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**The formation and migration of sulfur trioxide over  $V_2O_5/TiO_2$  catalyst in SCR of  $NO_x$  with  $NH_3$**

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The oxidation of sulfur dioxide ( $SO_2$ ) to sulfur trioxide ( $SO_3$ ) is undesirable during the selective catalytic reduction (SCR) of nitrogen oxides ( $NO_x$ ) found in the flue gas of power plants. Sulfur trioxide has toxicity over 10 times than that of  $SO_2$  and is highly corrosive, resulting in corrosion of equipment and pollution of the environment. The objectives of this research are to establish the fundamental kinetics and mechanism of  $SO_2$  oxidation over supported  $V_2O_5/TiO_2$  catalyst and establish sulfur balance, then, use these insights to propose a basic theory for collaboratively controlling  $SO_3$  emissions. A series of supported  $V_2O_5/TiO_2$  catalysts were prepared with various  $V_2O_5$  contents: 0.5%, 1%, 1.5%, 3%, 4% and 5%. Raman spectroscopy is used to determine the coordination of surface species. Isolated vanadia species were dispersed on the  $TiO_2$  surface as Ti-O-V bonds at VOx coverage below the monolayer. Sulfur dioxide temperature-programmed desorption and *in situ* diffuse reflectance infrared Fourier transform spectroscopy were conducted to study the interaction between  $SO_2$  and  $V_2O_5/TiO_2$ . It was found that Ti-O-V was demonstrated to play a critical role in the  $SO_2$  adsorption and oxidation. A possible reaction mechanism of  $SO_2$  oxidation over  $V_2O_5/TiO_2$  catalyst was established in this study. The turnover frequency for  $SO_2$  oxidation was very low, approximately  $10^{-5} s^{-1}$  at 320 °C. Over the range of conditions studied, the rate of  $SO_2$  oxidation is zero-order in oxygen and first-order in  $SO_2$  and the apparent activation energies for  $SO_2$  oxidation was obtained. The decrease of  $SO_2$ , the gaseous  $SO_3$  and the sulfate deposited on catalyst surface were to be quantified by Fourier transform infrared spectrometer (FTIR), PENTOL  $SO_3$  analyzer and ion chromatography (IC), respectively. Then, the sulfur balance was established.

**Biography**

Jin Xiong is currently pursuing his PhD in Environmental Engineering from University of Chinese Academy of Sciences, Beijing, China. He has completed his Bachelor's degree in Chemical Engineering and Technology from the Central South University, Changsha, China. His research interest is on  $SO_3$  formation mechanism and migration route during SCR DeNOx process and  $SO_3$  emissions collaboratively controlling.

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