

Aromatization of methane and other hydrocarbons over Mo₂C catalysts

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The upgrading of lower alkanes is an important subject of heterogeneous catalysis. Early studies focussed mainly on Pt catalysts, which exhibited an outstanding catalytic performance among the metals. All these studies were exclusively restricted to the reactions of C₂-C₇ hydrocarbons. The discovery of Wang et al.[1] that methane can be converted into benzene on MoO₃/ZSM-5 opened a new route for the utilization of methane. It turned out, however, that not the MoO₃ but Mo₂C is the key component for the activation and aromatization of methane, which is formed from MoO₃ during the induction period of the reaction. Methane can be converted into benzene on Mo₂C/ZSM-5 with ~80% selectivity at a conversion of ~15% at 973 K. In our laboratory, we have worked in two directions: (i) elaborating the effect of Mo₂C on the aromatization of C₁-C₇ hydrocarbons, and (ii) studying the chemistry of hydrocarbon species (CH₂, CH₃, C₂H₅, C₃H₅, C₄H₉ and C₅H₁₁), the primary products of the activation of the above compounds, on Mo₂C/Mo(100) in UHV by several spectroscopic methods.

[1] L.Wang, L. Tao, M.Xie, G.Xu, J.Huang, Y.Xu, Catal. Lett. 21 (1993) 35.