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Novel Drug Delivery System

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Commentary

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INTRODUCTION

Drug delivery is the process of administering the drug or pharmaceutical product, in order to achieve desired therapeutic effect. The method by which drug delivered is important, as it has significant effect on its efficacy. Novel drug delivery system involves various approaches like medical devices or drug-device combination products. Novel drug delivery system (NDDS) involves combining polymer science, pharmaceutics and molecular biology^[1-3]. Novel drug delivery systems are designed based on physical and biochemical mechanisms. Physical mechanism or controlled drug delivery system includes dissolution, osmosis, erosion and diffusion. Biochemical mechanism includes gene therapy, liposomes, nanoparticles, monoclonal antibodies. NDDS drugs are designed to target the site specific region, in order to achieve desired therapeutic effect, thereby reducing the side or toxic effects.

Novel drug delivery system

Various drug delivery and drug targeted systems are developed, in order to minimize drug degradation, drug adverse effect, and in order to increase the drug bioavailability (amount of drug available at site targeted region). Site specific drug delivery may be either active or passive process.

Liposomes

Liposome serves are lipoidal vesicles (lipid bilayer), which serves as drug carriers for improving the delivery for pharmaceutical drug.

Mechanism: Liposomes are small lipoidal vesicles enclosing aqueous solution inside a hydrophobic membrane, in order to deliver the molecules to targeted site, the lipid bilayer can fuse with other bilayers such as the cell membrane, thus liposomes act as drug carrier for drug delivery^[4-7].

They are various clinical approved liposomal drugs like: liposomal daunorubicin, doxorubicin, Liposomal amphotericin B, Liposomal cytarabine.

Liposomes have following advantages

1. Liposomes are non-toxic, biocompatible, biodegradable, and nonimmunogenic for systemic and non-systemic administrations.

2. The efficacy and therapeutic index of drug Actinomycin can be increased , by formulating it as liposomes.
3. Liposomes has flexibility to bind with site-specific ligands , in order to achieve active targeting.
4. Site-specific targeting of Anti-cancer, Anti-inflammatory drugs.
5. Has high penetration into tissues (Corticosteroids, anesthetics, and insulin).

Nano particles

Nanoparticles are amorphous or crystalline compounds ranging from 10-200 nm, which are used for novel drug delivery system ^[8 - 15]. Nano particles adsorb or encapsulate the drug, thus protecting against chemical or enzymatic degradation ^[16 - 21].

Various Nano devices for drug delivery includes: Nano tubes, quantum dots, Nano robots, dendrimers, Nano wires, Nano shells and Nano pores.

Mechanism: The release of the drug from the formulation is by controlled diffusion or erosion mechanism. Thus the release of the drug occurs from the core, across the polymer matrix or membrane. Thus the membrane acts as a barrier for drug release. Therefore solubility and diffusivity of drug in polymer membrane becomes the determining factor for drug release.

Advantages:

- Nanosomes offer uniform delivery of drug, with greater bioavailability.
- It can be administered through different routes.
- Smaller in size with high surface area.
- Low drug dose is required

Hydrogels

Hydrogels are three -dimensional cross linked water soluble polymers. Various combination of polymers are formulated as novel drug delivery system. Various drugs formulated as hydrogels include Riboflavin, Salicylic acid, Simvastatin ^[22 - 25].

Classification of hydrogels:

- Based on the methods of preparation- Homo-polymeric Hydrogel, Co-polymeric hydrogel, Inter Penetrating Network,
- Stimuli-sensitive hydrogels- Temperature-sensitive hydrogels, pH-sensitive hydrogels, Dual pH-thermal sensitive systems
- Based on mechanism of release-Diffusion controlled, swelling controlled.

Advantages of Hydrogels:

- Biocompatible, biodegradable and can be injected
- Hydrogels possess wide degree of flexibility similar to natural tissue.
- Have good transport properties and easy to modify.

Drug loaded erythrocytes

Drug loaded erythrocytes are novel drug delivery system, in which erythrocytes can be loaded with variety of biologically active substances. The mechanism of drug loading in erythrocytes involves various physical and chemical methods, in which cells are broken down, and the drug is entrapped into erythrocytes. Thus the entrapped drugs are resealed to form resealed erythrocytes. Resealed erythrocytes are used for targeting the drug delivery, and in treatment of parasitic diseases (antimalarial, antiamoebic drugs).

Advantages:

- Biodegradable, biocompatible and non-immunogenic.
- Wide variety of chemicals or drugs can be entrapped.
- Prevents degradation of drug from in activation by endogenous chemicals.

Fast dissolving tablets (FDT)

It is one of novel drug delivery system, in which the dosage form is administered, in absence of water or fluid intake. The administered dosage form dissolves or disintegrates in saliva within 60 seconds. Advantages of fast dissolving tablet include increased patient compliance with ease administration.

Iontophoresis

It is electrochemical process, which involves application of electric current or voltage, thus transport of solute molecules occurs through the skin, by creating potential gradient. Advantages of fast dissolving tablet include increased patient compliance with ease administration.

Phonophoresis

It involves transport of drug through the skin by ultra sound or ultra sonophoresis.

REFERENCES

1. Vijaya Shanti B et al. An Imperative Note on Novel Drug Delivery Systems. J Nanomedic Nanotechnol. 2011;2:125.
2. Agrawal P. Significance of Polymers in Drug Delivery System. J Pharmacovigil. 2015;3:e127.
3. Nikalje AP. Nanotechnology and its Applications in Medicine. Med chem. 2015;5:081-089.
4. Lee JH et al. Magnetically Triggered Drug Release from Liposome Embedded Gel. J Nanomedicine Biotherapeutic Discov. 2014;4:130.
5. Hu D et al. The Bright Future of Liposome Mediated Drug Delivery. Biochem Physiol. 2015;4:e133.
6. Fathalla D et al. Latanoprost Liposomes for Glaucoma Treatment Development and in vitro/in vivo Evaluation of Liposomal Gels for the Sustained Ocular Delivery of Latanoprost. J Clin Exp Ophthalmol. 2015;6:390.
7. Pawar HA and Bhangale BD. Phytosome as a Novel Biomedicine: A Microencapsulated Drug Delivery System. J Bioanal Biomed. 2015;7:006-012.
8. Jigar N Shah et al. Nanoparticulate Transscleral Ocular Drug Delivery. J Biomol Res Ther. 2014;3:116.
9. Hungund BS et al. Comparative Evaluation of Antibacterial Activity of Silver Nanoparticles Biosynthesized Using Fruit Juices. J Nanomed Nanotechnol. 2015;6:271.
10. Nia Y et al. Determination of Ti from TiO₂ Nanoparticles in Biological Materials by Different ICP-MS Instruments: Method Validation and Applications. J Nanomed Nanotechnol. 2015;6:269.

11. Aparna Mani KM et al. Evaluation of In-vitro Anti-Inflammatory Activity of Silver Nanoparticles Synthesised using Piper Nigrum Extract. *J Nanomed Nanotechnol.* 2015;6:268.
12. Le TTD et al. Novel Anti-HER2 ScFv Targeted-Docetaxel Nanoparticles in Therapy of HER2 Overexpressed Cancer. *J Nanomed Nanotechnol.* 2015;6:267.
13. Omprakash and Sharada. Progress in the Synthesis and Surface Modification of Superparamagnetic Iron Oxide Nanoparticles using Silica Nanoparticles. *J Nanomed Nanotechnol.* 2015;6:266.
14. El-Deeb NM et al. Novel Trend in Colon Cancer Therapy Using Silver Nanoparticles Synthesized by Honey Bee. *J Nanomed Nanotechnol.* 2015;6:265.
15. Aftabtalab A and Sadabadi H. Application of Magnetite (Fe₃O₄) Nanoparticles in Hexavalent Chromium Adsorption from Aquatic Solutions. *J Pet Environ Biotechnol.* 2015;6:200.
16. Kakran M et al. Fabrication of Nanoparticles of Silymarin, Hesperetin and Glibenclamide by Evaporative Precipitation of Nanosuspension for Fast Dissolution. *Pharm Anal Acta.* 2015;6:326.
17. Esposito E et al, Physico-Chemical Characterization and Biodistribution Studies of Lipid Nanoparticles. *J Nanomed Nanotechnol.* 2015;6:256.
18. El-Feky GS et al. Utilization of Crosslinked Starch Nanoparticles as a Carrier for Indomethacin and Acyclovir Drugs. *J Nanomed Nanotechnol.* 2015;6:254.
19. Manikandan A and Sathiyabama M. Green Synthesis of Copper-Chitosan Nanoparticles and Study of its Antibacterial Activity. *J Nanomed Nanotechnol.* 2015;5:251.
20. Hussein FH and Shaheed MA. Preparation and Applications of Titanium Dioxide and Zinc Oxide Nanoparticles. *J Environ Anal Chem.* 2015;2:e109.
21. Morris B and Behzad F. The Effects of Gold and Silver Nanoparticles on an Enzymatic Reaction Between Horseradish Peroxidase and 3,3',5,5'-Tetramethylbenzidine. *Biochem Pharmacol.* 2014;3:146.
22. Agostino AD et al. Semiinterpenetrated Hydrogels Composed of PVA and Hyaluronan or Chondroitin Sulphate: Chemico-Physical and Biological Characterization. *J Biotechnol Biomater.* 2012;2:140.
23. Chen Q et al. Hydrogels for Removal of Heavy Metals from Aqueous Solution. *J Environ Anal Toxicol.* 2012;S2:001.
24. Zhu J. Biomimetic Hydrogels as Scaffolds for Tissue Engineering. *J Biochips Tiss Chips.* 2012;2:e119.
25. Venturini M et al. Analysis of Operating Conditions Influencing the Morphology and In vitro Behaviour of Chitosan Coated Liposomes. *J Nanomed Nanotechnol.* 2014;5:211.