
Review on: Analysis of Pharmaceutical Polymers

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Abstract

Pharmaceutical polymers are the foundation of pharmaceutical preparation, both traditional and innovative drug delivery systems. Drug release is sustained, extended, modified, controlled, and targeted through the application of polymers in various medication methods of administration and prescription types. The formulation the polymers for a range of pharmaceutical applications, packaging materials, and medical instruments in both traditional and regulated medication systems for distribution and primary healthcare can benefit from an intelligent choice for the outer and bulk properties. Packaging, with particular emphasis on the Nigerian setting. The study discusses recent trends and translational insights, opportunities and problems associated with combining polymeric technology, the role of edible polymers in pharmaceutical development, and various methods to polymer drug delivery systems. Despite constraints such as cost, quality, and regulatory compliance, polymers have a substantial impact on pharmaceutical manufacturing and patient care, particularly in drug delivery, controlled release, and packaging. The future of polymer technology in the pharmaceutical business depends on continuous research and unleashing the full potential of polymers.

Keywords - Polymer, pharmaceutical aid, conventional drug systems, packaging materials

INTRODUCTION

One type of materials that is frequently utilized in medication formulation and delivery systems is medicinal polymers. Their responsibilities include maintaining appropriate medication targeting, improving the stability of active pharmaceutical ingredients (APIs), and regulating drug release rates. To guarantee the safety, effectiveness, and quality of medications, it is essential to analyze these Polymer. ^[1]

Importance of Pharmaceutical Polymers

Polymers in pharmaceuticals serve various functions:

Controlled Release

Polymers are used to modulate drug release by forming matrix systems or coatings, ensuring a sustained release over time. ^[2]

Drug Delivery

Polymers like hydrogels and nanoparticles enable targeted delivery to specific tissues or organs, improving therapeutic outcomes.

Stability

Polymers protect sensitive APIs from environmental factors like moisture, light, or pH, maintaining the drug's effectiveness.

Solubility Enhancement

Poorly soluble drugs can be encapsulated in polymer matrices to enhance solubility and bioavailability.^[3]

Classification of Pharmaceutical Polymers

Original Source

Natural Polymers

These come from natural sources such as bacteria, plants, and animals. They are frequently utilized in wound healing and medication administration systems since they are biocompatible and biodegradable.

Example

Polysaccharides: alginate, cellulose, and starch.

Proteins: collagen, gelatine^[4]

Nucleic acids: RNA and DNA

Synthetic Polymers

These polymers were produced artificially by chemical synthesis. They are very customisable since they provide a great deal of control on their molecular weight, composition, and characteristics.^[4]

Example

Polyvinyl alcohol (PVA)

Polyethylene glycol (PEG)

Polymethyl methacrylate (PMMA)^[5]

Structure Based

Linear polymer

Long chains of monomers joined end to end are known as linear polymers.

Example

Polyethylene oxide (PEO)^[6]

Branched polymer

Polymers that have side chains that split off from the main chain are known as branched polymers.

Example

Dextran

Cross linked polymer

Polymers with chains that are connected at different locations to create a network are known as cross-linked polymers.

Example

Polyacrylic acid^[7]

Considering Solubility

Water-soluble polymers

Water-soluble polymers are used as binders, suspension stabilizers, and viscosity regulators.

Example

Polyethylene glycol.

Water-insoluble polymers

Water-insoluble polymers are frequently seen in systems with regulated release.

Example

Ethyl cellulose Polymethacrylates ^[8]

Biodegradability-based Eco-friendly Polymers

Biodegradability-based Eco-friendly Polymers: break down inside the body into by products that are safe for biology.

Example

Polylactic acid

Polyglycolic acid ^[9]

Non-biodegradable polymers

Non-biodegradable polymers are those that are either eliminated unaltered or stay whole in the body.

Example

Polymethyl methacrylate

Methylcellulose ^[10]

Ideal Characters of Polymers

Chemical Resistance

The ability to withstand a variety of chemicals, such as bases, acids, and solvents.

Mechanical Strength

The ability to endure physical stressors due to high tensile strength, flexibility, and impact resistance. ^[11]

Thermal Stability

The capacity to withstand deterioration and continue operating at high temperatures. Lightweight: Because of their low density, they are simple to manage and move.

Durability

The ability to withstand age, wear, and fatigue to guarantee a lengthy service life. Biocompatibility: Adaptability to biological tissues is essential for medicinal applications. ^[12]

Electrical Insulation

They are appropriate for electrical insulation applications because to their low electrical conductivity.

Transparency

Clarity and transparency may be crucial for optical applications.

Processability

The simplicity of manufacturing using techniques such as 3D printing, injection molding, and extrusion. ^[13]

Cost-Effectiveness

Maintaining desirable qualities at reasonable production costs ^[14]

The Characteristics of Polymers

Crystalline polymers

Semi Crystalline polymers

Viscosity

Amorphous polymers

Polymer complex

Syneresis

Adsorption of Micro molecule

Bio adhesivity of water-soluble polymer

Polymer Dissolution

Crystalline polymer

Between crystalline and amorphous regions, light scattering results in polymers to become opaque, along with these kinds of polymers are referred to as crystalline. Transparency is increased for both low and High levels of crystallization and polymers with a because the density of barriers is reduced. For instance, polypropylene that is atactic normally Amorphous and translucent, whereas Synthetic polypropylene, with the presence of crystals is 50%.^[15]

Semi Crystalline polymers

Semi-crystalline materials have highly organized molecular structures and a sharp melting point. When a certain amount of heat is absorbed, semicrystalline materials swiftly convert to a low viscosity fluid and remain solid.

Viscosity

Large macromolecular solutes may have a noticeable impact when present in liquid. When viscous rises, the duration of the drugs sustained release increases.

Amorphous polymers

Amorphous polymers are ones that do not have a crystalline structure when exposed to x-rays or electrons. For example, utilizing straining-induced contrast enhancement in TEM.^[16]

Polymer complex

Complexes can form in solution easily thanks to polymers, as can be shown when polyglycol and high molecular weight polyacid are combined in an aqueous solution. Up until a certain point, mixed polyacid and propylene equimolar combination maintains the same viscosity and pH in response to an increase in oligomer chain length. Only after Does When the tiny particles of the polymer (the hormone ethylene ethanol) attain a specific size? These highly particular macromolecular reactions primarily depend on the structure of the molecules, translational heat, and a number of other factors. Small molecules found in animals often undergo an intricate process that's critical to their function. The specific relationship between hyaluronic acid and the proteoglycans that comprise cartilage's internal structure was demonstrated through research. The investigations. Both the promotion of development of gels in alkaline carbohydrate environments or the powerful binding of calcium and other multivalent ions in polysaccharide structures like alginic acid solutions, can be explained by the coordination of calcium between certain polysaccharides that contain uronic acid. This is being proven that these interactions possess physiological significance.^[17]

Syneresis

Syneresis is the process of removing fluid in gel form that has enlarged. This type of instability occurs in both non-aqueous and aqueous gels. The division of the liquid layer is thought to be caused by the elasticity compression of the copolymer atoms. The macromolecules involved in the swelling process during gel formation stretch, and as swelling increases, so do the elastic forces. The restoring power of the polysaccharides is balanced at optimum by the expanding actions, which are established by the pressure of osmotic fluid.

Adsorption of Micro molecule

The stability of suspensions and emulsions is taking use of certain macromolecules' capacity to absorb at interfaces. At the interface, proteins, acacia, and gelatine absorb. Adsorption of this kind is occasionally undesirable, as in the instance of glucose sticking to infusions vials made of glassware. It is currently standard procedure to add albumin in order to inhibit adsorption. Albumin attaches itself to the glass. Binding is believed to happen in a general way and can occur on materials like polythene and glass. Macromolecules, such as hyaluronic acid, may help with

lubrication in joint fluids because they create a more polar surface in the solution. This can lessen the sticking of proteins like insulin, but it doesn't always stop it completely. [18]

Bio adhesivity of water-soluble polymer

Bio adhesivity of polymers with water soluble develops on organic surfaces. Sticks to a hydrophilic polymer or to a surface that the polymer has attached to. This sticking occurs through the bio adhesive's polymer chains must interact well with the mucosal surface to ensure good adhesion. The charges of the molecules are important, especially when two negatively charged (anionic) polymers are involved. Are not charged, their contact will be at its highest. The pH of association and penetration must be adjusted. The adhesion capabilities of There are four types of polymers: good (Carbopal), fair (gelatin), poor (povidone), and excellent (carboxymethylcellulose). The FDA has approved cationic chitosans and anionic poly (acrylic acid) (carbophil) derivatives. At pH where they are undissociated, polycarbophil and carbomer (carbopol 934P) exhibit maximum mucoadheivity with pka values of around 4.5. [19]

Analytical Techniques for Pharmaceutical Polymers

Spectroscopy Method

Chromatographic Techniques

Thermal Analysis

Microscopic and imagine Techniques

Rheological Techniques

Spectroscopy Method

FTIRI (Fourier transform Infrared Spectroscopy)

This technique determines a polymer's functional categories by measuring the absorption of infrared light at various frequencies. It helps validate the chemical structure and detect contaminants. [20]

NMR (Nuclear Magnetic Resonance Spectroscopy)

NMR uses an analysis of the magnetic characteristics of atomic nuclei to offer information on the molecular structure of polymers. It helps in understanding the polymerization process and identifying monomers. [21] The receptor and ligand-receptor complex benefit from NMR. [22]

Chromatographic Techniques

GPC (Gel permeation Chromatography)

A commonly used method for figuring out the molecular weight distribution of polymers is GPC, sometimes referred to as Size Exclusion Chromatography (SEC). It facilitates comprehension of the polymer's polydispersity index (PDI) and molecular weight. [23]

DSC (Differential Scanning calorimetry)

DSC tracks the heat flow that results from a polymer's phase changes, like melting or crystallization. Understanding the Melting Temperature and thermal stability of polymers is crucial. [24]

Thermal Analysis

TGA (Term gravimetric Analysis)

Thermal Resistance and breakdown profile of the polymer are indicated by TGA, which measures weight changes in the polymer as a function of temperature or time.

Microscopic and imagine Techniques

Scanning Electron Microscopy

SEM Provides detailed images of the polymer's surface morphology, allowing the observation of particle size, shape, and surface characteristics

Atomic force microscopy

AFM Gives topographical information and measures surface Roughness and texture on a Nano scale level. [25]

Rheological Techniques

Rheological analysis measures the flow and deformation behaviour of polymers, providing information on viscosity, elasticity, and plasticity. It's crucial for understanding the processing characteristics and mechanical behaviour of pharmaceutical polymers [26]

Mechanisms of Polymer-Mediated Drug Release

Dissolution

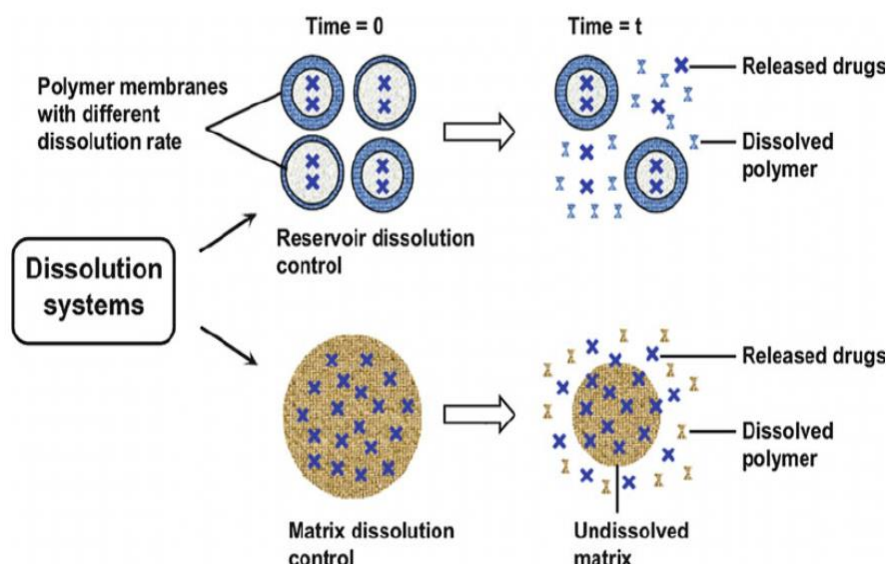


Figure 1: Release of Medication via A Dissolution-Controlled Polymerization Method

When a medication is released through the polymer dissolving and the dissolution fluid penetrating it. Drugs are either mixed into a matrix system or contained in single drug particles with slowly dissolving polymers in sustained or controlled delivery systems. The drug dissolves and is then released are determined by the rate at which the dissolving fluid penetrates the matrix. But the substrate has permeability, and presence of certain components determine how much dissolving fluid can penetrate. Hydrophobic additives, as well as the particle's surface and the system ability you can make a sustained release product by using coated particles with either a thin or thick coating in an encapsulated system. The thickness of the coating and its solubility in water affect how long it takes for the coating to dissolve. [27]

Diffusion

The In a controlled drug delivery system, the medicine disperses across the material's structure. and into the surrounding environment. The drug can be either released at a steady rate over time or triggered by specific conditions, such as pH or temperature changes. solid, diluted, or highly concentrated and is evenly spread across a compound that is matrices (monoliths matrices device) or it is encircled by a thin film and placed inside a polymer matrix (reservoir system). The medicine spreads as it leaves its plastic structure and affects the atmosphere around it. Since the bioactive agent has to travel over a considerable distance, the rate of diffusion and the properties of the surrounding medium can affect how quickly and effectively the drug reaches its target in these types of systems, the delivery rate often decreases with time and continuous drug release. Gradually, necessitating a longer diffusion period for the drug's final administration. Swelling-

controlled drug delivery method releases the medicine by causing the polymer to swell and then releasing the drug through diffusion with or without dissolution. [28]

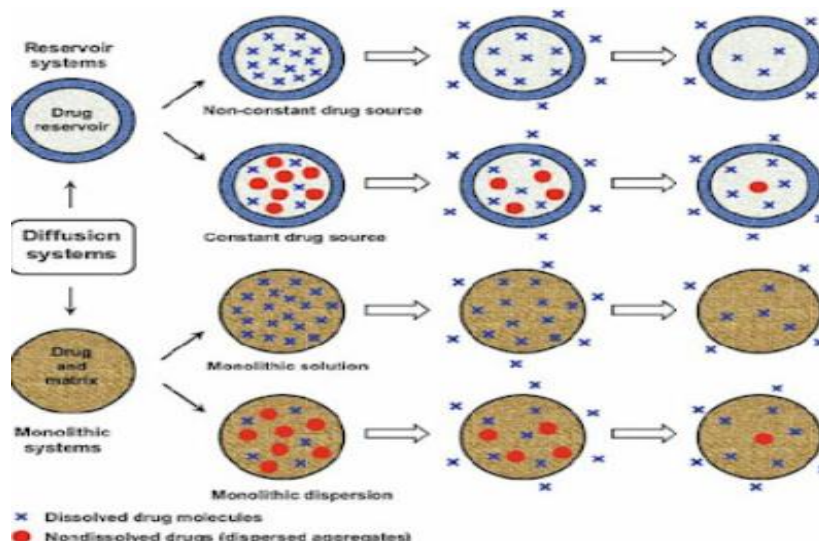


Figure 2: Release of medication by dispersion mediated method of material

Erosion

Active ingredient

This hydrolysed by bond A, which cleaves its covalent link to the polymer backbone and releases the active agent. Bond A should be much more reactive than bond B because we want the active agent to be released without any polymer fragments sticking to it.

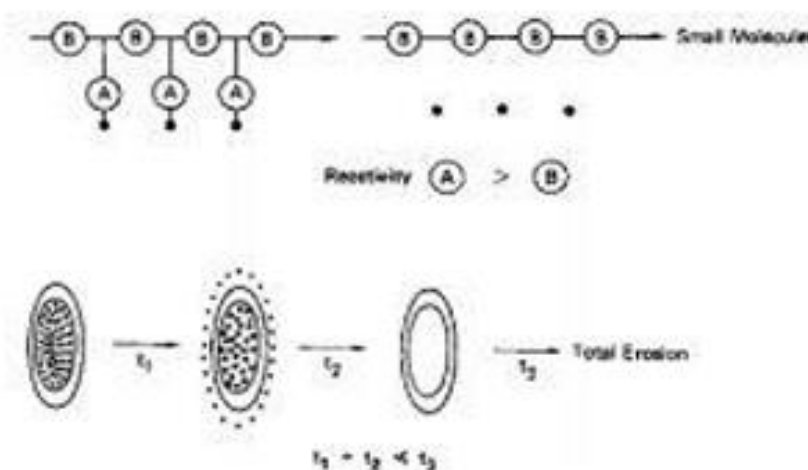


Figure 3: Drug Released Mechanism by Erosion

A bioerodible rate-regulating membrane envelops the active ingredient within a core. This kind of technology combines the benefits of erodibility—which causes bioerosion and eliminates requirement to surgically take out a device of the drug-depleted device with the characteristics such rate-controlling polymers layer guarantees consistent release of medications through a reservoir-type apparatus. The significant breakdown of the polymer must happen only after drug delivery is complete because the membrane needs to remain stable during the release process. remain It must remain unchanged during the delivery process to ensure a consistent drug release.

As a result, after therapy is finished, Polyester granules can stay inside the cells over varying periods as period.

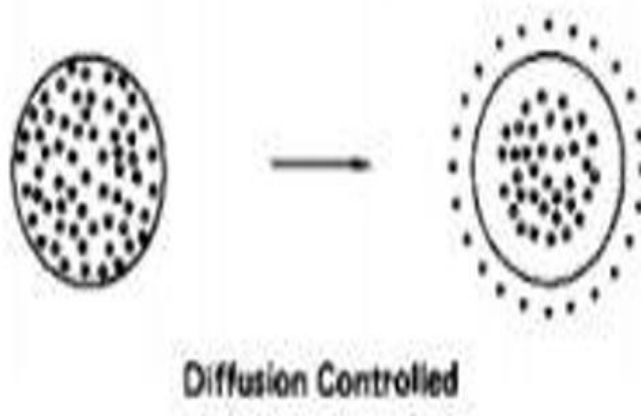


Figure 4: Drug distribution method by dissolution and propagation

The active component is equally distributed in a polymeric material, and the medication evaporates into the bulk via propagation, which consists of propagation and subsidence, or degradation alone. [29]

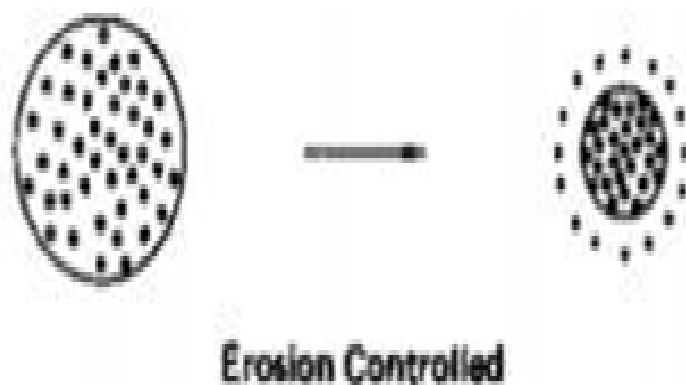


Figure 5: Drug distribution method through dissolution

Applications of Polymer in Pharmaceutical Delivery Systems

Sustained Release

Polymers help maintain a steady release of drugs over time, reducing the frequency of dosing.

Targeted Delivery

Polymers can be engineered to release drugs at specific sites in the body, increasing effectiveness and minimizing side effects.

Controlled Release

Polymers can regulate the timing and rate of Drugs are frequently released in reaction. to physiological changes (e.g., pH or temperature).

Microspheres and Nanoparticles

Polymers can form tiny particles that encapsulate drugs, enhancing solubility, stability, and bioavailability.

Hydrogels

These water-absorbing polymers can provide controlled release of drugs while maintaining moisture, suitable for applications like wound dressings.

Implants

Biodegradable polymer implants can deliver drugs locally over extended periods without the need for removal.

Transdermal Systems

Polymers can be used in patches that deliver drugs through the skin for systemic effects.

Oral Delivery Systems

Polymers can improve the stability and the absorption of medications occurs inside the digestive system.

Gene Delivery

Polymers can facilitate the delivery of genetic material for gene therapy, protecting it until it reaches the target cells.

Vaccines

Polymer-based formulations can enhance the delivery and effectiveness of vaccines by controlling release and targeting immune cells.

These applications leverage the unique properties of polymers to enhance drug efficacy and patient compliance. Researchers worldwide are exploring ways to improve the effectiveness of drugs by altering formulation techniques and using polymeric systems. Polymers specifically designed to address the limitations of traditional dosage forms have proven helpful. While novel polymers offer many advantages, they can also pose risks due to potential toxicity and incompatibilities. Therefore, careful selection of polymers for drug delivery systems is crucial. The main goal is to create biocompatible, multifunctional, low-toxicity, and cost-effective delivery systems.^[30]

CONCLUSION

Researchers worldwide are exploring ways to improve the effectiveness of drugs by altering formulation techniques and using polymeric systems. Polymers specifically designed to address the limitations of traditional dosage forms have proven helpful. While novel polymers offer many advantages, they can also pose risks due to potential toxicity and incompatibilities. Therefore, careful selection of polymers for drug delivery systems is crucial. The main goal is to create biocompatible, multifunctional, low-toxicity, and cost-effective delivery systems.

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