

Temperature programmed study of NO_x reduction with H₂ on Pt/Al₂O₃ catalyst in the presence of excess oxygen

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The reduction of NO_x by H₂ was studied over a 0.3 wt% Pt/Al₂O₃ catalyst as a function of temperature. The formation of N₂O began at low temperature due to the chemisorption of NO on the Pt metal and the resulting oxidation of the Pt active sites. The formation of N₂ was possible at higher temperatures, because the energy state of the reactants was high and the oxidized Pt active sites were reduced again by H₂. The formation of NO₂ was predominant above 150 oC, due to the excessive oxidation activity of the catalyst. The effects of the Pt loading and reaction conditions on the reductive activity were also investigated. The formation of N₂O at low temperature was strongly related to the Pt loading, and that of N₂ at high temperature was related to the temperature range in which the reductive activity was obtained. Therefore, for the selective formation of N₂ resulting from the reduction of NO on the Pt catalyst, both reductive conditions and an adequate reaction temperature were found to be important factors.