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Nota di contenuto PREFACE; INTRODUCTION; Chapter I. Brief information on types of

polymers and their chemical structure; Chapter II. Packing of macromolecules and polymer density; II.1. Increments method and basic physical assumptions; II.2. Relationship between free volume of polymers, coefficient of molecular packing and porous structure; Chapter III. Temperature coefficient of volumetric expansion; Chapter IV. Glass transition temperature of polymers; IV.1. Thermomechanical and other methods of evaluation of the glass transition temperature of

polymers; IV.2. Mechanism of glass transition

IV.3. Calculation of the glass transition temperature of linear polymersIV.4. Influence of plasticization on the glass transition temperature of polymers; IV.5. Calculation of the glass transition temperature of polymer networks; Chapter V. Temperature of transition into the viscous flow state for amorphous polymers; V.1 Estimation of temperature of transition into the viscous flow state of polymers; V.2 Dependence of Newtonian viscosity on molecular mass of polymer in a wide range of its change; Chapter VI. Melting point of polymers Chapter VII. Temperature of onset of intense thermal degradation of polymersChapter VIII. Optical and opto-mechanical properties of polymers; VIII.1 Refractive index; VIII.2 Stress-optical coefficient;

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Chapter XVII. Molecular design and computer synthesis of polymers with predetermined propertiesAppendix 1. Examples of solution of direct problems of polymer synthesis; Appendix 2. Examples of solving the reverse problem of polymer synthesis; Appendix 3. The example of solving the complex problem ... analysis of the chemical structure of phenol formaldehyde resin; Appendix 4. Application of the approach to multicomponent copolymers; Appendix 5. Influence of strong intermolecular interaction occurring between two dissimilar polymers on their miscibility

Appendix 6. On formation of super-molecular structures in amorphous polymers

## Sommario/riassunto

An approach to the quantitative analysis of the effect of the chemical structure of linear and network polymers on their properties is described. The approach is based on the representation of the repeating unit of the polymer in the form of a set of anharmonic oscillators which describe the thermal motion of atoms in the field of intra- and intermolecular forces, including weak dispersion forces, dipole-dipole interactions, hydrogen and valency bonds.