

## Additive polymerization of norbornene in the presence of highly effective catalytic systems on the basis of Ni(0)/HA/BF<sub>3</sub>·OEt<sub>2</sub>

© Vitaliy S. Tkatch,<sup>1,2,\*</sup> Dmitriy S. Suslov,<sup>1,2</sup> Olga V. Gubaidullina,<sup>1</sup>  
Mikhail V. Bykov,<sup>1,2</sup> and Marina V. Belova<sup>1</sup>

<sup>1</sup>Department of physical and colloidal chemistry. Chemical faculty.  
Irkutsk state university. K. Marx St., 1. Irkutsk, 664003. Russia.

Phone: +7 (3952) 52-10-82. E-mail: tkach@chem.isu.ru

<sup>2</sup>Institute of petro- and coal- chemical synthesis at Irkutsk state university. K. Marx St., 1. Irkutsk, 664003.  
Russia. Phone: +7 (3952) 52-10-82. E-mail: tkach@chem.isu.ru

\*Supervising author; <sup>+</sup>Corresponding author

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### Abstract

A number of catalytic systems on the basis of complexes Ni(0) and BF<sub>3</sub>·OEt<sub>2</sub> have been tested in polymerization of norbornene. It is established that the addition to catalytic system Ni (PPh<sub>3</sub>)<sub>4</sub> + 400BF<sub>3</sub>·OEt<sub>2</sub> of controllable quantities of water in the interval of molar relations from 2 to 104 leads to considerable growth of activity of catalytic systems (~27 times) and to the reduction of characteristic viscosity of polymers. At the use of catalytic systems (CH<sub>2</sub>=CH<sub>2</sub>)Ni(PPh<sub>3</sub>)<sub>2</sub>/6H<sub>2</sub>O/200BF<sub>3</sub>·OEt<sub>2</sub> the activity in polymerisation of norbornene increases 3 times as compared to the system Ni(PPh<sub>3</sub>)<sub>4</sub>/9H<sub>2</sub>O/400BF<sub>3</sub>·OEt<sub>2</sub>. Influence of molar relations H<sub>2</sub>O/Ni and norbornene/Ni, as well as the temperatures of reaction on the yield and characteristic viscosity of polynorbornenes, obtained in the presence of catalytic systems Ni(acac)<sub>2</sub>/2PPh<sub>3</sub>/5AlEt<sub>3</sub>/nC<sub>2</sub>H<sub>4</sub>/mH<sub>2</sub>O/200BF<sub>3</sub>·OEt<sub>2</sub> is studied. Among the studied proton donors (H<sub>2</sub>O, acacH, HBF<sub>4</sub>·OEt<sub>2</sub>, MeOH) methanol as the activator is the most effective. The structure of polynorbornenes has been studied with the methods of NMR <sup>1</sup>H, <sup>13</sup>C and IR spectroscopy.