

A Review on: Novel Approach In Targeted Drug Delivery System

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Abstract: . It is usually modified to increase the pharmacological and therapeutic character of standard drugs and to reduce problems such as bounded solubility, drug accumulation, poor bio distribution and absence of selectivity, governing drug release transporter and to decrease normal tissue damage present, there are some kinds of DDS using at test phase, such as slow collected discharge drug delivery system, targeted drug delivery systems, transdermal drug delivery system, adherence dosing system and so on. Targeted drug delivery seeks to concentrate the cure in the tissues of benefit while diminishing the relative concentration of the medication in the remaining tissues. This enhance efficacy of the while reducing side effects. It is very hard for a drug molecule to reach its target in the complex cellular network of an organism. The essential benefit of this technique has been the reduction in dose & side effect of the drug. Research related to the development of targeted drug delivery system is now a day is greatly selected and facilitating field of pharmaceutical world. Transdermal devices permit for pharmaceuticals to be delivered across the skin barrier.

Key Words: optics, photonics, light, lasers, templates, journals

1.INTRODUCTION (Size 11, Times New roman)

DDS is refers to the use of new technology and biological polymer information technology to transfer chromosome or drugs to the designated the body parts agree to the clinical necessity time and dose.DDS is the employment and growth in pharmacy by the advanced science and technical knowledge, it has become the theme of the modern pharmacy creation and improvement. Among the presently available delivery systems, which contain liposomes, emulsions, polymeric micelles and micro-particles, carbon nanotubes (CNTs) ^[04] and so on. Targeted drug delivery is a method of delivering treatment to a patient

in a kind that develop the concentration of the medication in some role of the body dependent to others. Targeted drug delivery seeks to concentrate the remedy in the tissues of attract while lowering the relative concentration of the treatment in the continuing tissues. This improves effect of the while decreasing side effects. Drug targeting is the delivery of drugs to receptors or agent or any other special part of the body to which one choose to deliver the drugs completely. The drug's therapeutic index, as calculated by its pharmacological feedback and safeness, relies in the entrance and particular initiation of the drug with its applicant receptor, whilst decreasing its introduction with non –target tissue.

The targeted or site- specific delivery of drugs is an even a very agreeable aim because this provides one of the practically possible ways to develop the therapeutic index of the drugs.^[04]

2.Targeted Drug Delivery System :-

2.1Liposomes :-

Liposomes, mostly nano-liposomes is mainly lipid bilayer membrane. Liposomes is protected and non-toxic, direct alteration asset are more apparent.Mammalian cell membrane, along with escape from endosomes, developing delivery vehicles to transport active protein to their intracellular target site is thus leading to increase protein-based genome editing. used the lipid-mixing organisation to constituent. Liposomes were modern in class to handle bacterial infections in cystic fibrosis with concern to the improved bactericidal activity of entrapped antibiotics discharged through their combination with bacterial membranes.to defeat the acute and developing problem of antibiotic immunity of bacteria to prevalent antibiotics which made it vital to develop new liposome formulations for antibiotics.

2.2Carbon Materials:-

The use of nanoscale substance has attracted considerable attention. Nanoscale materials have been demonstrated to offer a change of medicinal and diagnostic potential. The drug is usually made from natural polymer material and has been fully utilized to treatment course of tumor, diabetes and vascular

disease. Incidence of use of nano preparation technology can supply good development space for medical use and efficiently promote long-term development of nano biotechnology.

2.3 Metallic nanomaterials:-

Metallic nanomaterials as well as gold and silver nanocrystals, and Nano rods. Have been exhibited to induce confined hyper thermal heating through the absorption of occurrence optical radiation and surface Plasmon diversion to deal with the disease. Its importance is low- cost, readily complex nanoparticles and the particles are also hown to have a high thermal strength. This method will conduct to increased permeation for nanoparticles to beyond the vascular endothelium and execute enhanced collection in the tumor.

2.4 Semiconducting nanomaterials :-

Semiconducting nanomaterials have also accepted a considerable amount of recent care for hyper thermal theranostics, their structures commonly grant for substantial penetration of electromagnetic fields throughout the interiorl volume of the atom.

2.5 Iron oxide Nanocrystals:-

The difference between the magnetic particles and their metallic and semiconducting counterparts is the mechanism by which the atom is heated. Nanoscale matter have been demonstrated to extend a kind of curritive and theranostic capabilities.

2.6 Nano micelles:-

Polymeric Nano micelles are created by amphiphilic polymers with definite hydrophobic and hydrophilic division. The polymer self-assemble to form micelles in aqueous solution. The system of drug discharge from Nano micelles is charge on the nature and force of interactions between core-forming polymer and drug molecules, micelle strength.

2.7 Carbon nanotubes:-

the antifungal compound amphotericin B and doxorubicin have been delivered by Centrist benefit is defencive the cargo from absorption or deterioration, or through targeted delivery that can decreased side effects about the Nano toxicology of the CNT and their virtually dangerous effects on the environment . Lower multi-walled CNTs (MWCNTs, i.e., 1 μm) have been reputed to enter the cell membrane more accurately than the extensive CNTs, which can stop their uptake by self-arranging into a loop or wrap shape. Cai et al.. made a academic evaluation of perception route. They exhibit that amphiphilic nanotubes can enter into through synthetic lipid bilayers via an endocytosis pathway.

2.8 Mesoporous silica materials:-

Mesoporous silica nanoparticles (MSNPs) is evenly sized, porous and dispersible nanoparticles applying colloidal chemistry and evaporation-stimulate self-assembly .it usually becoming biocompatibility and low infection and can be used to track bio distribution, cancer cell targeting efficiency, internalization route, cytotoxicity, and the way of analysis. There have been generally vary reports regarding the infection of MSNP and amorphous silica in genere.

3. Recent approaches

3.1Quantam dots:-

A quantum dot is a semiconductor nanostructure that bound the movement of conduction band electrons, valence band holes. (confine couple of conduction band electrons and valence band holes) in all three spatial directions. Quantum dots are especially considerable for optical applications due to their probably high quantum yield. The capacity to tune the mass of quantum dots is favorable for many requisition and it is one of the most hopeful applicants for use in solid-state quantum computation and analysis,drug delivery, Tissue engineering, hydrolysis clarification and also fabrics technologies.^[03]

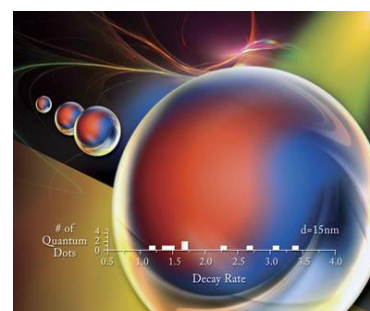


Fig -1: Quantum dots

3.2Transdermal Approach:

Transdermal drug delivery system is locally administered medicaments in the form of patches that deliver drugs for elemental effects at a prearranged and moderate rate. A transdermal drug delivery appliance, which may be of an active or a passive pattern, is a appliance which afford an substitute route for administering medication. These apparatus allow for pharmaceuticals to be delivered over the skin barrier. A drug is devoted in a relatively elevated dosage to the inside of a patch, which is worn on the skin for an elongated span of time. Since there is high compression on the patch and low compression in the blood, the drug will keep diffusing into the blood for a long span of time, maintaining the constant compression of drug in the blood stream.

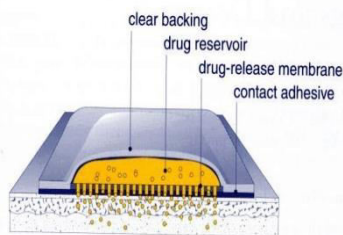


Fig -2: Transdermal Approach

3.3 Folate Targeting:-

Folate targeting is a process applied in biotechnology for drug transmission purposes. Based on the native elevated affinity of folate for the folate receptor protein (FR), which is generally conveyed on the surface of many human cancers, folate-drug conjugates also attach tightly to the FR and trigger cellular intake via endocytosis. The FR is also a well known tumor antisera/ biomarker. Because of this, systematic and corrective methods which abuse the FR's character are being advanced for cancer.

3.4 Brain targeted drug delivery system:-

The brain is a elegant organ, and development create very effective ways to secure it. The delivery of drugs to central nervous system (CNS) is a contest in the treatment of neurologic disturbance. Drugs may be administered straightly into the CNS or administered periodically (e.g., by intravenous injection) for targeted action in the CNS. The main investigate to CNS drug delivery is the blood-brain barrier (BBB).

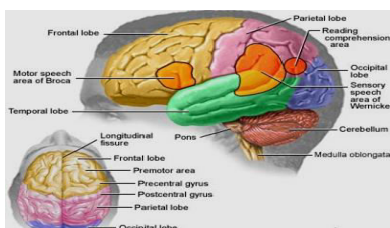


Fig -3: Brain targeted drug delivery system

Different planning that have been used for control the blood-brain barrier for drug delivery to the brain involve osmotic and synthetic opening of the blood-brain barrier as well as the benefit of transportation/carrier systems. Other planning for drug delivery to the brain comprise bypassing the BBB. Various pharmacological agents have been used to open the BBB and direct aggressive methods can establish curative agents into the brain stuff. disparate routes of administration as well as combination of

drugs, e.g., with liposomes and nanoparticles, are considered.

3.5 Liposomes:-

These are cellular coextensive arrangement, range in size from a nanometer to various micrometers, comprise a phospholipids bilayers and are biocompatible, decomposable and non-immunogenic. Liposomes have originated a excellent significance since of their flexibility and have present a suggestive role in composition of efficacious drugs to enhance their therapeutics.

4. Conclusion

Analysis associated to the circumstance of targeted drug delivery system is now a day is greatly superior and facilitating area of pharmaceutical world. It is very hard for a drug molecule to arrive at its purpose in the compound cellular network of structure. Targeted delivery of drugs, as the name recommend, is to further the drug molecule to achieve by preference to the desired location. Generally it may be decide with the vast database of different studies, the science of site particular or targeted delivery of this drug has become wiser. demonstration planning in a clinical now seems potential in near futures.

REFERENCES

- [1] Bennett E. Smith, Paden B. Roder, Xuezhe Zhou, et al. Nanoscale materials for hyperthermal theranostics. *Nanoscale*. 2015, 7(16): 7115–7126.
- [2] Wen S, Liu H, Cai H, et al. Targeted and pH-responsive delivery of doxorubicin to cancer cells using multifunctional dendrimer-modified multi-walled carbon nanotubes. *Adv healthc Mater*. 2013; 2(9):1267–1276.
- [3] R. D.K. Misra, "Quantum dots for tumor-targeted drug delivery and cell imaging," *Nanomedicine*, vol. 3, 2008.
- [4] M. Gupta and V. Sharma, "Targeted drug delivery system: A review," *Research Journal of Chemical Sciences*, vol. 1, 2011.
- [5] K. Rani and S. Paliwal, "A review on targeted drug delivery: Its entire focus on advanced therapeutics and diagnostics," *Scholars Journals of Applied Medical Sciences*, 2014.
- [6] J. Agnihotri, S. Saraf, and A. Khale, "Targeting: new potential carriers for targeted drug delivery system," *International Journal of Pharmaceutical Sciences Review and Research*, vol. 8, 2011.
- [7] R. Singh and J.W. Lillard Jr., "Nanoparticle-based targeted drug delivery," *ExpMolPathol*, vol. 86, 2009.
- [8] A. Swami *et al.*, "Nanoparticles for targeted and temporally controlled drug delivery," in *Multifunctional Nanoparticles for Drug Delivery Applications*, Sonke Svenson, R.K. Prud'homme, Springer US, 2012, Chapter 2
- [9] Aili, D., Mager, M., Roche, D. & Stevens, M. M. Hybrid

- nanoparticle-liposomedetection of phospholipase activity. *Nano Lett* .2011,11:1401–1405.
- [10] Ming Wang, John A. Zuris, Fantao Meng, et al. Efficient delivery of genome-editing proteins using bioreducible lipid nanoparticles *PNAS* .2016, 113(11) : 2868–2873
- [11] Cheng-Xiang Zhang, Wei-Yu Zhao, Lei Liu, et al. A nanostructure of functional targeting epirubicin liposomes dually modified with aminophenyl glucose and cyclic pentapeptide used for brain glioblastoma treatment. *Oncotarget*, 2015,6(32): 32681-32700
- [12] Carolina Garrido, Carrie A Simpson, Noelle P Dahl, et al. Gold nanoparticles to improve HIV drug delivery. *Future Med. Chem.* 2015,7(9), 1097–1107
- [13] Jain RK, Stylianopoulos T. Delivering nanomedicine to solid tumors. *Nature Reviews Clinical Oncology*. 2010; 7(11):653–664.
- [14] Gupta AK, Naregalkar RR, Vaidya VD, et al. Recent advances on surface engineering of magnetic iron oxide nanoparticles and their biomedical applications. *Nanomedicine* 2007; 2(1):23–39.
- [15] Ravi D. Vaishya, Varun Khurana, Sulabh Patel, et al. Controlled Ocular Drug Delivery with Nanomicelles. *Wiley Interdiscip Rev Nanomed Nanobiotechnol*. 2014 September ; 6(5): 422–437.
- [16] Mahdi Karimi, Navid Solati, Amir Ghasemi, et al. Carbon nanotubes part II: a remarkable carrier for drug and gene delivery. *Expert Opin Drug Deliv*. 2015 July ; 12(7): 1089–1105.
- [17] Cai D, Mataraza JM, Qin Z-H, et al. Highly efficient molecular delivery into mammalian cells using carbon nanotube spearing. *Nat Methods*. 2005; 2(6):449–454.
- [18] Pattinson SW, Ranganathan V, Murakami HK, et al. Nitrogen-induced catalyst restructuring for epitaxial growth of multiwalled carbon nanotubes. *ACS Nano*. 2012,6(9):7723-7730.
- [19] Tarn D, Ashley CE, Xue M, et al. Mesoporous Silica Nanoparticle Nanocarriers – Biofunctionality and Biocompatibility *Acc Chem Res*. 2013,19; 46(3): 792–801.
- [20] Lin Y-S, Haynes CL. Impacts of Mesoporous Silica Nanoparticle Size, Pore Ordering, and Pore Integrity on Hemolytic Activity. *Journal of the American Chemical Society*. 2010; 132:4834–4842.
- [21] Nicholas A. W. Bell and Ulrich F. Keyser Specific Protein Detection Using Designed DNA Carriers and Nanopores *J. Am. Chem. Soc.* 2015, 137, 2035-2041.