Reactivity of  $V_2O_5/TiO_2$  catalysts in  $NH_3$  selective catalytic reduction as a function of the oxidation state of vanadium precursor

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Marine SCR catalyst requires higher DeNOx activity in the low temperature range comparing to automobile. In this study, we aimed at finding the optimum condition to prepare  $V_2O_5/TiO_2$  catalyst known as low temperature SCR catalysts by changing oxidation states in vanadium precursor solution.  $V_2O_5$  catalysts supported by  $TiO_2$  (1, 3, 5 and 7 wt.%) were prepared by applying wet impregnation method using three precursor solutions with different oxidation states ( $V^{3+}$ ,  $V^{4+}$  and  $V^{5+}$ ). We utilized BET, ICP, XRD, Raman spectroscopy, XPS, UV-vis DRS and  $H_2$ -TPR to investigate the physicochemical properties catalysts. Also,  $NH_3$ -SCR reaction test was performed for these catalysts. It was found that  $V_2O_5/TiO_2$  catalysts prepared from the precursor solution of  $V^{3+}$  oxidation state (VT(3+)) was the most active and selective to  $N_2$  production simultaneously, produced the least amount of  $N_2O$ . The formation of highly coordinated polymeric vanadyl species on the sample attributes the highest NOx conversion and  $N_2$  selectivity of the VT(3+) catalyst.