

SO₂ Poisoning Mechanism

Subjects: Others

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The selective catalytic reduction (SCR) has been widely used in industrial denitrification owing to its high denitrification efficiency, low operating costs, and simple operating procedures. However, coal containing a large amount of sulfur will produce SO₂ during combustion, which makes the catalyst easy to be deactivated, thus limiting the application of this technology. This review summarizes the latest NH₃-SCR reaction mechanisms and the deactivation mechanism of catalyst in SO₂-containing flue gas. Some strategies are summarized for enhancing the poison-resistance through modification, improvement of support, the preparation of complex oxide catalyst, optimizing the preparation methods, and acidification. The mechanism of improving sulfur resistance of catalysts at low temperatures is summarized, and the further development of the catalyst is also prospected. This paper could provide a reference and guidance for the development of SO₂ resistance of the catalyst at low temperatures.

Keywords: selective catalytic reduction ; SO₂ resistance at low temperature ; catalyst modification ; NO_x ; NH₃-SCR

1. Introduction

Nitrogen oxide (NO_x) is a general term composed of nitrogen, oxygen, and other compounds. It is one of the major pollutants from the exhaust gas of thermal power plants, industrial furnaces, motor vehicles, ship exhaust emissions, and includes N₂O, NO, NO₂, etc.—among which NO and NO₂ account for the largest proportion^[1]. A large amount of NO_x emitted into the air will cause a series of environmental concerns. Therefore, exploring and developing efficient exhaust gas deNO_x technology has been an area of intense investigation. Among all flue gas denitrification technologies, selective catalytic reduction (SCR) is an extensively applied technology due to its low reaction temperature and high denitrification efficiency^{[2][3]}. Selective catalytic reduction (SCR) mainly refers to the reaction of NO_x using NH₃ as a reducing agent in the presence of O₂ to produce pollution-free N₂ and H₂O, whose core is the catalyst.

2. SO₂ Poisoning Mechanism of Low-Temperature Catalyst

At present, some power plants adopt wet desulfurization to remove SO₂ with lower flue gas temperature, failing to meet the reaction requirements of V₂O₅/TiO₂ catalyst. Therefore, there are many drawbacks such as low denitrification efficiency and catalyst waste. After desulfurization, tiny amounts of SO₂ still exist in the exhaust gas, bringing about the deactivation of SCR catalyst. Therefore, developing a vanadium-free catalyst with great denitrification performance and sulfur and water resistance at low temperatures is extremely necessary^[4]. To solve the sulfur poisoning of catalysts, many scholars have done a lot of research and elaborated on the poisoning mechanism in detail.

From the above studies, the SO₂ deactivation mechanism on the catalyst at low temperatures can be observed mainly in the following three aspects. (1) The ammonium sulfate and ammonium bisulfate are formed by the reaction of SO₂ and NH₃ in the presence of O₂ and attach to the catalyst surface, which can decrease the surface area, pore volume, and pore size of the catalyst, and then reduce the reaction rate. However, ammonium sulfate and ammonium bicarbonate will self-decompose when the NH₃-SCR reaction is carried out above 280 °C and 350 °C, respectively, so the catalytic activity is able to be restored by the washing method at low temperatures^[5]. (2) In the presence of O₂, SO₂ will react with the active component (mainly transition metal) on the catalyst surface to generate metal sulfate salt, which will cause irreversible deactivation of the catalyst. (3) SO₂ will compete with NO at the adsorption sites on the catalyst surface when these acidic gases are present in the reaction system, which would reduce the formation of SCR intermediate products and the catalytic efficiency of catalyst. [Figure 1-3](#) show the mechanism of catalyst sulfur poisoning.

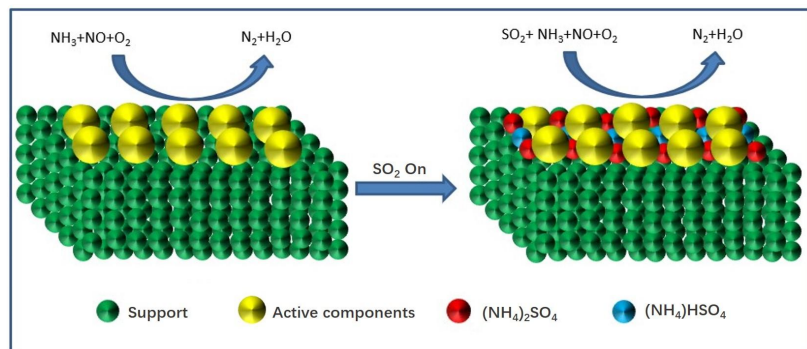


Figure 1. The formation process of $(\text{NH}_4)_2\text{SO}_4$ and NH_4HSO_4 .

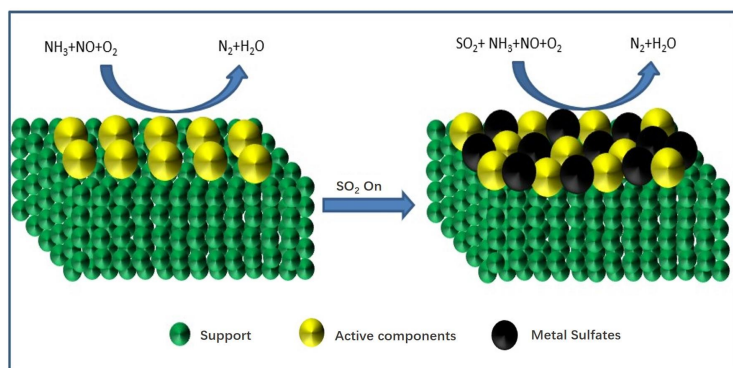


Figure 2. Sulfation of active components.

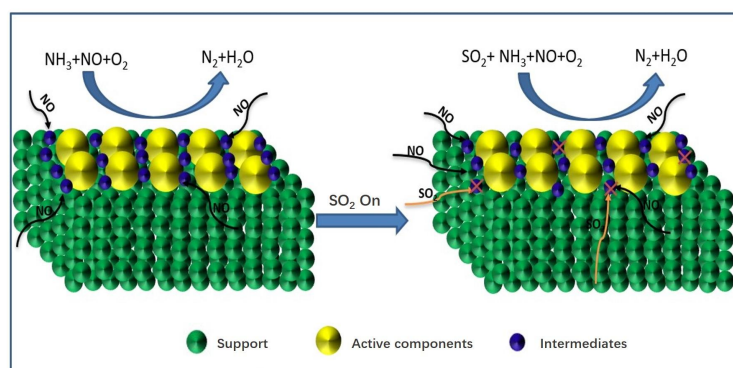


Figure 3. Competitive adsorption of NO and SO_2 .

3. Research Progress of SO_2 Resistance Catalyst at Low Temperatures

Catalyst is usually composed of active component and support, and there are other forms of catalysts such as composite oxide catalysts. To enhance the low-temperature sulfur resistance of catalysts, many scholars have focused their attention on the improvement of active components and supports. In addition, some scholars have found that the preparation method—the handling catalyst by acidification and reaction conditions—can make a difference in the SO_2 resistance of catalyst.

3.1. Effects of Active Components

The active component, which is composed of one or more substances, is the main unit of catalyst and affects the NH_3 -SCR reaction significantly. Using rare earth metals as well as transition metal oxides to improve active components is one of the most effective methods to improve sulfur resistance at low temperatures. We mainly summarized the SCR performance and/or SO_2 resistance mechanism of catalysts modified with Ce, Fe, Cu, W and other metal elements, as shown in [Figure 4-9](#).

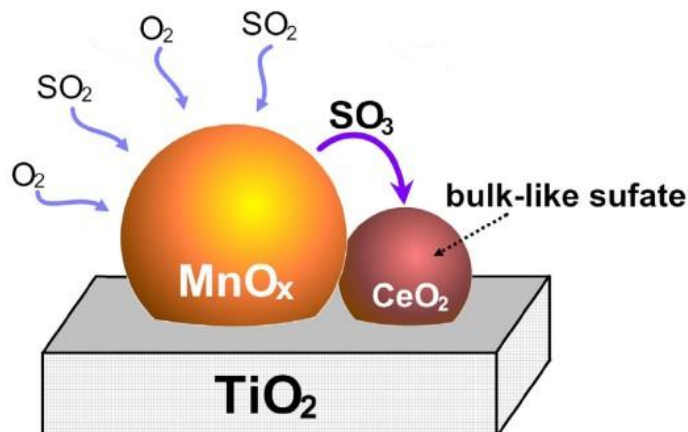


Figure 4. The sulfation mechanism of Ce-modified Mn–Ce/ TiO_2 catalyst^[6].

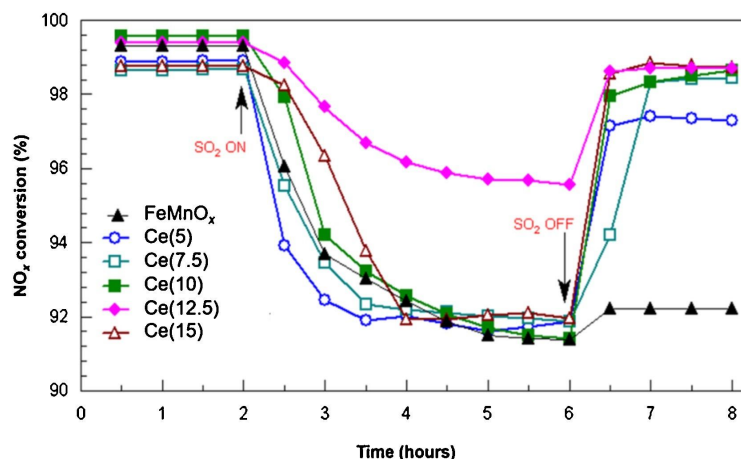


Figure 5. Influence of SO_2 on NO_x conversion of FeMnO_x and $\text{Ce}(y)$ catalysts. Reaction conditions: $[\text{NO}] = [\text{NH}_3] = 0.1\%$; $[\text{SO}_2] = 100 \text{ ppm}$; $[\text{O}_2] = 3\%$; N_2 balance, $\text{GHSV} = 30,000 \text{ h}^{-1}$; reaction temperature = 120°C ^[7].

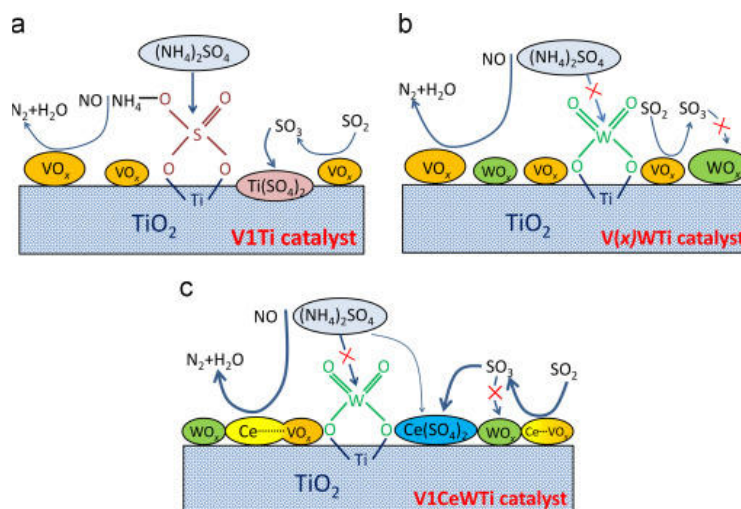


Figure 6. The mechanism of sulfur poisoning at low temperatures of catalysts (a) V1Ti; (b) V(x)WTi; (c) V1CeWTi^[8].

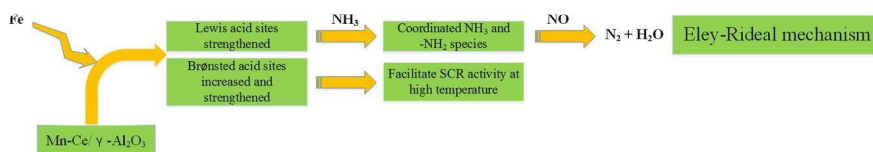


Figure 7. The reaction flow chart of Fe–Mn–Ce/ $\gamma\text{-Al}_2\text{O}_3$ catalyst^[9].

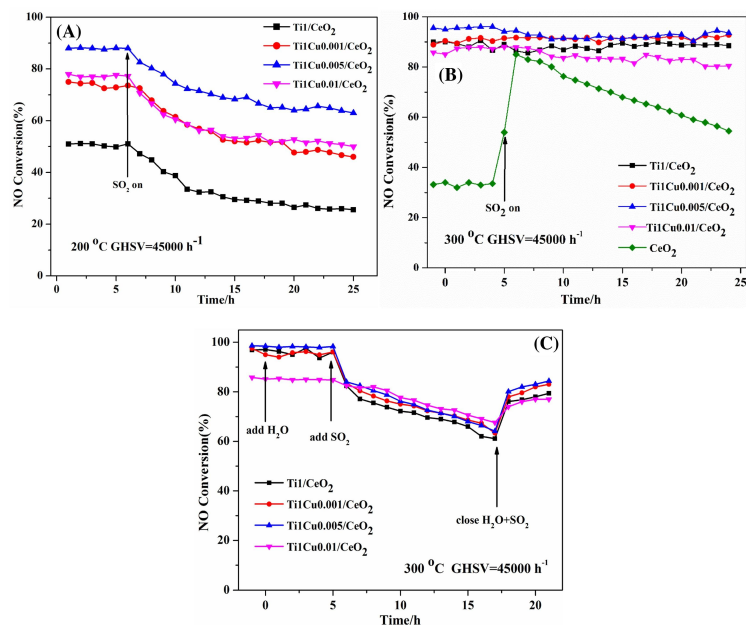


Figure 8. SO₂ durability of catalysts (A) SO₂ durability over Ti1/CeO₂ and Ti1Cuy/CeO₂ catalysts at 200 °C (B) SO₂ durability over Ti1/CeO₂ and Ti1Cuy/CeO₂ catalysts at 300 °C. (C) H₂O/H₂O + SO₂ durability over Ti1/CeO₂ and Ti1Cuy/CeO₂ at 300 °C^[10].

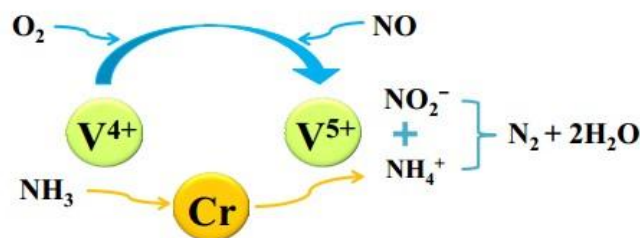


Figure 9. Mechanism of Cr doping improving sulfur resistance of Cr-V/TiO₂ catalyst^[11].

Above all, the content of oxygen adsorbed on the active material and catalyst surface—and even the number of acid sites—could be increased by adding Ce, Fe, Cu, W, and others to the active components, which would accelerate the rapid reaction of SCR and increase the catalytic performance of catalyst. Under the reaction conditions containing SO₂, these additives can preferentially react with SO₂ as the SO₂ catchers to avoid the sulfation of active substances. Besides, the stability of NH₄HSO₄ and (NH₄)₂SO₄ on the surface was reduced, and the low-temperature sulfur tolerance of catalyst was effectively improved. It is believed that transition metals and rare earth elements will have promising applications in improving sulfur resistance of catalyst at low temperatures. Figure 10 shows the typical SO₂-tolerant modified catalyst at low temperatures for selective catalytic reduction (SCR) reaction.

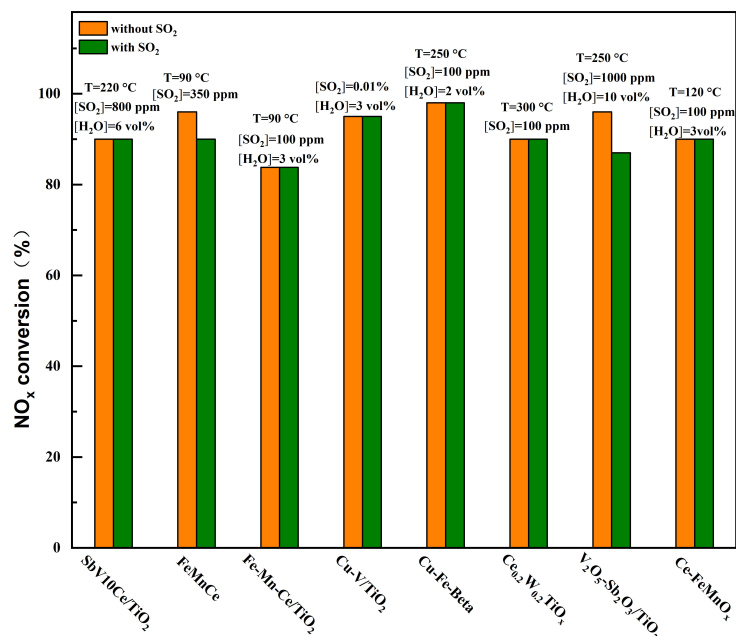


Figure 10. Typical SO₂-tolerant modified catalyst at low temperatures for selective catalytic reduction (SCR) reaction^{[7][12][9][13][14][15][16][17]}.

3.2. Effects of Supports

The support of catalyst is of vital importance in the catalytic activity. Loading the active components onto the support contributes to improving the specific surface area, thermal resistance, and mechanical strength of catalyst. Especially, the supports can slow down SO₂ poisoning on the catalytic activity of catalyst. As of now, there are TiO₂, Al₂O₃, activated carbon and zeolites, and other conventional supports in practical application, as shown in [Figure 11](#) and [Figure 12](#). However, different supports show different catalytic activity and sulfur resistance, which makes the research and improvement of the support become a part of the emphasis of research to improve the low-temperature SO₂ resistance of catalyst. [Figure 13](#) shows the typical SO₂-tolerant catalysts with different supports at low temperatures for SCR reaction.

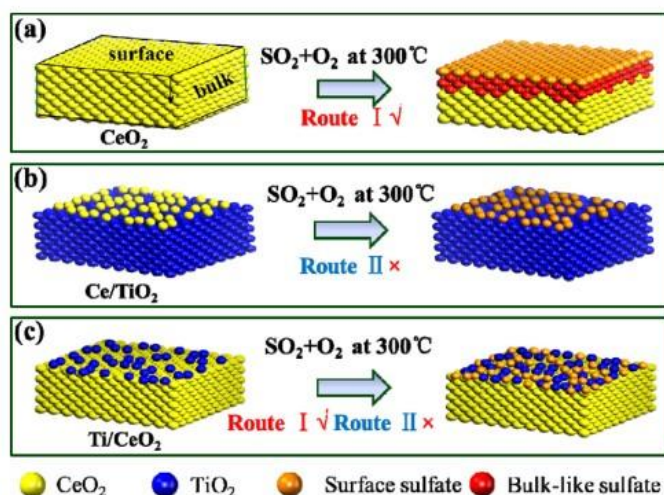


Figure 11. Sulfur resistance mechanism of (a) CeO₂, (b) Ce/Ti and (c) Ti/Ce catalyst^[18].

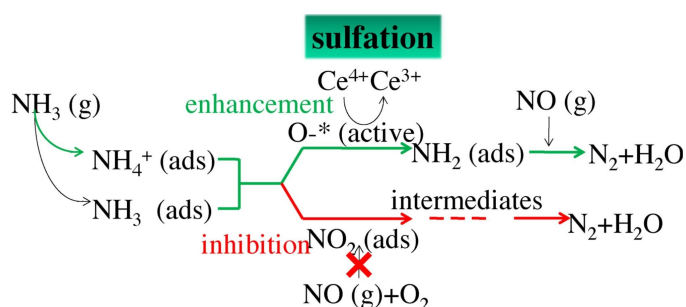


Figure 12. Reaction mechanism of Ce-Fe/WMH catalyst^[19].

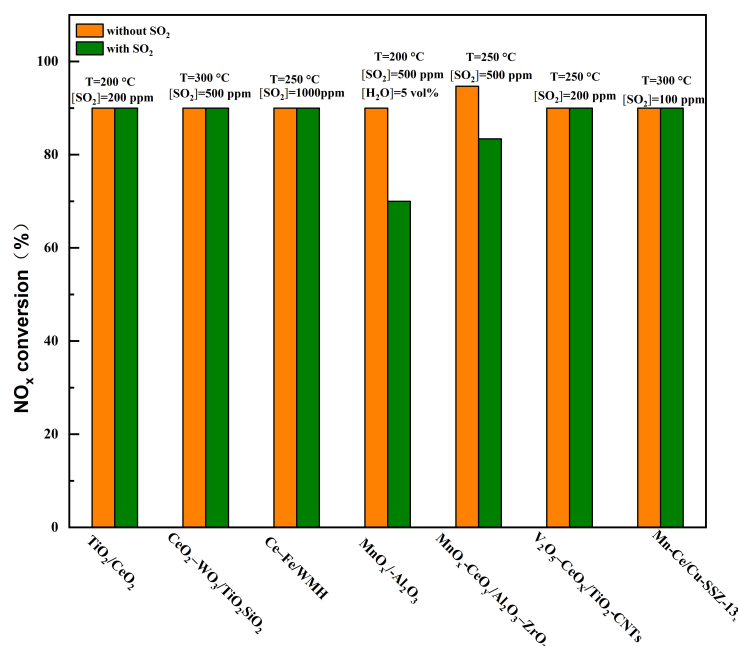


Figure 13. Typical SO₂-tolerant catalysts with different supports at low temperatures for SCR reaction^{[18][20][19][21][22][23][24]}.

3.3. Composite Oxide Catalysts

In recent years, extensive work has been done in the area of composite oxide catalyst. Compared with the supported catalysts, these catalysts all use metal oxides and have no clear support or active components. Many achievements have been made in the research of the catalytic performance and sulfur tolerance of catalyst.

3.4. Other Strategies to Improve the SO₂ Resistance

Among all the measures to enhance the sulfur resistance of the catalyst, some researchers have tried to find the influence of preparation methods, acidification, preparation and reaction conditions, further promoting the acidity and catalytic efficiency of catalyst.

4. Conclusions and Perspectives

Facing progressively strict legislation and policies to control NO emission, the research and design of low-temperature catalysts for NH₃-SCR have received a great deal of attention. Although the poisoning mechanism of catalyst at low temperatures has been studied thoroughly, how to maintain the high catalytic efficiency of catalyst at low temperatures and promote the SO₂-resistance-poisoning ability of catalyst to achieve practical application is still an urgent problem.

In the presence of O₂, SO₃ is easily formed on the catalyst due to the oxidation reaction of SO₂, and further combined with NH₃ to produce NH₄HSO₄ and/or (NH₄)₂SO₄, which can deposit on the surface of catalyst and inhibit the reaction gas to be adsorbed on the catalyst to participate in the SCR reaction. Besides, sulfate of active components can be formed and cause irreversible deactivation of catalyst. Therefore, it is effective to adopt several measurements to improve the SO₂ tolerance. Firstly, it is to reduce the adsorption of SO₂ on the catalyst. The highly acidic catalysts are effective to prevent the SO₂ adsorbing. Secondly, preventing the oxidation of SO₂ to SO₃ plays a significant role in high SO₂ tolerance by reducing the redox ability of catalyst, which can cut off the oxidation of SO₂ to some extent. Furthermore, the synergistic effect between catalyst components can also improve the sulfur resistance of catalyst, such as the construction of sacrificial sites, which is responsible for the reduction of active components sulfation. Along with these existing excellent sulfur resistant catalysts, it is expected that future studies will focus on optimizing the supports and preparation methods and concentrating on the application of new structures and technology, which are effective strategies to improve the low-temperature SO₂ tolerance of SCR catalysts.

References

1. Chuanmin Chen; Yue Cao; Songtao Liu; Jianmeng Chen; Wenbo Jia; Review on the latest developments in modified vanadium-titanium-based SCR catalysts. *Chinese Journal of Catalysis* **2018**, 39, 1347-1365, [10.1016/s1872-2067\(18\)63090-6](#).
2. Qi Zhao; Bingbing Chen; Jin Li; Xianbin Wang; Mark Crocker; Chuan Shi; Insights into the structure-activity relationships of highly efficient CoMn oxides for the low temperature NH₃-SCR of NO_x. *Applied Catalysis B: Environmental* **2020**, 277, 119215, [10.1016/j.apcatb.2020.119215](#).
3. Dongmei Meng; Qian Xu; Yunlei Jiao; Yun Guo; Yanglong Guo; Li Wang; Guanzhong Lu; Wang-Cheng Zhan; Spinel structured CoMnOx mixed oxide catalyst for the selective catalytic reduction of NO_x with NH₃. *Applied Catalysis B: Environmental* **2018**, 221, 652-663, [10.1016/j.apcatb.2017.09.034](#).
4. Xiaolong Tang; Yiran Shi; Fengyu Gao; Shunzheng Zhao; Honghong Yi; Zongli Xi; Promotional role of Mo on Ce_{0.3}FeO_x catalyst towards enhanced NH₃-SCR catalytic performance and SO₂ resistance. *Chemical Engineering Journal* **2020**, 398, 125619, [10.1016/j.cej.2020.125619](#).
5. Ya-Juan Shi; Hang Shu; Yu-Hua Zhang; Hong-Mei Fan; Ya-Ping Zhang; Linjun Yang; Formation and decomposition of NH₄HSO₄ during selective catalytic reduction of NO with NH₃ over V₂O₅-WO₃/TiO₂ catalysts. *Fuel Processing Technology* **2016**, 150, 141-147, [10.1016/j.fuproc.2016.05.016](#).
6. Ruiben Jin; Yue Liu; Yan Wang; Wanglai Cen; Zhongbiao Wu; Haiqiang Wang; Xiaole Weng; The role of cerium in the improved SO₂ tolerance for NO reduction with NH₃ over Mn-Ce/TiO₂ catalyst at low temperature. *Applied Catalysis B: Environmental* **2014**, 148, 582-588, [10.1016/j.apcatb.2013.09.016](#).
7. Liam John France; Qing Yang; Wan Li; Zhihang Chen; Jianyu Guang; Dawei Guo; Lefu Wang; Xuehui Li; Ceria modified FeMnO_x—Enhanced performance and sulphur resistance for low-temperature SCR of NO_x. *Applied Catalysis B: Environmental* **2017**, 206, 203-215, [10.1016/j.apcatb.2017.01.019](#).

8. Ziran Ma; Xiaodong Wu; Ya Feng; Zhichun Si; Duan Weng; Lei Shi; Low-temperature SCR activity and SO₂ deactivation mechanism of Ce-modified V₂O₅–WO₃/TiO₂ catalyst. *Progress in Natural Science: Materials International* **2015**, 25, 342-352, [10.1016/j.pnsc.2015.07.002](#).
 9. Fan Cao; Sheng Su; Jun Xiang; Pengying Wang; Song Hu; Lushi Sun; Anchao Zhang; The activity and mechanism study of Fe–Mn–Ce/γ-Al₂O₃ catalyst for low temperature selective catalytic reduction of NO with NH₃. *Fuel* **2015**, 139, 232-239, [10.1016/j.fuel.2014.08.060](#).
 10. Lulu Li; Lei Zhang; Kaili Ma; Weixin Zou; Yuan Cao; Yan Xiong; Changjin Tang; Lin Dong; Ultra-low loading of copper modified TiO₂/CeO₂ catalysts for low-temperature selective catalytic reduction of NO by NH₃. *Applied Catalysis B: Environmental* **2017**, 207, 366-375, [10.1016/j.apcatb.2017.02.041](#).
 11. Rui Yang; Haifeng Huang; Yijie Chen; Xixiong Zhang; Hanfeng Lu; Performance of Cr-doped vanadia/titania catalysts for low-temperature selective catalytic reduction of NO_x with NH₃. *Chinese Journal of Catalysis* **2015**, 36, 1256-1262, [10.1016/s1872-2067\(15\)60884-1](#).
 12. Kyung Ju Lee; Pullur Anil Kumar; Muhammad Salman Maqbool; Komateedi N. Rao; Kwang Ho Song; Heon Phil Ha; Ceria added Sb-V₂O₅/TiO₂ catalysts for low temperature NH₃ SCR: Physico-chemical properties and catalytic activity. *Applied Catalysis B: Environmental* **2013**, 142, 705-717, [10.1016/j.apcatb.2013.05.071](#).
 13. Boxiong Shen; Ting Liu; Ning Zhao; Xiaoyan Yang; Lidan Deng; Iron-doped Mn-Ce/TiO₂ catalyst for low temperature selective catalytic reduction of NO with NH₃. *Journal of Environmental Sciences* **2010**, 22, 1447-1454, [10.1016/s1001-0742\(09\)60274-6](#).
 14. Xin Zhao; L. Huang; Hongrui Li; Hang Hu; Jin Han; Liyi Shi; Dengsong Zhang; Highly dispersed V₂O₅/TiO₂ modified with transition metals (Cu, Fe, Mn, Co) as efficient catalysts for the selective reduction of NO with NH₃. *Chinese Journal of Catalysis* **2015**, 36, 1886-1899, [10.1016/s1872-2067\(15\)60958-5](#).
 15. Li Xu; Chuan Shi; Bingbing Chen; Qi Zhao; Yongjun Zhu; Hermann Gies; Feng-Shou Xiao; Dirk De Vos; Toshiyuki Yokoi; Xinhe Bao; et al. Improvement of catalytic activity over Cu–Fe modified Al-rich Beta catalyst for the selective catalytic reduction of NO with NH₃. *Microporous and Mesoporous Materials* **2016**, 236, 211-217, [10.1016/j.micromeso.2016.08.042](#).
 16. Wenpo Shan; Fudong Liu; Hong He; Xiaoyan Shi; Chang-Bin Zhang; A superior Ce-W-Ti mixed oxide catalyst for the selective catalytic reduction of NO_x with NH₃. *Applied Catalysis B: Environmental* **2012**, 115, 100-106, [10.1016/j.apcatb.2011.12.019](#).
 17. Tengfei Xu; Xiaodong Wu; Yuxi Gao; Qiwei Lin; Jianfeng Hu; Duan Weng; Comparative study on sulfur poisoning of V₂O₅–Sb₂O₃/TiO₂ and V₂O₅–WO₃/TiO₂ monolithic catalysts for low-temperature NH₃-SCR. *Catalysis Communications* **2017**, 93, 33-36, [10.1016/j.catcom.2017.01.021](#).
 18. Lei Zhang; Lulu Li; Yuan Cao; Xiaojiang Yao; Chengyan Ge; Bin Gao; Yu Deng; Changjin Tang; Lin Dong; Getting insight into the influence of SO₂ on TiO₂/CeO₂ for the selective catalytic reduction of NO by NH₃. *Applied Catalysis B: Environmental* **2015**, 165, 589-598, [10.1016/j.apcatb.2014.10.029](#).
 19. Yun Shu; Tanana Aikebaier; Xie Quan; Shuo Chen; Hongtao Yu; Selective catalytic reaction of NO_x with NH₃ over Ce–Fe/TiO₂-loaded wire-mesh honeycomb: Resistance to SO₂ poisoning. *Applied Catalysis B: Environmental* **2014**, 150, 630-635, [10.1016/j.apcatb.2014.01.008](#).
 20. Yue Peng; Caixia Liu; Xueying Zhang; Junhua Li; The effect of SiO₂ on a novel CeO₂–WO₃/TiO₂ catalyst for the selective catalytic reduction of NO with NH₃. *Applied Catalysis B: Environmental* **2013**, 140, 276-282, [10.1016/j.apcatb.2013.04.030](#).
 21. Xiaojiang Yao; Tingting Kong; Shuohan Yu; Lulu Li; Fumo Yang; Lin Dong; Influence of different supports on the physicochemical properties and denitration performance of the supported Mn-based catalysts for NH₃-SCR at low temperature. *Applied Surface Science* **2017**, 402, 208-217, [10.1016/j.apsusc.2017.01.081](#).
 22. Long Qu; Caiting Li; Guangming Zeng; Mengying Zhang; Mengfan Fu; Jinfeng Ma; Fuman Zhan; Diqiang Luo; Support modification for improving the performance of MnO_x–CeO_y/γ-Al₂O₃ in selective catalytic reduction of NO by NH₃. *Chemical Engineering Journal* **2014**, 242, 76-85, [10.1016/j.cej.2013.12.076](#).
 23. Qian Li; Xiaoxu Hou; Hangsheng Yang; Zhaoxia Ma; Junwei Zheng; Fu Liu; Xiaobin Zhang; Zhongyong Yuan; Promotional effect of CeOX for NO reduction over V₂O₅/TiO₂-carbon nanotube composites. *Journal of Molecular Catalysis A: Chemical* **2012**, 356, 121-127, [10.1016/j.molcata.2012.01.004](#).
 24. Qingling Liu; Zhenchao Fu; Lei Ma; Hejingying Niu; Caixia Liu; Junhua Li; Ziyin Zhang; MnO–CeO₂ supported on Cu-SSZ-13: A novel SCR catalyst in a wide temperature range. *Applied Catalysis A: General* **2017**, 547, 146-154, [10.1016/j.apcata.2017.08.024](#).
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