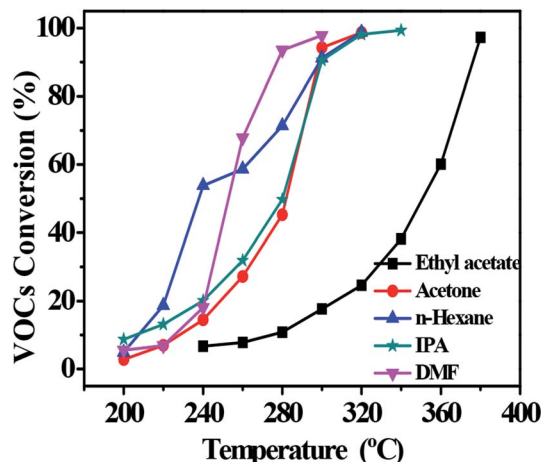


Fig. 8 Light-off curves of VOC catalytic combustion over 0.1Pt/FeCrAl fiber. Other conditions: VOC concentration of 2500 ppm and WHSV of 10 000 mL h⁻¹ g⁻¹.

Table 2 Comparison of catalytic activity of 0.1Pt/FeCrAl fiber for VOC combustion

VOCs	$T_{10}/^{\circ}\text{C}$	$T_{50}/^{\circ}\text{C}$	$T_{90}/^{\circ}\text{C}$
Toluene	213	235	242
DMF	228	250	276
n-Hexane	214	238	295
IPA	212	280	298
Acetone	230	282	300
Ethyl acetate	280	350	374

activities of our catalyst for VOC combustion are hydrocarbon (nitrogen) > oxygenated compounds because hydrocarbon VOCs were strongly absorbed on the Pt catalyst and were quickly activated. These behaviors are similar to those reported in the literature.^{7,38} In addition, it can be illustrated that our catalyst can be applied to the catalytic combustion of different types of VOCs.

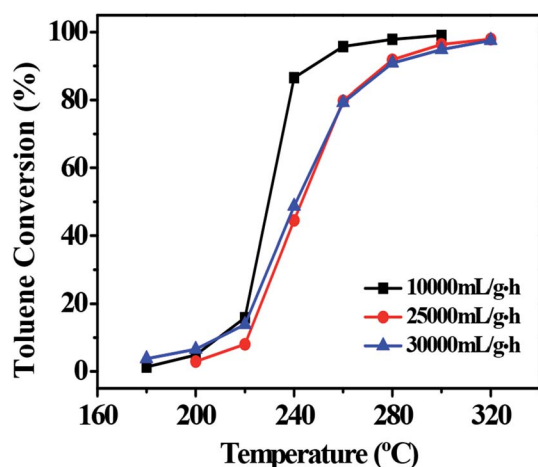


Fig. 9 Effect of space velocity on toluene combustion over 0.1Pt/FeCrAl fiber. Other conditions: toluene concentration of 2500 ppm.

3.7 Effect of space velocity

Space velocity is an important parameter for the practical application of catalysts.³⁹ To achieve high VOC combustion efficiency a high space velocity is required for catalysts.³⁶ Hence, the catalytic behavior of the 0.1Pt/FeCrAl fiber in the catalytic combustion of toluene under three different WHSV values was tested (Fig. 9).⁴⁰ Increasing the WHSV values decreased the toluene conversion at the same reaction temperature. This result may be attributed to the shortened residence time of toluene in the catalyst bed at high WHSV. Nevertheless, the result suggested that the catalyst could be used to catalyze the conversion of high VOC fluxes. Thus, our catalyst has considerable application potential in the environmental catalysis field.

4. Conclusions

Highly dispersed Pt NPs dispersion liquid was prepared *via* the solvent-thermal method. A well-adhered alumina layer was formed on the surfaces of FeCrAl fibers through thermal treatment. Then, the supported catalyst was fabricated by simply spraying the dispersion liquid on the FeCrAl fibers. Active species were randomly loaded on the surfaces of FeCrAl fibers by spraying. In addition, the catalyst with Pt loading of 0.1 wt% delivered 100% toluene conversion and high CO₂ selectivity at the space velocity of 10 000 mL h⁻¹ g⁻¹. Moreover, the catalyst was stable for at least 600 min without deactivation. The intrinsic activity of the catalyst is intimately related to the high dispersion of 2–4 nm Pt NPs on the surfaces of the alumina coating. Furthermore, we believe that this strategy will provide a new route for the preparation and application of environmental metallic monolithic catalysts.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This study was supported by the Natural Science Foundation of China (No. 21506194, No. 21676255), the Natural Science Foundation of Zhejiang Province (No. Y16B070011), and the Commission of Science and Technology of Zhejiang province (No. 2017C03007, No. 2017C33106).

Notes and references

- 1 H. Wang, W. Yang, P. Tian, J. Zhou, R. Tang and S. Wu, *Appl. Catal., A*, 2017, **529**, 60–67.
- 2 H. Lu, Y. Zhou, W. Han, H. Huang and Y. Chen, *Catal. Sci. Technol.*, 2013, **3**, 1480–1484.
- 3 S. Scire, S. Minico, C. Crisafulli, C. Satriano and A. Pistone, *Appl. Catal., B*, 2003, **40**, 43–49.
- 4 H. Huang, Y. Xu, Q. Feng and D. Y. Leung, *Catal. Sci. Technol.*, 2015, **5**, 2649–2669.
- 5 B. Li, J. Wang and H. Gong, *Catal. Today*, 2009, **148**, 81–87.



- 6 S. Xie, Y. Liu, J. Deng, J. Yang, X. Zhao, Z. Han and H. Dai, *Catal. Sci. Technol.*, 2018, **8**, 806–816.
- 7 M. S. Kamal, S. A. Razzak and M. M. Hossain, *Atmos. Environ.*, 2016, **140**, 117–134.
- 8 Y. Wang, L. Guo, M. Chen and C. Shi, *Catal. Sci. Technol.*, 2018, **8**, 459–471.
- 9 W. Zhang and D. Wu, *Ceram. Int.*, 2016, **42**, 16563–16570.
- 10 K. S. Yang, J. S. Choi and J. S. Chung, *Catal. Today*, 2004, **97**, 159–165.
- 11 N. Burgos, M. Paulis, M. M. Antxustegi and M. Montes, *Appl. Catal., B*, 2002, **38**, 251–258.
- 12 W. Fei, S. C. Kuiry and S. Seal, *Oxid. Met.*, 2004, **62**, 29–44.
- 13 L. M. Martinez, O. Sanz, M. I. Dominguez, M. A. Centeno and J. A. Odriozola, *Chem. Eng. J.*, 2009, **148**, 191–200.
- 14 M. Valentini, G. Groppi, C. Cristiani, M. Levi, E. Troncoin and P. Forzatti, *Catal. Today*, 2001, **69**, 307–314.
- 15 S. Cimino, A. Gambriasi, L. Lisi, G. Mancino, M. Musiani, L. Vázquez-Gómez and E. Verlato, *Chem. Eng. J.*, 2016, **285**, 276–285.
- 16 H. Sun, X. Quan, S. Chen, H. M. Zhao and Y. Zhao, *Appl. Surf. Sci.*, 2007, **253**, 3303–3310.
- 17 O. Sanz, F. J. Echave, M. Sánchez, A. Monzón and M. Montes, *Appl. Catal., A*, 2008, **340**, 125–132.
- 18 C. Wang, L. Han, P. Chen, G. Zhao, Y. Liu and Y. Lu, *J. Catal.*, 2016, **337**, 145–156.
- 19 C. Wang, L. Han, Q. Zhang, Y. Li, G. Zhao, Y. Liu and Y. Lu, *Green Chem.*, 2015, **17**, 3762–3765.
- 20 Y. Diaz, A. Sevilla, A. Mónaco, F. J. Méndez, P. Rosales, L. Garcia and J. L. Brito, *Fuel*, 2013, **110**, 235–248.
- 21 J. E. Samad, J. A. Nychka and N. V. Semagina, *Chem. Eng. J.*, 2011, **168**, 470–476.
- 22 B. P. Barbero, L. Costa-Almeida, O. Sanz, M. Morales, L. E. Cadus and M. Montes, *Chem. Eng. J.*, 2008, **139**, 430–435.
- 23 I. Yuranov, N. Dunand, L. Kiwi-Minsker and A. Renken, *Appl. Catal., B*, 2002, **36**, 183–191.
- 24 L. Wu, A. Mendoza-Garcia, Q. Li and S. Sun, *Chem. Rev.*, 2016, **116**, 10473–10512.
- 25 P. Hou, H. Liu, J. Li and J. Yang, *CrystEngComm*, 2015, **17**, 1826–1832.
- 26 C. Shen, C. Hui, T. Yang, C. Xiao, J. Tian, L. Bao, S. Chen, H. Ding and H. Gao, *Chem. Mater.*, 2008, **20**, 6939–6944.
- 27 Y. Wang, Y. Xue, C. Zhao, D. Zhao, F. Liu, K. Wang and D. D. Dionysiou, *Chem. Eng. J.*, 2016, **300**, 300–305.
- 28 W. Zhang, L. Hu, F. Wu and J. Li, *Catal. Lett.*, 2017, **147**, 407–415.
- 29 L. Pérez-Mirabet, E. Solano, F. Martínez-Julián, R. Guzmán, J. Arbiol, T. Puig, X. Obradors, A. Pomar, R. Yáñez, J. Ros and S. Ricart, *Mater. Res. Bull.*, 2013, **48**, 966–972.
- 30 W. Wang, L. Zhuang, Y. Zhang and H. Shen, *Mater. Res. Bull.*, 2015, **69**, 61–64.
- 31 P. Avila, M. Montes and E. E. Miró, *Chem. Eng. J.*, 2005, **109**, 11–36.
- 32 S. Zhao, J. Zhang, D. Weng and X. Wu, *Surf. Coat. Technol.*, 2003, **167**, 97–105.
- 33 S. Govender and H. B. Friedrich, *Catalysts*, 2017, **7**, 62.
- 34 J. M. Zamaro, M. A. Ulla and E. E. Miró, *Appl. Catal., A*, 2006, **308**, 161–171.
- 35 J. Jia, J. Zhou, J. Zhang, Z. Yuan and S. Wang, *Appl. Surf. Sci.*, 2007, **253**, 9099–9104.
- 36 L. Lin and H. Bai, *Chem. Eng. J.*, 2016, **291**, 94–105.
- 37 Z. Zhang, Z. Jiang and W. Shangguan, *Catal. Today*, 2016, **264**, 270–278.
- 38 L. F. Liotta, *Appl. Catal., B*, 2010, **100**, 403–412.
- 39 H. Li, C.-Y. Wu, Y. Li and J. Zhang, *Appl. Catal., B*, 2012, **111**, 381–388.
- 40 C. W. Ahn, Y. W. You, I. Heo, J. S. Hong, J. K. Jeon, Y. D. Ko, Y. H. Kime, P. Hosik and J. K. Suh, *J. Ind. Eng. Chem.*, 2017, **47**, 439–445.

