

Electronic Supplementary Information (ESI)

Enhanced Catalytic Performance of $V_2O_5-WO_3/Fe_2O_3/TiO_2$ Microspheres for Selective Catalytic Reduction of NO by NH_3

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1. The morphology of 1V10W3FeTMS catalysts

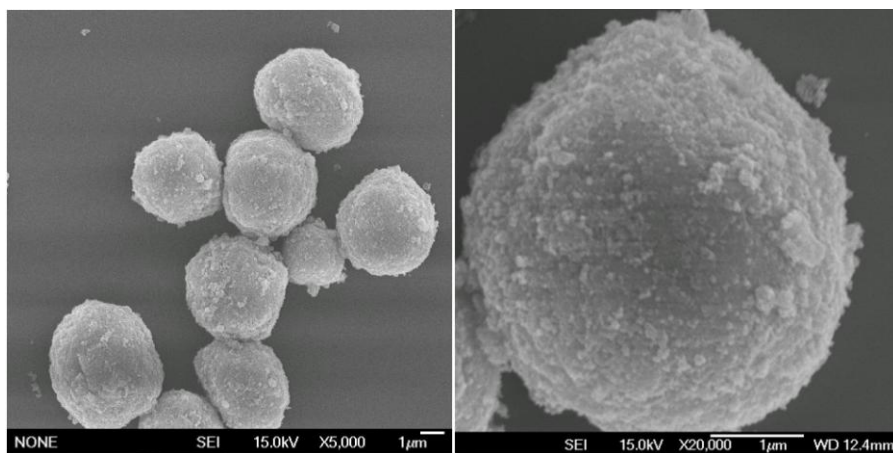


Fig. S1 SEM images of the 1V10W3FeTMS catalyst.

As shown in Fig. S1, after the TMS was impregnated with tungsten and vanadium precursor and calcination at 550 °C, they still maintained the spherical shape, and showed the uniform size distribution with the average diameter of 3 μm. This is a characteristic structural feature of the titania microspheres [1, 2].

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2. The NH₃-TPD profiles of 1V10WTMS and 1V10WFeTMS catalysts

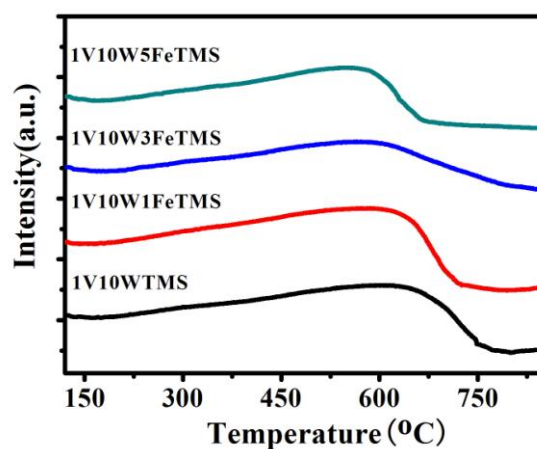


Fig. S2 NH₃-TPD profiles of 1V10WTMS and 1V10WFeTMS catalysts.

The NH₃-TPD results of the different catalysts are showed in Table 1 and discussed in the main text.

3. Effect of the different components

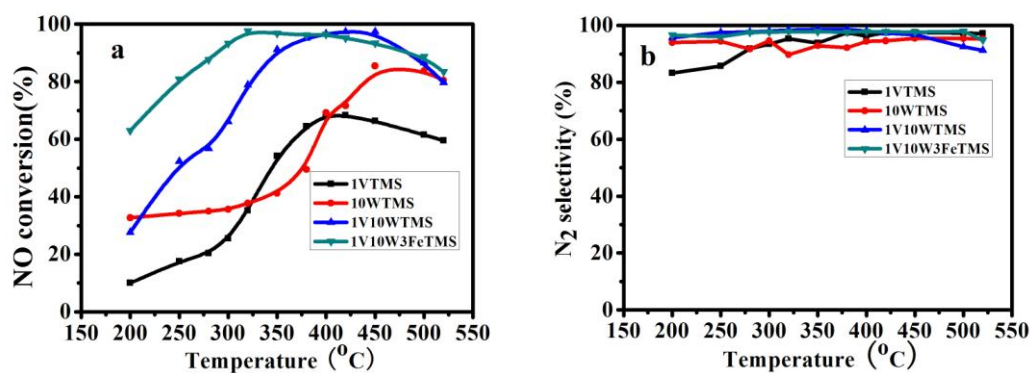


Fig. S3 (a) NO conversion against temperature under SCR condition over different samples (b)N₂ selectivity against temperature under SCR condition over different samples: (B)1VTMS; (D)10WTMS;(F) 1V10WTMS; (H)1V10W3FeTMS.

To know the effect of each element in the ternary VVFeTMS catalyst, the SCR

activity of monadic and binary catalysts was investigated. The results are provided in Fig. S3. When one or two elements among V, W, and Fe were removed from V1W10Fe3Ti in the catalyst preparation, the NO conversion decreased. Monadic catalyst V1Ti showed higher NO conversion when the temperature is higher than 380 °C, whereas W10Ti showed higher NO conversion with temperature higher than 400°C. Binary catalysts V1W10Ti increased the NO conversion below 300 °C. Ternary VWFeTMS catalyst increases further the NO conversion below 300 °C. These results indicate that Fe, V, and W elements supported on TiO₂ have a positive synergetic interaction and accelerate the SCR reaction.

4. Effect of the supports

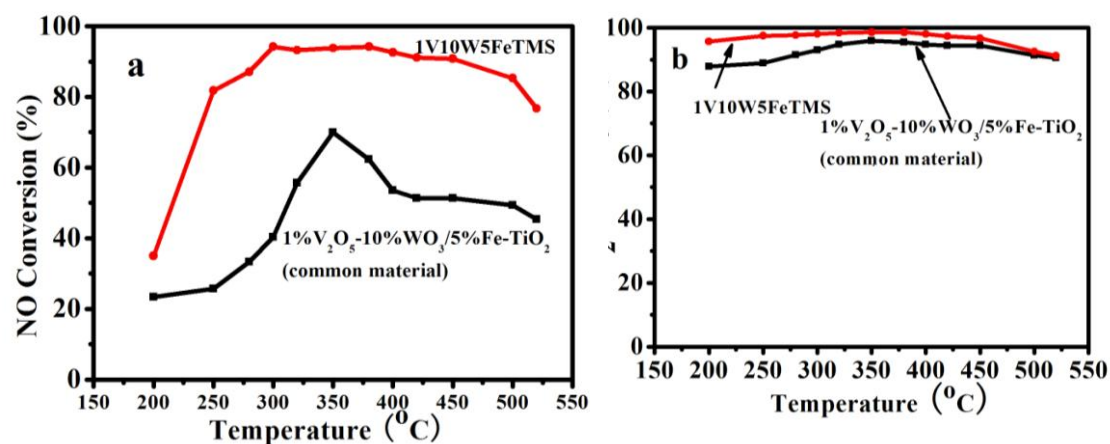


Fig. S4 (a) NO conversion against temperature under SCR condition over different samples (b)N₂ selectivity against temperature under SCR condition over different samples: (B) 1V10W5FeTMS; (D) 1%V₂O₅-10%WO₃/5%Fe-TiO₂ (common material).

The results of SCR over 1V10W5FeTMS, and TiO₂(common material) are shown in Fig. S4. Comparison of 1%V₂O₅-10%WO₃/5%Fe-TiO₂ (common material)

reveals that the performance of 1V10W5FeTMS is higher than that of 1% V₂O₅-10%WO₃/5%Fe-TiO₂ (common material) . This phenomenon uncovers that the mesoporous spheres and the unique structure of TiO₂ (microshpere) are more favorable for the title reaction than the TiO₂ (common material) counterpart.

5. Effect of the calcination temperature

Fig. S5 characterizes the effect of calcination temperature on the catalytic behavior of the 1V10W3FeTMS catalyst. The catalyst exhibited its best performance at a calcination temperature of 550 °C. Calcination temperatures above or below 550 °C led to decreased activity. These results can be explained by the gradual decomposition of the intrinsic structure of the active species on the catalyst at calcination temperatures of 650 °C . Thus, 550 °C was chosen as the optimum calcination temperature for the 1V10W3FeTMS catalyst, because the vanadium oxide species, tungsten oxide species and iron oxides species could not be fully activated at lower calcination temperature.

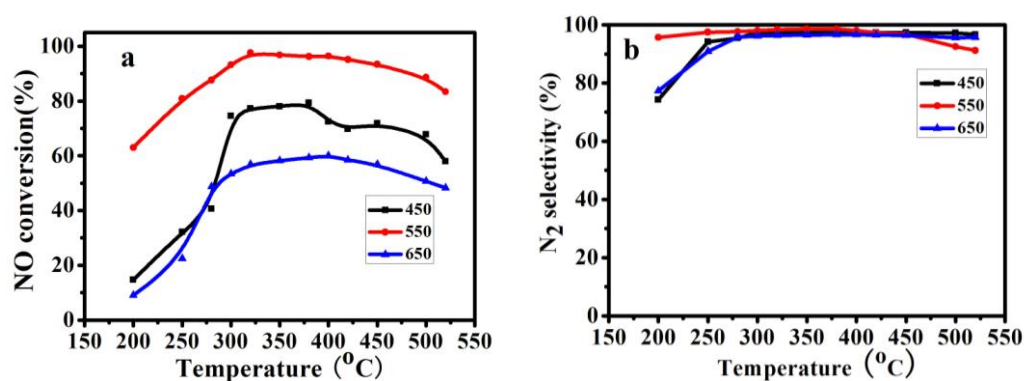


Fig. S5 (a) NO conversion against temperature under SCR condition over different samples (b)N₂ selectivity against temperature under SCR condition over different samples: (B)

1V10W3FeTMS-450°C; (D) 1V10W3FeTMS-550°C (D) 1V10W3FeTMS-650°C.

6. The reusability of 1V10W3FeTMS catalyst

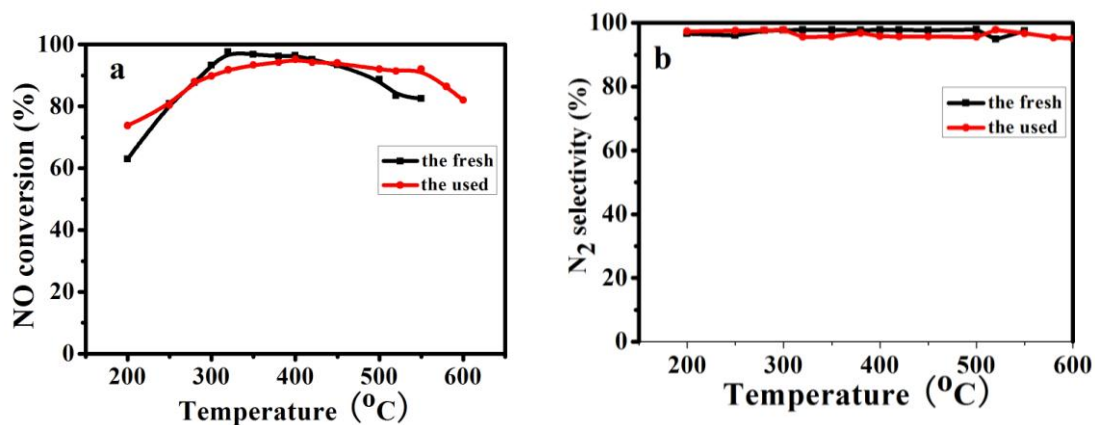


Fig. S6 (a) NO conversion against temperature under SCR condition over the 1V10W3FeTMS and the used catalysts (b)N₂ selectivity against temperature under SCR condition over the 1V10W3FeTMS and the used catalysts.

The reusability of 1V10W3FeTMS is also listed in Fig. S5. The NO conversion and selectivity of N₂ over the 1V10W3FeTMS catalyst still keep above 80% and 92% after the second cycle. This finding clearly indicates that the 1V10W3FeTMS catalyst shows good stability.

7. The conversion of NO to NO₂ over the different catalysts

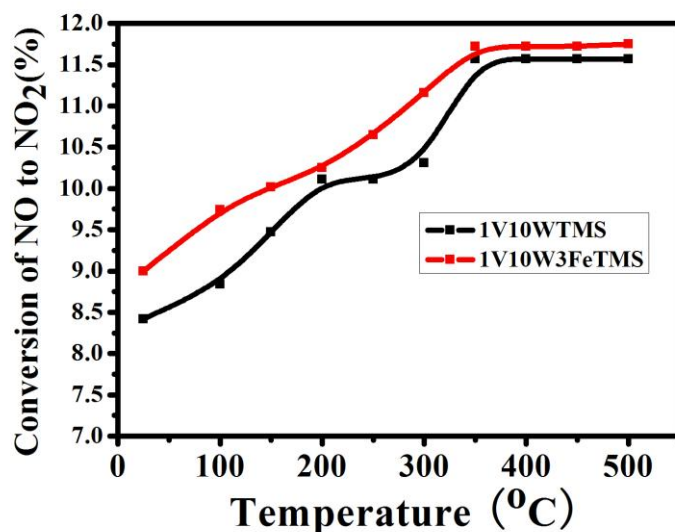


Fig. S7 Oxidation activity of NO to NO₂ by O₂ over the 1V10WTMS and 1V10W3FeTMS catalysts.

The conversion of NO to NO₂ over the different catalysts was measured and listed in **Fig. S7**. It is obvious that the introduction of Fe additive to the catalyst increases the oxidation activity of NO to NO₂. It has been accepted that NO is more readily reduced to N₂ along with portion NO₂ than single NO by NH₃. The introduction of iron oxide increases the oxidation rate of NO to NO₂, which increase the SCR activity.

References

- [1] C. W. Guo, Y. Cao, S. H. Xie, W. L. Dai, K.-N. Fan, *Chem. Commun.*, 2003, 700.
- [2] X.L. Yang, W.L. Dai, C. Guo, H. Chen, Y. Cao, H. Li, H. He, K. Fan, *J. Catal.*, 2005, **234**, 438.