

# Effect of pore structure of TiO<sub>2</sub> on the SO<sub>2</sub> poisoning over V<sub>2</sub>O<sub>5</sub>/TiO<sub>2</sub> catalysts for selective catalytic reduction of NO<sub>x</sub>

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Selective catalytic reduction (SCR) of NO<sub>x</sub> is a principal technology for the removal of NO<sub>x</sub> from stationary and mobile sources. In practical SCR system, vanadium catalyst is widely used for DeNO<sub>x</sub> treatment [1]. SCR catalysts are generally deactivated in the presence of SO<sub>2</sub>. However, a question about how the activity of the catalyst is affected in the presence of SO<sub>2</sub> remains unclear.

We prepared vanadium catalysts supported on TiO<sub>2</sub> containing 5wt% V<sub>2</sub>O<sub>5</sub>, denoted as 5wt% VT. Mesoporous TiO<sub>2</sub> (DT-51) and microporous TiO<sub>2</sub> [2] were used as support with anatase phase. In this work, two types of vanadium catalysts supported on TiO<sub>2</sub> with different pore structure were investigated when they were exposed to SO<sub>2</sub>. Thus we aimed at examining the effect of pore structure of TiO<sub>2</sub> support on SO<sub>2</sub> poisoning when NH<sub>3</sub> SCR was carried out.

NO<sub>x</sub> conversion of 5wt% VT(DT-51) decreased by 15% whereas only under 5% decline was observed on 5wt% VT(micro) at 350°C when SO<sub>2</sub> existed in the feed. Also, Fig.1 showed the amount of SO<sub>2</sub> desorption from pre-adsorbed sulfur species. Although all catalysts were treated with 300ppm SO<sub>2</sub> for 3h, 5wt% VT(DT-51) generated 2.5 times higher SO<sub>2</sub> than 5wt% VT(micro). In other words, different affinity with SO<sub>2</sub> is observed depending on the TiO<sub>2</sub> type.

Ammonium bisulfate (NH<sub>4</sub>HSO<sub>4</sub>: ABS) is known to form on the SCR catalysts via the reaction between NH<sub>3</sub> and H<sub>2</sub>SO<sub>4</sub>, generated from H<sub>2</sub>O and SO<sub>2</sub>. To distinguish the sulfate species formed on catalysts, model sulfate

compounds, NH<sub>4</sub>HSO<sub>4</sub> as diluted solution, was intentionally impregnated on VT catalysts [3]. Such approach allowed us to analyze the sulfur species of each catalyst. 0.1M ABS impregnated 5wt% VT samples had different weight loss patterns at similar temperature range during TG analysis. According to Fig. 2, sulfate species on the catalysts decomposed with different weight loss (%) at 420 - 440°C and 546 - 547°C. At low temperature range around 430°C, larger amount of sulfate species decomposed on 5wt% VT(micro) than 5wt% VT(DT-51).

Therefore, it is summarized that V<sub>2</sub>O<sub>5</sub>/TiO<sub>2</sub> (micro) has lower affinity with SO<sub>2</sub> and is also able to decompose sulfate species more readily than V<sub>2</sub>O<sub>5</sub>/TiO<sub>2</sub> (DT-51), which can explain the less decline of SCR activity in the presence of SO<sub>2</sub> over the former catalyst.

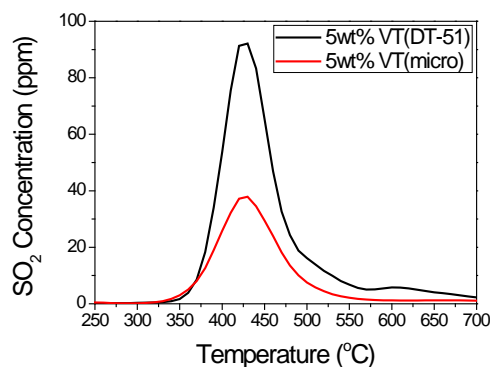


Fig. 1. Desorption of SO<sub>2</sub> of pre-sulfated 5wt% V<sub>2</sub>O<sub>5</sub>/TiO<sub>2</sub> catalysts.

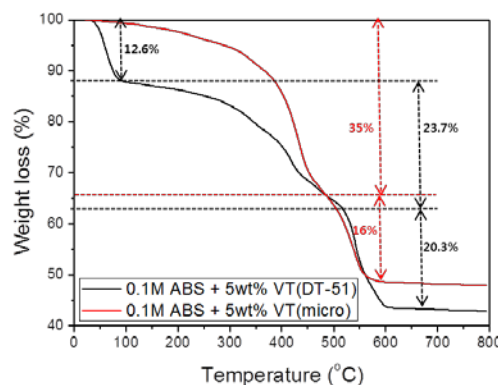


Fig.2. TG-DTA of ammonium bisulfate impregnated V<sub>2</sub>O<sub>5</sub>/TiO<sub>2</sub> catalysts.

## REFERENCES

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