**Chapter 02**

**Polymers in Controlled Release Drug Delivery Systems: An Exploration of Classification, Properties, and Applications**

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**Abstract**

This chapter explores the pivotal role of polymers in the development and application of controlled release medication delivery systems. These systems aim to optimize therapeutic benefits while minimizing side effects by gradually releasing medicinal substances. The chapter delves into the classification of polymers, including natural, synthetic, and semi-synthetic varieties, highlighting their unique characteristics and applications in various drug delivery routes. The versatility of polymers enables the creation of sustained release, biodegradable, targeted, and tunable drug delivery systems. Additionally, the chapter discusses the classification of polymers and their characteristics, emphasizing the importance of safety, biocompatibility, and degradation rates. The extensive applications of polymer-based controlled release systems are explored, covering oral, transdermal, injectable, ocular, and targeted drug delivery. The chapter provides insights into the diverse uses of natural polymers like chitosan and alginate, synthetic polymers such as PLGA and PVA, and semi-synthetic polymers like cellulose derivatives. Furthermore, it compares biodegradable and non-biodegradable polymers, highlighting their eco-friendly aspects. The working mechanism of controlled release systems based on polymers is detailed, emphasizing drug incorporation, matrix or reservoir formation, diffusion or erosion mechanisms, and release profiles. Environmental triggers, biodegradability, targeted delivery, and monitoring/control aspects are also discussed. The importance of controlled drug delivery systems in enhancing patient adherence, providing steady drug delivery, minimizing adverse effects, improving patient comfort, and reducing healthcare expenses is underscored. The chapter concludes by highlighting the role of controlled release technologies in research, innovation, and optimizing pharmacokinetics.

*Keywords: Controlled release, polymers, drug delivery, biodegradability, targeted delivery, sustained release, pharmaceutical technology.*

1. **Introduction**

Pharmaceutical technology has advanced significantly with the introduction of controlled release medication delivery devices. These systems are made to gradually release medicinal substances, such medications, so as to minimize side effects and keep concentrations of the agents within a therapeutic window. Among the most important factors that make controlled release systems successful is the application of polymers. This chapter delves into the crucial function that polymers fulfill in the development and use of controlled release medication delivery systems (1). One key idea that has transformed medicine administration is controlled release of pharmaceuticals. Conventional immediate-release formulations frequently cause the body's medication concentration to fluctuate between peaks and troughs, which can have negative side effects and minimize therapeutic benefits. This chapter explores the complex realm of polymers in controlled release drug delivery systems, including their characteristics, categorization, and wide range of uses. Natural, synthetic, and semi-synthetic polymers are all included in the categorization; each has special qualities that meet distinct medicinal requirements. The mechanical strength, drug-polymer compatibility, biocompatibility, and degradation rates of these polymers are important factors that determine how well the drug delivery system works (2).

Furthermore, a variety of delivery routes, including oral, transdermal, injectable, and ocular, are utilized in the applications of polymer-based controlled release systems. These systems provide flexible ways to reduce the frequency of dose, improve patient compliance, and provide sustained therapeutic benefits. Furthermore, the field delves into targeted medication delivery, which shows how polymers facilitate the accurate administration of medicinal substances to certain bodily locations (3).One of the main areas of pharmaceutical and medical research and development is polymer-based controlled release medication delivery systems. These systems provide a number of benefits, such as increased therapeutic efficacy, less side effects, and greater patient compliance. They do this by controlling the release of medications over an extended period of time using different polymers. An outline of polymers' function in controlled release medication delivery systems is provided below:

1. **Sustained Release:** Polymers are utilized to make reservoirs or matrices for medication delivery. These polymers have the ability to deliver the medication over a lengthy period of time at a regulated pace. Drug release may be customized to meet the unique requirements of the treatment by choosing the right polymer and modifying its characteristics.
2. **Biodegradability:** A large number of polymers utilized in medication delivery systems are biodegradable, which means that the body may gradually break them down. This is beneficial because, after the medicine is discharged, there is no longer a need to surgically (5,6).
3. **Targeted Delivery:** Polymers can be designed to specifically target bodily tissues or cells. This is frequently accomplished by binding to certain receptors with ligands or antibodies, which guarantees that the medication is delivered just where it is required.
4. **Tunable Release Profiles:** Various release profiles may be produced by varying the formulation of the medication, the polymer used, and its molecular weight. For example, according on the therapeutic requirements of the medicine, you can develop systems for pulsatile, sustained, or instantaneous release.
5. **Decreased Administration Frequency**: By lowering the frequency of medication administration, controlled release systems can enhance patient compliance and lessen variations in the body's drug concentrations (10,11).
6. **Reduced adverse Effects:** Controlled release systems can reduce adverse effects related to excessive drug concentrations in non-targeted tissues by regulating the release rate and directing the medication to certain areas.
7. **Protection of Sensitive Drugs:** Certain medications are susceptible to changes in temperature, humidity, or light. These medications may be kept in a protected environment by polymers, which increase their stability and shelf life.
8. **Drug Solubility:** By increasing the solubility and bioavailability of medications that are not very soluble in water, polymers can help enhance the body's capacity to absorb these drugs.
9. **Versatility:** A large range of polymers are accessible, and researchers can select the best polymer based on the particular needs of the medicine and the planned use (4,5).

**Classification of Polymers in Drug Delivery**

1. **Natural polymers**
2. **Synthetic polymers**
3. **Semi-synthetic polymers**
4. **Biodegradable vs. non-biodegradable polymers.**
5. **Natural Polymers:**

These polymers come from living things found in the natural world, such plants, animals, and marine life. Their biocompatibility and biodegradability have led to their usage in a variety of medication delivery methods.

**As an illustration:**

**Chitosan**: Chitosan is made from chitin, which is present in crab exoskeletons. Because of its mucoadhesive qualities, it can be used for both nasal and oral medication administration. Applications for chitosan include transdermal patches, microspheres, and wound dressings.

**Alginate**: Brown seaweed is used to extract alginate. When calcium ions are present, it gels, which makes it helpful for tissue engineering, wound healing, and controlled release systems.

**Collagen**: Found in connective tissues, collagen is a naturally occurring polymer based on proteins. It is employed in cosmetic applications, wound healing, and tissue regeneration medication delivery.

**Biocompatibility:** The body often tolerates natural polymers well, lowering the possibility of negative responses.

**Biodegradability:** They spontaneously disintegrate inside the body, lowering chronic toxicity.

**Mucoadhesion:** A lot of natural polymers stick to mucosal surfaces to distribute drugs in a targeted manner.

**Adaptable Mechanical Properties:** The viscosity, strength, and flexibility of natural polymers differ.

**Applications:**

 Chitosan microspheres are one example of a controlled drug release mechanism.

Tissue engineering with scaffolds made of collagen.

Distribution of drugs topically, such as alginate dressings for wounds (6,7).

1. **Synthetic Polymers:**

 Man-made and providing exact control over their characteristics, synthetic polymers are ideal for a variety of medication delivery applications (8).

**As an illustration:**

The biodegradable polymer known as poly (lactic-co-glycolic acid) (PLGA) is extensively utilized. It is utilized in injectable microspheres and implants and provides regulated medication release over prolonged periods of time.

Polyvinyl alcohol, or PVA, is a water-soluble material that finds use in hydrogels, capsules, and drug delivery films (9,10).

**PEG (polyethylene glycol):** PEG is utilized to increase medicine solubility and lengthen bodily circulation periods because of its hydrophilicity.

**Qualities:**

**Exactness:** It is possible to customize synthetic polymers to get certain medication release characteristics.

**Stability:** They maintain their qualities and have a lengthy shelf life.

Controlled Degradation: A predictable pace of degradation is possible for certain synthetic polymers (11).

**Applications:**

1. Extended duration of medication release, such as PLGA in injectable depots.
2. Increasing solubility, such as by PEGylating medications.
3. Film-forming, such as PVA patches, for transdermal medication administration (12,13).
4. **Semi-Synthetic Polymers:**

These materials combine the advantages of natural and synthetic sources to create a blend of the two.

**As an illustration:**

**Cellulose Derivatives:** Natural cellulose is changed to create cellulose ethers and esters. They are utilized in coatings and tablets with controlled release.

**Starch-Based Polymers:** For oral administration, controlled release techniques employ modified starches.

Combines the qualities of synthetic and natural polymers.

Both biodegradable and biocompatible.

adaptable to a variety of situations (14).

**Applications:**

Oral tablets with modified release, such as those made with cellulose derivatives.

Matrix based on starch for controlled release taken orally.

1. **Comparing Biodegradable vs Non-Biodegradable Polymers.**

Eco-Friendly Polymers: The body may convert these polymers into non-toxic metabolites, which lessens the requirement for elimination. They are frequently used because of their sustainability and safety for medication administration (15,16).

Non-Biodegradable Polymers: The body is unable to quickly break down certain polymers. They may need to be removed after usage; however they are employed when long-term medication release or device integrity is necessary.

**Drug Delivery: The Characteristics of Polymers**

1. Both safety and biocompatibility.
2. Physical characteristics.
3. The rate of degradation.
4. Compatibility of drugs with polymers.
5. Features of erosion and swelling.
6. Kinetics and processes of release (17).



Figure 1: Nanosystem of polymeric particle

**Application of Polymer-Based Systems of Controlled Release**

1. **Oral Drug Delivery:**

**Extended-Release Tablets:** Polymers may be utilized to create oral tablets with a progressive release of medication, which lowers the frequency of dosage and maintains therapeutic levels in the bloodstream (18).

**Oral Capsules:**

Polymeric coatings on controlled release capsules help to adjust medication release, enhancing patient compliance and reducing adverse effects.

**Drug-Loaded Microspheres:**

For oral delivery, sustained release of medication is provided using microspheres encapsulated in polymer matrices (19, 20).

1. **Transdermal drug delivery:**

 **Transdermal Patches:** Made of polymers, transdermal patches gradually release pharmaceuticals via the skin. This method is easy and painless for administering medications such as hormone treatments, nicotine replacement therapy, and painkillers.

**Microneedle Arrays:** Drugs, vaccines, and biologics can be delivered subcutaneously under control using microneedle arrays with polymeric carriers.

1. **PLGA Microspheres for Injectable Medication Delivery:** As injectable long-acting drug depots, poly (lactic-co-glycolic acid) (PLGA) microspheres are widely utilized, especially for hormones, anticancer medications, and contraceptives (21).
2. **Ocular Drug Delivery:**

**Contact Lens-Based Drug Delivery:** Drugs for ocular disorders, such as glaucoma and dry eye, can be released via polymers built into contact lenses, offering long-lasting comfort.

**Ocular Implants:** By introducing biodegradable polymer implants into the eye, medication can be released gradually, minimizing the frequency of eye drops (22).

1. **Targeted Drug Delivery:**

 **Polymer-Based Nanoparticles:** Therapeutic drugs can be incorporated into polymer-based nanoparticles, which can then be engineered for targeted drug delivery. By delivering medications only to tumor locations and reducing systemic exposure, they are utilized in cancer therapy.

**Antibody-Linked Polymer Transporters:** Polymer carriers that have been coupled with antibodies or ligands that attach to certain receptors on target cells can improve the accuracy of medication delivery (23).

1. **Pulmonary medication Delivery:**

**Polymeric Inhalation Formulations:**

To enable regulated release in the lungs, polymers are used in inhalable medication formulations for ailments such as asthma and chronic obstructive pulmonary disease (COPD).

1. **Intravaginal Drug Delivery:**

**Polymeric Rings:** Polymeric intravaginal rings have the ability to distribute medications over a longer period of time, including antivirals and contraceptives.

**Drug Delivery to the Central Nervous System:** Biodegradable Polymer Implants and Wafers: These can be inserted into the brain or spinal cord to deliver medication locally in cases of neurological disorders (24).

1. **Periodontal Drug Delivery:** Polymeric Gels and Films: In periodontal drug delivery systems, polymers help to release antimicrobial medicines under regulated conditions to treat gum disease.
2. **Veterinary and agricultural:** Slow-release fertilizers, animal health products, and pest control are among the applications for polymer-based controlled release systems in veterinary medicine and agriculture.Polymer-based controlled release systems provide a flexible and inventive method of delivering drugs, meeting a range of therapeutic requirements. They can enhance therapeutic efficacy in a variety of medical and pharmaceutical applications by lowering adverse effects, increasing patient compliance, and optimizing medication pharmacokinetics (1, 25).

**The working mechanism of controlled medication delivery systems based on polymers:**

**Drug Incorporation:** The polymer matrix is first filled with the drug of interest. Several techniques, such as solvent casting, physical mixing, or drug encapsulation in the polymer, can be used to accomplish this (5, 6).

**Matrix or Reservoir Formation:** A regulated drug delivery system is created by processing the drug-polymer combination. Various forms, including solid matrices, hydrogels, microspheres, nanoparticles, and coatings coated with drugs, can be employed, contingent on the intended release profile and use (8, 24).

**Diffusion or Erosion:** In controlled delivery systems, medication release happens primarily through two mechanisms:

1. **Diffusion-Controlled Release**: This process uses diffusion to transfer the drug molecules through the polymer matrix. Drug release is driven by the gradient in drug concentration between the interior and outside of the polymer. Creating barriers that regulate the pace of diffusion or employing low-permeability polymers are two ways to accomplish slow, sustained release.
2. **Erosion-Controlled Release:** The polymer erodes or deteriorates progressively over time in erosion-controlled systems. The drug is released when the polymer breaks down and exposes more drug particles or encapsulations to the outside world. By choosing polymers with certain rates of breakdown, one may regulate the erosion rate (15, 16).

**Release Profiles:**

1. **Zero-Order, First-Order, or Alternative:** Depending on the therapeutic needs, the release kinetics can be designed to follow various patterns. For instance:
2. **Zero-Order Release:** Over time, a steady rate of drug release is sustained.
3. **First-Order Release:** Over time, the pace of drug release slows down.
4. **Pulsatile Release:** The medication is delivered at predetermined intervals in distinct bursts.
5. **Biphasic Release:** The release profile might consist of a continuous release phase after an initial burst release (19,20).

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Figure 2: Drug release profile

**Environmental Triggers:** Certain pH, temperature, and/or enzyme variations at the target location are examples of environmental triggers that are included into some controlled drug delivery systems. To produce therapeutic effects that are site- or time-specific, these triggers have the ability to adjust drug release rates (23).

**Biodegradability:** The polymer utilized in the controlled release technique is frequently biodegradable, which means that it progressively transforms into innocuous byproducts. This is especially helpful when it is not desirable to remove the delivery system surgically.

**Targeted Delivery:** The medication delivery system may be made to specifically target one or more bodily tissues or cells. To guarantee that the medication is delivered at the intended location, this might be accomplished by adding ligands or antibodies that identify and bind to certain receptors on the target cells (9, 24).

**Monitoring and Control:** By modifying the polymer's composition, the drug's concentration, and the delivery systems architecture, scientists and medical experts can keep an eye on and regulate the release of pharmaceuticals (22, 25).

**Importance of Controlled Drug Delivery System**

In contemporary medicine, controlled release medication delivery devices are essential and have a big impact on patients and the healthcare sector. Controlled release medication delivery systems are crucial for the following main reasons (29, 30):

1. **Enhanced Patient Adherence**: One of the main advantages of controlled release systems is their ability to streamline medication schedules. Patients frequently struggle to follow intricate dosage regimens. Patients will find it simpler to adhere to their treatment programs when controlled release devices minimize the need for frequent dosage. Better illness management and therapy results result from this.
2. **Steady and Prolonged Drug delivery**: These methods deliver a steady and regulated medicine delivery over a lengthy time. By keeping medication concentrations within the therapeutic window, this lessens the peaks and troughs that come with formulations for quick release. It has fewer adverse effects and improves symptom management.
3. **Fewer adverse Effects**: Compared to traditional immediate-release formulations, controlled release medication delivery methods can lower the chance of adverse effects. Drugs are released gradually to reduce the possibility of overdosing or underdosing, which can result in side effects or ineffective therapy.
4. **Improved Patient Comfort**: Painless and practical medication delivery is provided by some administration methods, such as transdermal patches. Patients who have trouble swallowing medicines or those who need long-term therapy would particularly benefit from this.
5. **Decreased Hospitalization and Healthcare expenses**: Lower hospitalization and healthcare expenses can be attained through improved patient adherence and fewer side effects. Controlled release systems assist stop disease progression, problems, and the need for further treatments by making sure patients take their prescriptions as directed.
6. **Customized medication Release Profiles**: With the help of these systems, medication release rates may be precisely controlled. Pharmaceutical experts are able to create formulations that are tailored to the pharmacokinetics of certain medications, guaranteeing the drug's continued effectiveness over the whole prescribed term(28,29).
7. **Extended-Acting Formulations**: The creation of extended-acting formulations requires these systems. Long-acting controlled release systems can greatly enhance patients' quality of life when used for drugs that need to be taken often, such as HIV or diabetes.
8. **Research and Innovation**: The pharmaceutical industry's innovation is propelled by the advancement and enhancement of controlled release technologies. Drug treatments progress as a result of researchers' constant hunt for novel materials and formulations to improve drug delivery methods.
9. **Targeted Drug Delivery**: By designing these systems to release medications at certain bodily locations, illnesses may be precisely targeted and the amount of medicine that is exposed to healthy tissues is reduced.
10. **Pharmacokinetics optimized**: Targeting certain body areas is possible with controlled release methods. Drugs that must work locally or prevent systemic circulation should pay particular attention to this. For instance, by administering a medication straight to the site of action, they can lessen systemic adverse effects (26, 27, 28).

**Conclusion**

To sum up, the use of polymers in controlled release medication delivery systems is a cutting edge area of pharmaceutical research that is constantly evolving. Customized methods to drug delivery system design are made possible by the wide categorization of polymers, which spans from synthetic to natural, and their distinct features. These systems demonstrate the durability and flexibility of polymers in the pharmaceutical industry. They are distinguished by their targeted distribution, prolonged release, and enhanced patient compliance. Investigating new polymer-based formulations has great potential to improve treatment results, customized medicine, and transform drug delivery as the area develops.

**References**

1. Adepu, S., & Ramakrishna, S. (2021). Controlled Drug Delivery Systems: Current Status and Future Directions. *Molecules*, *26*(19). https://doi.org/10.3390/MOLECULES26195905
2. Heng, P. W. S. (2018). Controlled release drug delivery systems. *Pharmaceutical Development and Technology*, *23*(9), 833. https://doi.org/10.1080/10837450.2018.1534376
3. Begines, B., Ortiz, T., Pérez-Aranda, M., Martínez, G., Merinero, M., Argüelles-Arias, F., & Alcudia, A. (2020). Polymeric Nanoparticles for Drug Delivery: Recent Developments and Future Prospects. *Nanomaterials*, *10*(7), 1–41. https://doi.org/10.3390/NANO10071403
4. Giammona, G., & Craparo, E. F. (2019). Polymer-Based Systems for Controlled Release and Targeting of Drugs. *Polymers*, *11*(12). https://doi.org/10.3390/POLYM11122066
5. Craparo, E. F., Porsio, B., Mauro, N., Giammona, G., & Cavallaro, G. (2015). Polyaspartamide-Polylactide Graft Copolymers with Tunable Properties for the Realization of Fluorescent Nanoparticles for Imaging. *Macromolecular Rapid Communications*, *36*(15), 1409–1415. https://doi.org/10.1002/MARC.201500154
6. Siafaka, P. I., Üstündağ Okur, N., Karavas, E., & Bikiaris, D. N. (2016). Surface modified multifunctional and stimuli responsive nanoparticles for drug targeting: Current status and uses. *International Journal of Molecular Sciences*, *17*(9). https://doi.org/10.3390/IJMS17091440
7. Cavallaro, G., Sardo, C., Craparo, E. F., Porsio, B., & Giammona, G. (2017). Polymeric nanoparticles for siRNA delivery: Production and applications. *International Journal of Pharmaceutics*, *525*(2), 313–333. https://doi.org/10.1016/J.IJPHARM.2017.04.008
8. Sung, Y. K., & Kim, S. W. (2020). Recent advances in polymeric drug delivery systems. *Biomaterials Research*, *24*(1). https://doi.org/10.1186/S40824-020-00190-7
9. Ahmed, T. A., & Aljaeid, B. M. (2016). Preparation, characterization, and potential application of chitosan, chitosan derivatives, and chitosan metal nanoparticles in pharmaceutical drug delivery. *Drug Design, Development and Therapy*, *10*, 483–507. https://doi.org/10.2147/DDDT.S99651
10. Tien, N. D., Lyngstadaas, S. P., Mano, J. F., Blaker, J. J., & Haugen, H. J. (2021). Recent developments in chitosan-based micro/nanofibers for sustainable food packaging, smart textiles, cosmeceuticals, and biomedical applications. *Molecules*, *26*(9). https://doi.org/10.3390/MOLECULES26092683
11. Hoare, T., Zurakowski, D., Langer, R., & Kohane, D. S. (2010). Rheological blends for drug delivery. I. Characterization in vitro. *Journal of Biomedical Materials Research - Part A*, *92*(2), 575–585. https://doi.org/10.1002/JBM.A.32392
12. Macosko, C. W., Guégan, P., Khandpur, A. K., Nakayama, A., Marechal, P., & Inoue, T. (1996). Compatibilizers for melt blending: Premade block copolymers. *Macromolecules*, *29*(17), 5590–5598. https://doi.org/10.1021/MA9602482
13. Macha, I. J., Ben-Nissan, B., Vilchevskaya, E. N., Morozova, A. S., Abali, B. E., Müller, W. H., & Rickert, W. (2019). Drug delivery from polymer-based nanopharmaceuticals-an experimental study complemented by simulations of selected diffusion processes. *Frontiers in Bioengineering and Biotechnology*, *7*(MAR). https://doi.org/10.3389/FBIOE.2019.00037
14. Imre, B., & Pukánszky, B. (2013). Compatibilization in bio-based and biodegradable polymer blends. *European Polymer Journal*, *49*(6), 1215–1233. https://doi.org/10.1016/J.EURPOLYMJ.2013.01.019
15. Kaur, M., Sharma, A., Puri, V., Aggarwal, G., Maman, P., Huanbutta, K., Nagpal, M., & Sangnim, T. (2023). Chitosan-Based Polymer Blends for Drug Delivery Systems. *Polymers*, *15*(9). https://doi.org/10.3390/POLYM15092028
16. Martinho, N., Damgé, C., & Reis, C. P. (2011). Reis, Recent advances in drug delivery systems. *J Biomater Nanobiotechn*, *2*(05), 510–526. https://doi.org/10.4236/jbnb.2011.225062
17. Langer, R., & Peppas, N. A. (2003). Advances in biomaterials, drug delivery, and bionanotechnology. *AICHE J*, *49*(12), 2990–3006. https://doi.org/10.1002/aic.690491202
18. JM Anderson, S. K. (1989). Advances in Drug Delivery Systems (3), Book Review. *J Pharm Sci*, *78*(7), 608–609. https://doi.org/10.1002/jps.2600780723
19. Sung, Y. K., & Kim, S. W. (2020). Recent advances in polymeric drug delivery systems. *Biomaterials Research 2020 24:1*, *24*(1), 1–12. https://doi.org/10.1186/S40824-020-00190-7
20. Sinha, V. R., & Khosla, L. (1998). Bio-absorbable polymers for implantable therapeutic systems, Drug Dev. *Ind Pharm*, *24*(12), 1129–1138. https://doi.org/10.3109/03639049809108572
21. Din, F. U., Aman, W., Ullah, I., Qureshi, O. S., Mustapha, O., Shafique, S., & Zeb, A. (2017). Effective use of nano-carriers as drug delivery systems for the treatment of selected tumors. *Int J Nanomedicine*, *12*, 7291–7309. https://doi.org/10.2147/ijn.s146315
22. Tiwari, G., Tiwari, R., Bannerjee, S., Bhati, L., Pandey, S., Pandey, P., & Sriwastawa, B. (2012). Drug delivery systems: An updated review. *Int J Pharm Investig*, *2*(1), 2–11. https://doi.org/10.4103/2230-973x.96920
23. Heller, A. (2005). Integrated medical feedback systems for drug delivery. *AICHE J*, *51*(4), 1054–1066. https://doi.org/10.1002/aic.10489
24. Begines, B., Ortiz, T., Pérez-Aranda, M., Martínez, G., Merinero, M., Argüelles-Arias, F., & Alcudia, A. (2020). Polymeric Nanoparticles for Drug Delivery: Recent Developments and Future Prospects. *Nanomaterials*, *10*(7), 1–41. https://doi.org/10.3390/NANO10071403
25. Chen, Z. Y., Wang, Y. X., Lin, Y., Zhang, J. S., Yang, F., Zhou, Q. L., & Liao, Y. Y. (2014). Advance of molecular imaging technology and targeted imaging agent in imaging and therapy. *BioMed Research International*, *2014*. https://doi.org/10.1155/2014/819324
26. Jain, K. K. (2020). Role of Nanobiotechnology in Drug Delivery. *Methods in Molecular Biology*, *2059*, 55–73. https://doi.org/10.1007/978-1-4939-9798-5\_2
27. Macedo, A. S., Castro, P. M., Roque, L., Thomé, N. G., Reis, C. P., Pintado, M. E., & Fonte, P. (2020). Novel and revisited approaches in nanoparticle systems for buccal drug delivery. *Journal of Controlled Release*, *320*, 125–141. https://doi.org/10.1016/J.JCONREL.2020.01.006
28. Giri, T. K., Kumar, K., Alexander, A., Ajazuddin, Badwaik, H., & Tripathi, D. K. (2012). A novel and alternative approach to controlled release drug delivery system based on solid dispersion technique. *Bulletin of Faculty of Pharmacy, Cairo University*, *50*(2), 147–159. https://doi.org/10.1016/J.BFOPCU.2012.07.002
29. Liechty, W. B., Kryscio, D. R., Slaughter, B. v., & Peppas, N. A. (2010). Polymers for Drug Delivery Systems. *Annual Review of Chemical and Biomolecular Engineering*, *1*, 149. https://doi.org/10.1146/ANNUREV-CHEMBIOENG-073009-100847
30. *Polymers for Drug Delivery Systems - PMC*. (n.d.). Retrieved January 5, 2024, from https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3438887/