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# Influence of Technological Factors on the Decomposition Process of Nitrous Oxide (I)

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#### **ABSTRACT**

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#### **Mini Review**

Nitric acid production represents the largest source of  $\rm N_2O$  in chemical industry, with global annual emission of 23.7 million tons of  $\rm N_2O$  per year.  $\rm N_2O$  has a high impact on the environment as greenhouse gas and stratospheric ozone depletory [1].

In this production, the formation of nitrous oxide is possible along two chemical routes [2]:

**1.** During the oxidation of ammonia in the contact reactor nitrous oxide  $(N_2O)$  is formed as by-product.

$$4NH_3 + 4O_2 = 2N_2O + 6H_2O \tag{1}$$

**2.** In the process of low-temperature catalytic reduction of nitrogen oxides with ammonia.

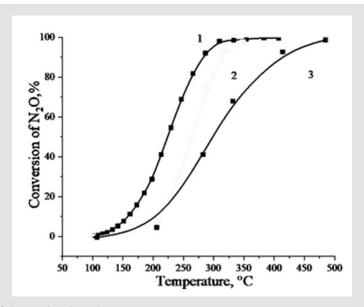
$$2NH_3 + 8NO \rightarrow 5N_2O + 3H_2O \tag{2}$$

$$4NH_3 + 3O_2 + 4NO \rightarrow 4N_2O + 6H_2O$$
 (3)

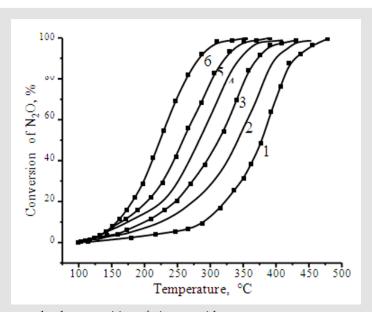
The efficiency of the catalytic reduction of nitrous oxide (I) is determined primarily by the activity of the catalyst used. The active site of CoO -exchanged catalyst for  $\rm N_2O$  decomposition is well known as divalent CoO (II) cations which closely collaborate on the  $\rm N_2O$  splitting. Thus, the formation of CoO-O is a key to achieve better performance in the  $\rm N_2O$  decomposition. The catalyst preparation method plays an important role in controlling the catalyst activity [3]. The catalyst's activity towards  $\rm N_2O$  decomposition is often significantly affected by the various components such as  $\rm H_2O$ ,  $\rm O_2$  in the flue gas. The catalyst activity inhibition from  $\rm H_2O$  and  $\rm O_2$  should be paid serious attention to. Therefore, the effects of the

components in the flue gas on the catalyst's activity have also been investigated in the study. Catalysts based on cobalt ferrite are highly active in the decomposition of nitrous oxide in the production of nitric acid. The residual nitrous oxide content depends on the selected process conditions. On the same catalyst, catalytic activity may vary. The paper presents the dependence of the degree of conversion on almost all technological factors: temperature, pressure, concentration of water vapor,  $H_2$  and  $O_2$ .

As shown in Figure 1, when pressure increases to 1.1 MPa, the influence of the ratio of components in the catalyst becomes more significant in terms of catalytic activity. So, on cobalt ferrite of stoichiometric composition, a 50% degree of decomposition of N<sub>2</sub>O is observed at 478 °C, and 100% at 548 °C. The increase in pressure leads to a significant decrease in the temperature range of the catalyst. At 255 °C, a 50% conversion of N<sub>2</sub>O is observed, and at 347 °C nitrous oxide is completely decomposed. Increasing the content of cobalt oxide to 44% in the catalyst makes it possible to reduce the temperature by half and to achieve complete decomposition of N<sub>2</sub>O at 36-45 °C, respectively. The reduction of nitrous oxide (I) contained in exhaust gases in various technologies can be carried out at pressure from 0.4 to 1.1 MPa as it's presented in Figure 2. For this reason, we conducted detailed studies on the effect of pressure on the decomposition of nitrous oxide. Figure 2 shows the dependences of the degree of decomposition of N<sub>2</sub>O on temperature and pressure. It is shown that increase in pressure from 0.1 MPa to 1.1 MPa causes increase of degree of decomposition of nitrous oxide by 6 times. So, for example, at 310 °C the degree of decomposition is 17% and 99.5% for 0.1 and 1.1 MPa respectively.



**Figure 1:** Catalytic activity of the catalyst based on  $CoFe_2O_4$ . Note: 1-Fe<sub>2</sub>O<sub>3</sub>: CoO = 1.1; 2-Fe<sub>2</sub>O<sub>3</sub>: CoO = 0.6; 3-Fe<sub>2</sub>O<sub>3</sub>: CoO = 1.



**Figure 2:** The effect of pressure on the decomposition of nitrous oxide. Note: The pressure in the reactor: 1 - atm; 2 - 2 atm; 3-4 atm; 4 to 6 atm; 5 to 8 atm; 6 -11 atm.

Also, the effect of oxygen and water on the decomposition of  $\rm N_2O$  was observed. Water inhibition effect and oxygen one differs. Water simply blocks a catalyst's site. Role of oxygen could be presented in two stages. At the first stage of  $\rm N_2O$  decomposition reaction, oxygen competes with nitrous oxide for active sites. At the second stage  $\rm N_2O$  mixes with oxygen and starts recombination, which creates idle cycles. "The work was carried out as part of the state assignment for the implementation of research. Theme number FZZW-2020-0010 ».

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