

either of the compounds comprising the catalyst is increased. The apparent activation energy, E_{app} , is 15.3 kcal/mole.

Effect of Solvents on the Rate of Radical Formation in the Thermal Decomposition of azo-bis-Isobutyronitrile

L. M. ANDRONOV AND G. E. ZAYKOV

*Institute of Chemical Physics,
Academy of Sciences, USSR*

The authors have measured the rates of initiation of chain reactions in thermal decomposition of azo-bis-isobutyronitrile in media comprising mixtures of methyl ethyl ketone with benzene, nitrobenzene, or water at 40–70°C. In methyl ethyl ketone, the activation energy for removal of a radical from the cage, E_s , is 1.9 kcal/mole. Addition of nitrobenzene does not affect E_s , but that of benzene decreases it and that of water increases it.

Catalytic Activity of Alkali Metal Ions in Oxidative Dissolution of Diamonds

A. P. RUDENKO, I. I. KULAKOVA,
AND A. A. BALANDIN

*Department of Chemistry, The Lomonosov
State University, Moscow*

The authors have studied the oxidative dissolution of diamonds in superheated steam and in carbon dioxide at 870–880° over alkali metal carbonates or hydroxides. Li^+ , Na^+ , K^+ , Rb^+ , and Cs^+ promote this oxidation in superheated steam and suppress it in CO_2 . The highest activity is exhibited by K^+ , and the lowest by Li^+ and Na^+ ; the activities of Rb^+ and Cs^+ are lower than that of K^+ , but much higher than those of Li^+ and Na^+ . It appears that the activity of these ions is induced by the formation of an intermediate alkoxide-resembling complex.

Kinetics of Biphotonic Reactions. The System Carbazole-Ethanol

Z. A. SINITSYNA AND KH. S. BAGDASARYAN

The Karpov Institute of Physical Chemistry

The authors have studied the kinetics of biphotonic reactions in which the second photon is absorbed when the molecule is a triplet. They have shown that in the system carbazole-ethanol at 77°K, there are considerable deviations from the I^2 relationship at low (~3%) degrees of filling of the triplet level. Data obtained on discontinuous illumination are well correlated by a

theoretically derived equation containing a factor describing this deviation.

Temperatures and Rates of Reduction of PbO_2 , Pb_3O_4 , and PbO by Hydrogen and Carbon Monoxide

D. K. LAMBIYEV AND M. S. KURCHATOV

*Institute of General and Inorganic Chemistry,
Bulgarian Academy of Sciences, Sofia*

A method allowing direct measurement of sample temperature while also measuring the reaction rate was used to study the reduction of lead oxides with gases. The reduction has no induction period. Its rate decreases from the very beginning, even though the sample temperature rises quite high during the initial period of the reaction. The authors have studied the effect of addition of oxygen on the reduction, and have shown that in some cases the reduction process starts inside the solid sample.

Catalysis Occurring during Thermal Decomposition of Silver Oxide

M. M. DUBININ, P. KADLETS, AND V. PONETS

*Institutes of Physical Chemistry of the Academies
of Sciences of the USSR and Czechoslovakia*

Direct quantitative measurements of liberated O_2 and CO_2 proved that silver oxide decomposes above 280°C. The reaction has a long induction period. Metallic dusts of silver or nickel deposited on the surface are very active catalysts of this reaction.

Effect of Additives on the Thermal Decomposition of Ammonium Perchlorate

A. V. BOLDYREVA, B. N. BEZRUKOV,
AND V. V. BOLDYREV

*Institute of Chemical Kinetics and
Combustion, Siberian Department,
Academy of Sciences, USSR*

The authors have studied the effects of particle size and specific surface on the catalytic activity of NiO or CuO in the low-temperature (thermally induced) decomposition of NH_4ClO_4 . Specific surface has a strong effect on the activity, while variation of particle size from 100 to 300 μ has virtually no effect. Oxides of copper, nickel, chromium, cobalt, etc., catalyze processes occurring during gas-phase decomposition of NH_4ClO_4 . It appears that these oxides not only catalyze the decomposition in the solid but also substantially affect the gas-phase processes.