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Conference Proceedings:
CHEMICAL ENGINEERING JOURNAL (Elsevier)
CHEMICAL & ENGINEERING PROCESSING:
PROCESS INTENSIFICATION (Elsevier)

VP-9. Ivashkina E.N., Nurmakanova A.E., Boichenko S.S.

PROCESS PARAMETERS' INFLUENCE ON EFFICIENCY OF ALKYLATE COMPOSITION AND OCTANE NUMBER

Tomsk Polytechnic University, Tomsk, Russia

VP-10. Kukueva V.V.

COMPUTATIONAL CHEMISTRY AS A TOOL FOR STUDYING OF THE PROCESSES IN THE CHEMICAL REACTORS ON THE MOLECULAR STAGES LEVEL

State Institution "Institute of Environmental Geochemistry" NAN Ukraine, Kiev, Ukraine

VP-11. Mityanina O.E., Samborskaya M.A.

OUTPUT MULTIPLICITY ANALYSIS OF METHYL TERT-BUTYL ETHER REACTIVE DISTILLATION SYNTHESIS

Tomsk Polytechnic University, Tomsk, Russia

VP-12. Nurmakanova A., Ivashkina E., Boychenko S.

ANALYSIS OF THE DEACTIVATION OF SULFURIC ACID DURING THE ALKYLATION OF ISOBUTANE WITH BUTENES

Tomsk Polytechnic University, Tomsk, Russia

VP-13. Pisarev M., Dolganov I., Dolganova I., Ivashkina E.

IMPROVING THE EFFICIENCY OF TECHNOLOGICAL GAS LOW-TEMPERATURE SEPARATION UNIT DEVICES USING SIMULATION DYNAMIC MODEL

Tomsk Polytechnic University, Tomsk, Russia

VP-14. Putin A.Y., Katsman E.A., Temkin O.N., Bruk L.G.

THE HYDROCARBOXYLATION OF ALKENES COUPLED WITH CARBON MONOOXIDE OXIDATION IN THE SYSTEM PdBr₂-CuBr₂-TETRAHYDROFURAN-H₂O

Moscow Technological University, Moscow, Russia

VP-15. Sinitsin S.A., Petrov A.Yu., Nefedova N.V., Shulyaka S.E.

PROSPECTIVE FERRITE CATALYSTS FOR FLUE GASES CLEANING FROM CARBON MONOXIDE

D. Mendeleyev University of Chemical Technology of Russia, Moscow, Russia

VP-16. Tataurshchikov A.A., Ivanchina E., Krivtcova N.

APPLICATION OF INDUSTRIAL DATA OF LG-24/7 IN MODEL OF SULFUR COMPOUND KINETICS IN THE HYDROTREATING PROCESS

Tomsk Polytechnic University, Tomsk, Russia

VP-17. Tungatarova S., **Baizhumanova T.,** Zheksenbaeva Z., Abdukhalykov D., Zhumabek M., Kassymkan K.

SILICON-TUNGSTEN HETEROPOLY COMPOUNDS - THE ACTIVE CATALYST COMPONENTS OF INCOMPLETE OXIDATIVE CONVERSION OF C_1 - C_2 ALKANES

D.V. Sokolsky Institute of Fuel, Catalysis and Electrochemistry, Almaty, Kazakhstan

VP-18. Tungatarova S., Baizhumanova T., Zheksenbaeva Z., Zhumabek M., Kassymkan K. CONVERSION OF LIGHT ALKANES INTO OXYGENATES

D.V. Sokolsky Institute of Fuel, Catalysis and Electrochemistry, Almaty, Kazakhstan

VP-19. Tungatarova S., Baizhumanova T., Zhumabek M.

OXIDATION OF LIGHT ALKANES TO HYDROGEN-CONTAINING COMPOSITIONS OVER SUPPORTED BLOCK CATALYSTS

D.V. Sokolsky Institute of Fuel, Catalysis and Electrochemistry, Almaty, Kazakhstan

Oxidation of Light Alkanes to Hydrogen-Containing Compositions over Supported Block Catalysts

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Tests of the ceramic block catalysts in oxidation of CH₄ were conducted in flow apparatus at atmospheric pressure in a quartz reactor with internal diameter of 0.025 m. The prepared catalyst block is wrapped in fiberglass and placed in a reactor. Initial reaction mixture was introduced at a rate of 50 ml/min after purging the system with inert gas (Ar) for 30 min at varying the ratio of reaction gases CH₄ and O₂ in a mixture from 1 : 1 to 4 : 1 and gradually increasing the temperature from 700 to 875°C and volume rate from 1000 to 10000 h⁻¹.

The process of oxidative conversion of hydrocarbon mixtures which are close to real while gradually increasing the reaction temperature from 700 to 875°C was investigated. Increase of methane conversion from 28 to 48% is observed with the growth of temperature. The amount of the produced H₂ increases from 33% to 84% at 850°C and again decreases to 70.2% at 875°C. A small amount of CO is formed at a temperature 700°C which increases with increasing temperature from 0.6 to 40.5%. Formation of C₂H₄ is observed at a reaction temperature 750°C in an amount of 13.0% and with the growth of temperature is lowered to 4%. Thus, the temperature 850°C is optimal for the formation of hydrogen, and 875°C and 750°C - for CO and C₂H₄, respectively. Effect of space velocity on direction of oxidative conversion of hydrocarbon mixture, which is close to the real, at variation of space velocity from 1000 to 10000 h⁻¹ was studied. It was found that the hydrogen yield was 40.5% at 1000 h⁻¹. Amount thereof increased to 84.0% at 5000 h⁻¹ and then reduced to 45.1% with increasing space velocity up to 10,000 h⁻¹. The same dependence is observed for the formation of CO and ethylene. The amount of CO is increased from 10.5% to 38.0% and ethylene - from 7.8% to 13.0% at 5000 h⁻¹. Product yield gradually decreases with further increase in space velocity to 10000 h⁻¹. Increasing the space velocity from 1000 h⁻¹ to 5000 h⁻¹ positively affects on the conversion of initial mixture

to desired products. 5000 h⁻¹ is optimal space velocity for the formation of desired products. Thus, optimum ratios of main gases in the reaction mixture were also determined.

Thus, optimization of technological modes of oxidative conversion of hydrocarbon mixture was carried out. Effect of varying the reaction temperature, space velocity and ratio of gases for selective production of desired products was investigated. The temperature ranges of $725-875^{\circ}$ C and space velocity 5000 h⁻¹ are optimal process parameters to produce hydrogen compositions. It is determined that the oxidative conversion of methane to hydrogen-containing mixture is carried out at the optimum ratio of components $CH_4: O_2 = 2: 1$ at a concentration of gases (50%: 25%), respectively.

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