



## Polymerization: It's Fundamentals and Mechanism

Liu Zhou\*

Department of Chemistry, Shanghai University, Shanghai, China

### DESCRIPTION

With a long history, cationic polymerization is one of the most important polymerization methodologies in both academia and industry. The significant synthetic butyl rubber, first reported in 1937 and commercially produced in the early 1940s, was produced through cationic copolymerization of IsoButylene (IB) and IsoPrene (IP) in a slurry or solution process at low temperatures ranging from 50°C to 105°C. Academic and industrial efforts have been made to promote the development of butyl rubber due to its superior properties compared to natural rubber, wide range of applications, and remarkable global market growth. The highly active unstable centre of the carbocation, on the other hand, causes a large exothermic rapid polymerization and a variety of side reactions.

Miniemulsion polymerization has recently exploded in terms of publications and the development of a diverse range of useful polymer materials that are only available *via* this polymerization technique. The fundamental aspects involved in the preparation and polymerization of monomer miniemulsions are reviewed here. The use of miniemulsion polymerization for the creation of high solids low viscosity latexes, controlled radical polymerization for the dispersion of polymers with well-defined microstructures, catalytic polymerization in aqueous media, the preparation of hybrid polymer particles, anionic and step polymerization in aqueous dispersed media, and more are all covered.

Carbocationic polymerizations fundamentals and general characteristics, as well as a timeline and current references. Following general considerations, the fundamental elements of carbocationic polymerization are discussed, including monomers, initiating systems, solvents, and temperature. Carbocation stability, monomer nucleophilicity, and dynamic interactions are

all given special consideration. Current knowledge of the fundamental reactions, including copolymerization, initiation, propagation, termination, and transfer reactions, is presented in the section on the kinetics of carbocationic polymerization. It is discussed how to control (living) carbocationic polymerization.

As indicated by the numerous applications in both conventional fields like coatings, inks, and adhesives as well as high-tech fields like optoelectronics, laser imaging, stereolithography, and nanotechnology, photoinitiated polymerization is increasingly used in industry. The most recent advances in photoinitiating systems for free radical and cationic polymerizations are presented in this context. Moreover, strategies for resolving issues with efficiency, wavelength flexibility, environmental concerns, and safety concerns in all photoinitiating systems for diverse forms of activation are intended to be demonstrated. In the last ten years, much progress has been made in the preparation of complex and nano-structured macromolecules using photoinitiated polymerizations. Thus, new and emerging applications of photoinitiated polymerizations in biomaterials, surface modification, block and graft copolymer preparation, and nanocomposites have been addressed.

They describe the mechanism of direct oxidation of 3-alkylthiophenes with ferric chloride (FeCl<sub>3</sub>) as the polymerization oxidant/catalyst to produce high molecular weight Poly(3-Alkyl Thiophenes) (P3ATs), conjugated polymers with electrical and optical properties. This study shows that the presence of solid FeCl<sub>3</sub> in the reaction mixture is necessary for it to function as an oxidant during the polymerization of P3AT. On the basis of the crystal structure of FeCl<sub>3</sub> and quantum chemical computations of thiophene derivatives, a feasible polymerization mechanism for 3-alkylthiophene was developed. The polymerization is thought to take place *via* a radical mechanism rather than a radical cation mechanism.

**Correspondence to:** Liu Zhou, Department of Chemistry, Shanghai University, Shanghai, China, E-mail: zhou@gmail.com

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