A Review on Biomedical Applications of Polymeric Nanoparticles

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Abstract

Splendid achievements have been made in management of disease through invention of drugs over past decade. The present conventional drug delivery systems often have side-effects and complications due to their wide distribution throughout the body fluids. The localization of drug action in injured tissue is a promising way to solve this problem. The objective of drug targeting is to achieve a desired pharmacological response at a selected site without undesirable interaction at other sites. This is especially important in cancer chemotherapy and rheumatoid arthritis treatment. Recently invention of drugs has been generated by various drug delivery systems like microspheres, liposomes, noisomes, nanocapsules and nanoparticles. Among all colloidal drug carriers’ nanoparticles are gaining more popularity because of their stability, easy preparation, for achieving reduced toxicity, for increased drug efficacy & site targeted action. Nanoparticulation is a very useful strategy towards targeted drug delivery and also for enhancement of bioavailability of low soluble drugs. In this article nanoparticles preparation methods and their biomedical applications were discussed in detail.

Keywords: Nanoparticles; Polymeric Materials; Bottom Up Approach

Introduction

Nanotechnology has achieved breakthrough in therapeutics, bioengineering, diagnostics, imaging, and optics in recent vintage [1]. The development of nanosystems by tailoring the macromolecules is the recent topic of interest. As nanoparticles possess extraordinary, often tunable properties dramatically different from the bulk materials, such as high surface to volume ratio, particle size and so forth there is an enormous demand for the tailor-made functional nanoparticle systems. Inorganic, organic or hybrid nanoparticulate materials are used in various applications fields as medicine, pharmaceuticals, analytics, catalysis, coating, and several others. Nanoparticles are efficient and versatile devices for drug delivery as they can improve crucial properties of a drug entity such as solubility, pharmacokinetic, biodistribution and in vivo stability [2]. Due to their tailoring properties they can overcome physiological barriers and can help to guide their payload to specific cells or intercellular compartments. By which side effects can be minimized and therapeutic benefits of a drug can be increased. By virtue of their small size and by functionalizing their surface with polymers and appropriate ligands, polymeric nanoparticles can also be targeted to specific cells and locations in the body. Depending on the polymer characteristics, polymeric nanocarriers can also be engineered in such a way that they can be activated by changes in the environmental pH, chemical stimuli, or temperature. Macrophages are well recognized phagocytic cells of the reticuloendothelial system (RES) and one of the main cells responsible for the uptake and clearance of administered drug loaded nanoparticles. In general, once nanoparticles are opsonised, endocytosis/phagocytosis occurs, and the nanoparticles are incorporated in an endolysosome/phagolysosome and degrade [3]. However, the ability of various nanoparticles to escape the endolysosomal compartment allows incorporated drugs to be delivered to the cytoplasm and finally to the nucleus. Thus, this property of the nanoparticles to be easily taken up by phagocytic cells makes them feasible to carry proteins, genes and other biological macromolecules as well [4]. Other applications include cytoplasmic release of plasmid vectors and therapeutic agents (e.g. for cytoplasmic infections and for slow cytoplasmic release of drugs that act on nuclear receptors) [5]. Depending on the preparation methods used, two different types of nanoparticles can be obtained, namely nanospheres and nanocapsules [6,7]. Nanoparticles are drug loaded particles with diameter ranging from 1 to 1000nm.
Necessity of Nanoparticulate Drug Delivery Systems [8-10]

Controlled drug delivery systems are those type of devices in which therapeutic agents may be released at controlled rates for long periods of time, ranging from days to months. The control exercised over the nature of drug delivery may be in temporal nature (rate controlled) or of a spatial nature (site controlled) or both. Currently there are a limited number of formulations approaches available for the compounds that are soluble in water; that includes, solubilisation, cosolvency, complexation with beta-cyclodextrines and solid dispersions that can enhance the dissolution of the drugs. Another classical formulation approach for poorly soluble drugs is micronisation, that means the transfer of coarse drug powder into ultrafine powder. It’s a technology for BCS classified class II drugs that are having a poor solubility but a low bioavailability due to their poor solubility. But for the colonic drug delivery, micronisation often results in a low and variable bioavailability. Hence, the next step taken to improve the saturation solubility, dissolution velocity and bioavailability of drugs is by reducing the particle size from microns to nano size levels that can be termed as Nanonisation. Hence, it was thought that nanoparticle could be used as an ideal drug delivery.

The major goals in designing nanoparticles as a delivery system are to control particle size, surface properties and release of pharmaceutically active agents in order to achieve the site-specific action of the drug at the therapeutically optimal rate and dose regimen. Though liposomes have been used as potential carriers with unique advantages including protecting drugs from degradation, targeting to site of action and reduction toxicity or side effects, their applications are limited due to inherent problems such as low encapsulation efficiency, rapid leakage of water-soluble drug in the presence of blood components and poor storage stability. On the other hand, polymeric nanoparticles offer some specific advantages over liposomes. For instance, they help to increase the stability of drugs/proteins and possess useful controlled release properties. Nanoparticles have a relatively large surface which promotes its ability to bind, adsorb and carry other compounds such as drugs and proteins. Although the definition identifies nanoparticles as having dimensions below 0.1 micro meter and 100nm, especially in the area of the drug delivery relatively large sized nanoparticles (greater than 100nm) may be needed for loading sufficient amount of drug onto the particles.

Method of Synthesis

Nanomaterials exhibit unique properties at nanoscale of 1 to 100 nanometer (nm). However, achieving sizes <100nm is more feasible with hard materials (like Si/ ca, Metal oxides and diamonds) compared to drug and polymer molecules, which are soft materials. Production of nanoparticles for drugs that are usually soft materials with melting point below 300°C is much more challenging than that of hard materials because of high stickiness of the former. For this reason, it is a reasonable goal to aim at <300nm for drug and polymer materials. Hence two basic approaches are employed for the synthesis of nanostructures in the 50-300nm range for drug delivery, irrespective of the field or discipline. The two approaches are ‘Bottom-Up’ approach and ‘Top-Down’ approach [2].

'Bottom-Up' Approach

The building of nanostructures is achieved by growing or assembling of atoms or molecules which are the building blocks. The building blocks may be manipulated through controlled chemical reactions to self-assemble and make nanostructures such as nanotubes and quantum dots. Atoms or molecules may also be physically manipulated to form nanostructures using minute probes. Self-assemblying of atoms or molecules can be achieved by templating and non-templating. Templating involves the interaction of bio macromolecules under the influence of a specific sequence, pattern, structure, external force or spatial constraint. Non-templating is the formation of nanostructures from atoms or molecules with external influence.

'Top-Down' Approach

Bulk materials are reduced by some processes to form nanostructures. 'Top down' is achieved by breaking or etching techniques which is achieved by bulk machining, surface machining and mold machining employing lithography.

Bio Medical Applications of Nanoparticle [11-13]

Medicine: The biological and medical research communities have exploited the unique properties of nanomaterials for various applications (e.g., contrast agents for cell imaging and therapeutics for treating cancer). Terms such as biomedical nanotechnology, bio nanotechnology, and nanomedicine are used to describe this hybrid field. Functionalities can be added to nanomaterials by interfacing them with biological molecules or structures. The size of nanomaterials is similar to that of most biological molecules and structures; therefore, nanomaterials can be useful for both in vivo and in vitro biomedical research and applications. Thus far, the integration of nanomaterials with biology has led to the development of diagnostic devices, contrast agents, analytical tools, physical therapy applications, and drug-delivery vehicles.

Diagnostics: Magnetic nanoparticles bound to a suitable antibody are used to label specific molecules, structures or microorganisms. Gold nanoparticles tagged with short segments of DNA can be used for detection of genetic sequence in a sample. Multicolor optical coding for biological assays has been achieved by embedding different-sized quantum dots, into polymeric micro beads. Nanopore technology for analysis of nucleic acids converts strings of nucleotides directly into electronic signatures.
Drug Delivery: The overall drug consumption and side-effects can be lowered significantly by depositing the active agent in the morbid region only and in no higher dose than needed. This highly selective approach reduces costs and human suffering. An example can be found in dendrimers and nanoporous materials. They could hold small drug molecules transporting them to the desired location. Some potentially important applications include cancer treatment with iron nanoparticles or gold shells. A targeted or personalized medicine reduces the drug consumption and treatment expenses resulting in an overall societal benefit by reducing the costs to the public health system.

Delivery of Anticancer Drugs: Polyalkyl cyanoacrylate nanoparticles have been studied for targeting drugs to specific sites in the body. The small colloidal carriers are biodegradable and drug substances can be incorporated by a process of surface adsorption.

Eg: Doxorubicin associated with polyisohexylcyanoacrylate nanoparticles, Mitoxantrone in Polybutyricyanoacrylate nanoparticles, aclacinomycinA in polyisobutycyanoacrylate nanoparticles, acyclovir in Polybutylicyanoacrylate nanoparticles and doxorubicin in polyalkyl cyanoacrylate nanoparticles, granulocyte colony stimulating factor in polyalkyl cyanoacrylate nanoparticles.

Nanoparticles Can Be Used for Targeting Inflammatory Bowel Disease: ALF Lanprecht, Nathalie Urban and co-workers in a series of studies advocated nanoparticles for brain delivery. They reported transport of the hex peptide dalargin across the blood brain barrier using poly(butyl cyanoacrylate) nanoparticles which were coated with polysorbate 80. Some neuropeptides are also delivered across blood-brain barrier using nanoparticle technology.

Work Done on The Nanoparticle Drug Delivery System

Hasaan A et al. [14] aimed to prepare anti-glucomatous Dorzolamide hydrochloride-(Dorzo) loaded nanoparticles as a controlled release system. Eudragit RS 100 (RS) and/or RL 100 (RL) were used in formulations by an opportunely adapted Quasi-emulsion solvent diffusion technique. The formulations were evaluated in terms of particle size, zeta potential, drug entrapment, and release profile. All formulations showed tiny particle size varying from 114 to 395 nm for RS and 65 to 277 nm for RL. Positive zeta potential was +19 to +32 mV for RS and +23 to +42 mV for RL formulations. It was demonstrated that increasing polymer concentration lead to increase the percentage of drug entrapped in all batches, to a certain extent (drug: polymer 1:4). Nanoparticles prepared using RL showed lower entrapment efficiency than RS. A prolonged drug release was shown by all the formulations. Increasing the polymer concentration caused a decrease in the release rate. M Bharath et al. [15] prepared and investigated the Valsartan nanoparticles by nanoprecipitation method and the various formulations were prepared by optimizing. The prepared nanoparticles were characterized by FTIR, DSC, SEM, particle size analysis. In vitro diffusion and in vivo studies are been performed. The particle sizes of the prepared nanoparticles were ranging from 175 to 232 nm. From three formulations, F2 formulation showed best release of 60.38 % at the end of 24th h. In vivo studies revealed that in case of free drug, 40.9 mcg/ml drug of maximum dose was

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recovered but in case of nanoparticles the dose recovered in serum was 16.02 mcg/ml after 6hr.

S Tamizharsi et al. [16] aimed to prepare and evaluate polymethacrylic acid nanoparticles containing Lamivudine in different drug to polymer ratio by nanoprecipitation method. SEM indicated that nanoparticles have a discrete spherical structure without aggregation. The average particle size was found to be 121nm. The particle size of the nanoparticles was gradually increased with increase in the proportion of polymethacrylic acid polymer. The drug content of the nanoparticles was increasing on increasing polymer concentration up to a particular concentration. No appreciable difference was observed in the extent of degradation of product during 60 days in which, nanoparticles were stored at various temperatures. The in-vitro release behavior from all the drug loaded batches was found to be zero order and provided sustained release over a period of 24 h. The developed formulation overcome and alleviates the drawbacks and limitations of Lamivudine sustained release formulations and could possibility be advantageous in terms of increased bioavailability of Lamivudine. Anbarasan B et al. [17] aimed to study and optimize the formulation and the In-vitro evaluation of Chloroquine Phosphate loaded Chitosan Nanoparticles. The drug content of Nanoparticles increased on increasing the polymer concentration up to a particular level. Entrapment efficiency of 92.87% was achieved with drug to polymer ratio 1:6. In-vitro release of Chloroquine Phosphate from Chitosan Nanoparticles was 85.13% within 24 h. Good stability was observed at refrigeration condition compared to other temperature conditions during eight weeks of storage.

Paresh N Patel et al. [18] formulated and evaluated chitosan nanoparticles of Tamoxifen citrate for cancer therapy. Nanoparticles of TMX were prepared using chitosan using ionic gelation method. The concentration of the polymers Chitosan was selected based on their investigation it was concluded that the repaglinide loaded CN nanoparticles is an effective carrier for the design of a controlled drug delivery system (Table 1).

Table 1: List of marketed nanodrug delivery systems.

| List of Marketed Nano drugs | Composition          | Company      | Administration | Indication          |
|-----------------------------|----------------------|--------------|----------------|---------------------|
| Abelect                     | Amphotericin B       | Enzon        | Intravenous    | Fungal infection    |
| Epaxel                      | IRIV vaccine         | Berna Biotech | Intramuscular  | Hepatitis A         |
| Estrasorb                   | Micellar Estradiol   | Novavax      | Topical        | Menopausal therapy  |
| Pegasys                     | PEG-a-interferon 2g  | Nektar        | Subcutaneous   | Hepatitis B, Hepatitis C |
| Rapamune                    | Nanocrystalline sirolimus | Elan, Wyeth | Oral           | Pharmaceuticals Immunosuppressant |
| Emend                       | Nanocrystalline aprepitant | Elan, Mercck | Oral           | Antiemetic          |
| Tricor                      | Nanocrystalline fenofibrate | Elan, Abbott | Oral           | Anti-hyperlipidemic |

Conclusion

Nanoparticle drug delivery system was found to be very important drug delivery system for targeting of anti-neoplastic agents, anti-rheumatic agents, vaccines and genes. Scientist can target brain, lungs, eye and arteries more effectively by applying nanotechnology principles. Toxicity caused by these nanoparticles is now becoming a great problem. Measurements should be taken to avoid toxicity generated by the nanoparticles.

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