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# "A Review Paper On Nanoparticles For Targeted Drug Delivery"

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### **Abstract:-**

Nanoparticles are tiny carriers designed to deliver drugs directly to affected tissues, improving treatment efficacy while minimizing side effects. Their minute size allows them to cross biological barriers, and their surfaces can be decorated with ligands for precise targeting. Both biodegradable (like PLGA, chitosan) and inorganic materials (such as gold, iron oxide) are being researched for applications in cancer, neurological, and cardiovascular therapies. While promising, challenges like manufacturing consistency, safet§, and regulatory approval remain significant.

**Keywords:-** Nanoparticles, Targeted Drug Delivery, Nanomedicine, Liposomes Polymeric Nanoparticles, Passive Targeting, Active Targeting, Controlled Release, Theranostics, Blood-Brain Barrier

## 1. Introduction:-

Targeted drug delivery aims to direct therapeutic agents exactly where needed, reducing dosage and collateral damage. Nanoparticles or "nano medicines" offer a revolutionary approach, enabling improved treatment precision and reduced systemic toxicity.

Targeted drug delivery is about moving medicine directly to the intended site, like

tumor cells. Nanotechnology is revolutionizing this field by enabling highly precise delivery systems sometimes called nano medicines that aim to reduce dosage and side effects. (1)

# 2. Types of Nanoparticles:-

**Lipid-based systems:** These include liposomes and solid lipid nanoparticles (SLNs), which encapsulate both water-loving and water-repelling drugs.

**Polymeric nanoparticles:** Made of biodegradable polymers like PLGA and PLA, they allow controlled drug release.

- **Metallic nanoparticles**: Gold or iron oxide types, used for drug delivery, imaging, and hyperthermia treatment.
- **Dendrimers**: Tree-like structures that offer multiple places for drug attachment.
- **Liposomes**: These are made of lipids (fats) and can carry both water-loving and water-hating drugs.
- Others: Quantum dots, nanotubes, and nanocapsules are also under research.

# 3. Mechanisms of Targeting:-

- **Passive targeting**: Nanoparticles accumulate in tumors due to leakier blood vessels (EPR effect).
- **Active targeting**: Particle surfaces are tagged with molecules (like antibodies or peptides) that bind to specific cell receptors, improving drug delivery accuracy.
- Stimuli-responsive systems: Designed to release drugs when triggered by conditions like pH changes, heat, or electric signals. (2)

# Agglomeration R group Polar group Shapes Nanoparticles Nanoparticles Nanoparticles Nanoparticles Surface charge Nanoparticles Nanoparticles Nanoparticles Surface charge Nanoparticles Nanoparticles Nanoparticles Liposomes Liposomes Charge Nanoparticles Nanop

Fig no.1 Nanoparticles as a drug delivery system

# 4. Advantages;-

- Direct drug delivery to diseased tissue.
- Less harm to healthy cells
- Possibility for controlled (slow or on-demand) release.
- Facilitates theranostics—using the same particles for treatment and imaging.

# 5. Challenges;-

- Toxicity: Some nanoparticles (especially metals)
   may be harmful depending on their size,
   composition, and surface characteristics.
- Stability and manufacturing: Issues like premature drug release, difficulty in keeping size uniform, and scaling up production.
- **Biodistribution**: Nanoparticles may accumulate in organs like the liver or spleen rather than target tissues.
- Clinical translation and regulation: The actual clinical benefit in humans is often unclear, and standard manufacturing and safety guidelines are lacking. (3)

# 6. Future Perspectives;-

- **Programmable nanoparticles**: Designs like the four-domain model (architecture, interface, payload, dispersal) offer precise control.
- Nanorobots: Tiny, self-driving particles could revolutionize targeted treatment.
- Green synthesis: Eco-friendly methods using plant extracts or microbes may reduce toxicity and production costs.
- **Brain-targeted systems**: Crossing the bloodbrain barrier to treat neurological conditions.
- AI-designed formulations: Machine learning can speed up nanoparticle optimization.
- Nanoparticles are a very promising tool in modern drug delivery. They help carry medicines directly to the place in the body where they are needed, like cancer cells, which makes the treatment more effective and reduces side effects. Because of their tiny size and ability to be modified, they can even cross difficult barriers like the blood-brain barrier.
- Different types of nanoparticles like liposomes, polymeric particles, and

metallic nanoparticles have their own special advantages. Some are already being used in real treatments, and many more are being tested in research and clinical trials.<sup>(4,5)</sup>

## 7. References;-

- Immordino, M. L., Dosio, F., & Cattel, L. (2006). Stealth liposomes: review of the basic science, rationale, and clinical applications, existing and potential. *International Journal of Nanomedicine*, **1**(3), 297–315.
- Shi, J., Xiao, Z., Kamaly, N., & Farokhzad, O. C. (2011).
   Self-assembled targeted nanoparticles: evolution of technologies and bench-to-bedside translation. *Accounts of Chemical Research*, 44(10), 1123–1134.
- Mura, S., Couvreur, P., & Nicolas, J. (2013). Stimuli-responsive nanocarriers for drug delivery. *Nature Materials*, **12**, 991–1003.
- Alexis, F., Pridgen, E., Molnar, L. K., & Farokhzad, O. C. (2008). Factors affecting the clearance and biodistribution of polymeric nanoparticles. *Molecular Pharmaceutics*, **5**(4), 505–515.
- Danhier, F., Ansorena, E., Silva, J. M., Coco, R., Le Breton, A., & Préat, V. (2012). PLGA-based nanoparticles: an overview of biomedical applications. *Journal of Controlled Release*, 161(2), 505–522.
- Kumari, A., Yadav, S. K., & Yadav, S. C. (2010). Biodegradable polymeric nanoparticles based drug delivery systems. *Colloids and Surfaces B: Biointerfaces*, **75**(1), 1–18.

