

## Investigation of the strain behavior of amorphous polyurethane elastomers

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### Abstract

The strain behavior of two series of amorphous polyurethane elastomers on the base of olygoetherdiols and oligodienediols under stretching with a constant rate was investigated. Samples of the each series being varied in the network density value only demonstrated high-elastic behavior with some hardening effect in the final stage of the strain process. The stress versus strain dependence of investigated objects was described using the early proposed approach, taking into account high-elastic properties of elastomers and relaxation properties as well. A significant feature of the mentioned approach was separate accounting contributions to the stress value of stabile (chemical) cross-linking bonds and labile ones; this allowed taking into account the hardening effect on the total stress value in strained material. This accounting was provided using the transformation function which type depended on the type of considered polymer chains. The use of the mentioned function provided an adequate accounting varying in the polymer chains elasticity within the strain process. It was experimentally found that the chemical network density of investigated samples did not vary with the strain process up to the rupture allowing use of elements of basic equations of modern approaches in the framework of high-elasticity theory. The closeness of the experimental values of relaxation parameters for elastomers for the same series of samples allowed using average values of these parameters that lead to significant simplification of the applied calculations. Thus, bases for developing an engineer-theoretic approach to predict strain behavior of elastic materials under non-equilibrium conditions were created. The obtained parameters allowed adequately describing stress versus strain relationships for all investigated objects.

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