Available online on 15.7.2017 at <http://ujpr.org>
Universal Journal of Pharmaceutical Research
 An International Peer Reviewed Journal

Open access to Pharmaceutical research
 This is an open access article distributed under the terms of the Creative Commons Attribution-Non Commercial Share Alike 4.0 License which permits unrestricted non commercial use, provided the original work is properly cited

Volume 2, Issue 3, 2017

Open Access

REVIEW ARTICLE

NANOSPONGES: A NEW ERA OF VERSATILE DRUG DELIVERY SYSTEM

Neha Rahi^{1D}, Kapil Kumar^{1D}

Department of Pharmaceutics, Global Institute of Pharmaceutical Education and Research, Jaspur Road, Kashipur, US Nagar, Uttarakhand, India.

ABSTRACT

Nanotechnology is a multi disciplinary science which is getting considerable attention in development of new chemical entities, diagnosis and treatment of several ailments. Nanomedicine technology has developed many drug delivering systems like nanoparticles, nanoemulsions, nanosuspensions, nanosponges etc., which are associated with many advantages including improved bioavailability. Nanosponges are tiny sponges having size of about a virus and can be filled with variety of drugs. This sponge can circulate around the body until interact with specific target site and stick on surface and start releasing drug in a controlled manner. Nanosponge play vital role in targeting drug delivery in a controlled manner. Both lipophilic and hydrophilic drugs are incorporated in nanosponges. The outer surface is typically porous, allowing controlled release of drug. Important characteristic of these sponges is their solubility in aqueous form and suitable for the drugs with poor solubility. This review is focusing on the preparation methods, applications of nanosponges in the field of drug delivery.

Keywords: Nanosponges, nano delivery, targeted site.

Article Info: Received 9 May 2017; Revised 5 June; Accepted 8 July, Available online 15 July 2017

Cite this article-

Rahi N, Kumar K. Nanosponges: a new era of versatile drug delivery system. Universal Journal of Pharmaceutical Research 2017; 2(3): 30-33.

DOI: <http://doi.org/10.22270/ujpr.v2i3.RW4>

Address for Correspondence:

Neha Rahi, Department of Pharmaceutics, Global Institute of Pharmaceutical Education and Research, Jaspur Road, Kashipur, US Nagar, Uttarakhand, India, E-mail: neharahi1245@gmail.com

INTRODUCTION

The drug delivery technology has certainly a new interest for drugs by providing them new life through their therapeutic targets. Targeting drug delivery is the major problem which is being faced by the researchers. Target oriented drug improvements in therapeutic efficacy, reduction in side effects and optimized dosing regimen, shall be the leading trends in the area of therapeutics¹. Targeted drug delivery implies for selective and effective localization of pharmacologically active moiety at preidentified (preselected) target in therapeutic concentration, while restricting its access to non-target normal cellular linings and thus minimizing toxic effects and maximizing therapeutic index of the drug². They provide excellent topical delivery of drugs³. These embraces nanotechnology which is applied to pharmacy as nano materials, diagnosing and focusing right place in the body and controlling release of the drug⁴. Nanosponges are about the size of virus which has been backed by naturally degradable polyester. These tiny sponges can circulate around the body they encounter the specific target site and stick on the surface and began to release the drug in a controlled and predictable manner. Owing to their small size and porous nature they can bind poorly-

soluble drugs within the matrix and improve their bioavailability. These are solid in nature and it can be formulated as oral, parenteral, topical or inhalational dosage forms. For oral administration, nanosponges may be dispersed in a matrix of excipients, diluents, lubricants and anti-caking agents which is suitable for the preparation of tablets or capsules.

Figure 1: Structure of Nanosponge.

For parenteral administration, these can be simply mixed with sterile water, saline or other aqueous solutions. For topical administration, they can be effectively incorporated into topical hydrogel⁵.

Advantages

1. This technology offers entrapment of ingredients and reduces side effects⁶.
2. Improved stability, increased elegance and enhanced formulation flexibility.
3. These formulations are stable over range of pH 1 to 11.
4. These formulations are stable at the temperature up to 130°C.
5. These formulations are compatible with most vehicles and ingredients⁷.
6. These are self sterilizing as their average pore size is 0.25µm where bacteria cannot penetrate.
7. They increase the bioavailability of drug⁸.
8. These modify the release of drug.
9. They increase the solubility of poorly soluble drug.

Disadvantages:

- 1) Nanosponges include only small molecules⁹.
- 2) Depend only upon loading capacities.

Table 1: Chemicals used for the synthesis of nanosponges.

Polymers	Hyper cross linked Polystyrenes, Cyclodextrins and its derivatives like Methyl β-Cyclodextrin, Alkyloxycarbonyl Cyclodextrins, 2-Hydroxy Propyl β-Cyclodextrins and Copolymers like Poly(valerolactone-allylvalerolactone)
Crosslinkers	Diphenyl Carbonate, Diarylcarbonates, Diisocyanates, Pyromellitic anhydride, Carbonyldiimidazoles, Epichloridrine, Glutaraldehyde, Carboxylic acid dianhydrides, 2,2- bis(acrylamido) Acetic acid and Dichloromethane
Apolar solvents	Ethanol, Dimethylacetamide, Dimethyl formamide

METHODS OF PREPARATION

It is one of the important criteria for the formation of product obtained activity in β-cyclodextrin, titanium oxide.

1. Solvent method

The solvent required is mixed with the polymer mainly in a polar aprotic solvent, for example dimethyl formide, dimethyl sulfoxide then add this mixture to cross linker in a exceed quantity, the ratio for cross linker/ molar ratio is preferred as 4 to 16¹⁰. The reaction is completed and solution is allow to cool at room temperature. Finally product is dried under vaccum and grinded in a mechanical mill to obtain homogeneous powder¹¹.

2. Ultrasound assisted synthesis

Nanosponges are obtained by reacting polymer with cross linkers without adding or without using solvent and sonification is maintained. The polymer is mix with a cross linkers in a balanced ratio in a flask. The flask is placed in a molar ratio in an ultrasound bath field with water and temperature maintained at 90 °C, the mixture is sonicated for 5 hrs¹². The product is dried under vaccum at 25°C until its further use is utilized¹³.

3. Loading of drug into nanosponges

Nanosponges obtained should be pretreated to maintain mean particle size blow 500nm. Nanosponges are suspended in water and were sonicated to avoid presence of aggregates and particles and got centrifuged to obtain colloidal fraction, then supernatant is separated and dried sample by freezing by drying¹⁴. Para-crystalline nanosponges revealed different loading capacities when compared to crystalline nanosponges poorly crystalline nanosponges had act drug loading as a mechanical mixture rather than inclusion complex¹⁵.

Factors influence nanosponge formation

1. Type of polymer

Type of polymer used can influence the formation as well as the performance of Nanosponges. For complexation, the cavity size of nanosponge should be suitable to accommodate a drug molecule of particular size¹⁶.

2. Type of drugs

Drug molecules to be complexed with nanosponges should have certain characteristics mentioned below¹⁷

- Molecular weight between 100 and 400
- Drug molecule consists of less than five condensed rings
- Solubility in water is less than 10mg/ml
- Melting point of the substance is below 250°C

3. Temperature

Temperature changes can affect drug/nanosponge complexation. In general, increasing in the temperature decreases the magnitude of the apparent stability constant of the drug/ nanosponge complex may be due to a result of possible reduction of drug/ nanosponge interaction forces, such as Van-der Waal forces and hydrophobic forces with rise of temperature¹⁸.

Physicochemical characterization of nanosponge

1. Particle size determination

Free-flowing powders with fine aesthetic attributes will possible to obtain by controlling the size of particles during polymerization. Particle size analysis of loaded and unloaded nanosponges will performed by laser light diffractometry or Malvern Zeta sizer. Particles larger than 30 m can impart gritty feeling and hence particles of sizes between 10 and 25 m are preferred to use in final topical formulation¹⁹.

2. Determination of loading efficiency and production yield

The prepared nanosponge loading efficiency is determined by subtracting the un-entrapped drug from the total amount of drug. The drug entrapment efficiency will be determined by separating un-entrapped drug estimated by any suitable method of analysis²⁰. The method used for separation of un-entrapped drug by gel filtration, dialysis and ultra centrifugation²¹.

3. Porosity

Porosity study is performed to check the extent of nanochannels and nanocavities formed. Porosity of nanosponges is assessed with a helium pycnometer, since helium gas is able to penetrate inter- and intra-particular channels of materials. The true volume of material isdetermined by the helium displacement method. Owing to their porous nature, nanosponges

exhibit higher porosity compared to the parent polymer used to fabricate the system²².

4. Swelling and water uptake

For swellable polymers like polyamidoamine nanosponges, water uptake can be determined by soaking the prepared nanosponges in aqueous solvent²³. Swelling and water uptake can be calculated using equations:

$$\% \text{ Swelling} = \frac{\text{Marking of cylinder at a specified time point}}{\text{Initial marking before soaking}} \times 100$$

$$\% \text{ Water uptake} = \frac{\text{Mass of hydrogel after 72 hrs}}{\text{Initial mass of dry polymer}} \times 100$$

5. Resiliency (Viscoelastic properties)

Resiliency of sponges can be modified to produce beadlets that is softer or firmer according to the needs of the final formulation. Increased crosslinking tends to slow down the rate of release²⁴.

6. Zeta Potential

Zeta potential is a measure of surface charge. The surface charge of nanosponges can be determined by using Zeta sizer²⁵.

7. In vitro release studies

Dissolution profile of Nanosponge can be studied by use of the dissolution apparatus usp XXIII with a modified basket consisted of 5m stainless steel mesh. Speed of the rotation is 150 rpm. Samples from the dissolution medium can be analyzed by a suitable analytical method²⁶.

8. Permeation studies

The diffusion studies of the prepared nanosponge can be carrying out in Franz diffusion cell for studying the dissolution release of nanosponge through a cellophane membrane. Nanosponge sample (0.5g) can taken in cellophane membrane and the diffusion studies were carried out at 37±1°C using 250 ml of phosphate buffer (pH 7.4) as the dissolution medium²⁷.

APPLICATIONS OF NANOSPONGES

1. Nanosponges for drug delivery

Because of their nonporous structure, nanosponges can advantageously carry water insoluble drugs (Biopharmaceutical Classification System class-II drugs). These complexes can be used to increase the dissolution rate, solubility and stability of drugs, to mask unpleasant flavors and to convert liquid substances to solids²⁸.

2. Nanosponges as chemical sensors

Nanosponges which are the type of "metal oxides" act as a chemical sensors which is used in highly sensitive detection of hydrogen using nanosponge titania. Nanosponge structure initially have no point of contact so there is less hinderance to electron transport and it results in higher 3D interconnect nanosponges titania which is sensitive to H₂ gas²⁹.

3. Nanpsponge for oral delivery

In oral application it forms the nanosponge system consist of pores which increase the rate of solubilization of poorly water soluble drugs which get entrapped the drug in pores. The surface area is increased due to nanosize form and increase rate of solubilization³⁰.

4. Solubility enhancement

β-cyclodextrin based nanosponges of itraconazole have enhance solubility of poorly soluble drug. The solubility increased by 50 folds compared to ternary dispersion system e.g. - copolyvidonum³⁰.

5. Nanosponges as a carrier for biocatalysts and release of enzymes, proteins, vaccines and Antibodies

It includes the process applied in industry which correlates with operational condition. Reactions which are not specific give rise to low yields and require high temperatures and pressures which consume large amount of energy and cooling water in down-stream process³¹.

6. Antiviral application

Nanosponges used in nasal, pulmonary route of administration. It provide specificity to deliver antiviral drug on RNA to lungs or nasal route through nanocarriers for targeting virus which may cause infection to RTI such as influenza virus, rhinovirus. Drugs used as nanocarriers are- Zidovudine, Saquinavir³².

7. Cancer

Targeting drug to specific site avoiding the obstacle created by immune system. Different cancer cells had been treated by nanosponges like breast cancer or fast acting glioma type with help of single dose of injections³³.

8. Oxygen Delivery System

Characterized by using α, β and γ cyclodextrins and this are suspended in water and get saturated with water. A silicone form of membrane can also be used for oxygen permeation with the help of nanosponge/hydrogel system³⁰.

CONCLUSION

Nanosponges are tiny mesh-like structures that may revolutionize the treatment of many diseases and this technology is five times more effective at delivering drugs for cancer than conventional methods. Because of their small size and spherical shape nanosponges can be developed as different dosage forms like parenteral, aerosol, topical, tablets and capsules. Nanosponge are nano sized colloidal carrier so they easily penetrate through skin. The nanosponges have the ability to include either lipophilic or hydrophilic drugs and release them in a controlled and predictable manner at the target site. Nanosponge can be effectively incorporated into topical drug delivery system for retention of dosage form on skin. Hence present study concludes that nanosponges may play an important role for the treatment of different diseases.

AUTHOR'S CONTRIBUTION

All authors have worked equally for this work.

ACKNOWLEDGEMENTS

The authors extend their thanks and appreciation to the Global Institute of Pharmaceutical Education and Research, Uttarakhand, India to provide necessary facilities for this work.

CONFLICT OF INTEREST

No conflict of interest associated with this work.

REFERENCES

- Vyas SP, Khar RK. Targeted and controlled drug delivery-novel carrier systems. molecular basis of targeted drug delivery. CBS Publishers and Distributors. New Delhi. 2008; 38- 40.<https://doi.org/10.20959/wjpps20177-9607>
- Nacht S, Kantz M. The Microsponge: A Novel Topical Programmable Delivery System. Chapter 15, In: Topical Drug Delivery Systems 1992; 42: 299-325.
<https://doi.org/10.4103/0110-5558.72416>
- Cavalli R, Trotta F, and Tumiatti W, Cyclodextrin-based nanosponges for drug delivery. J Incl Phenom Macro Chem 2006; 56(1-2):209-213
<https://doi.org/10.1007/s10847-006-9085-2>
- Herbert AL, Martin MR, Gilbert SB, Pharmaceutical dosage forms: Disperse Systems. Marcel dekker, inc. 2005; 2(3): 88-105.
<https://doi.org/10.1002/jps.2600790927>
- Selvamuthukumar S, Anandam S, Kannan K, Manavalan R. Nanosponges: A novel drug delivery system-review. J Pharm Pharmaceut Sci 2012; 15(1): 103 -111.
<https://doi.org/10.18433/J3K308>
- Liang L, De-Pei L, Chih-Chuan L. Optimizing the delivery systems of chimeric RNA DNA Oligonucleotides beyond general oligonucleotide transfer. Eur J Biochem 2002; 269: 5753-5758
<https://doi.org/10.1046/j.1432-1033.2002.03299.x>
- Aritomi H, Yamasaki Y, Yamada K, Honda, Khoshi M, Development of sustained release formulation of chlorpheniramine maleate using powder coated microsponges prepared by dry impact blending method. J Pharma Sci Tech 1996; 56(1): 49-56
<https://doi.org/10.1208/s12249-011-9663-5>
- Yurtdas G, Demirel M, Genc L. Inclusion complexes of fluconazole with beta-cyclodextrin: physicochemical characterization and *in vitro* evaluation of its formulation. J Incl Phenom Macrocycl Chem 2011; 70: 429-435;
<https://doi.org/10.1007/s10847-010-9908-z>
- Zuruzi S, MacDonald NC, Moskovits M, Kolmakov A. Metal oxide nanosponges as chemical sensors: Highly sensitive detection of hydrogen using nanosponge titania; Angewandte Chemie 2007; 46 (23): 4298-4301.
<https://doi.org/10.1002/anie.200700006>
- Swaminathan S, Vavia PR, Trotta F. Formulation of beta cyclodextrins based nanosponges ofitraconazole. J Incl Phenom Macro Chem 2007; 57:89-94.
<https://doi.org/10.1007/s10847-006-9216-9>
- Wong VN, Fernando G, Wagner AR, Zhang J, Kinsel GR, Zauscher S, Dyer DJ. Separation of peptides with polyionic nanosponges for MALDIM Sanalysis. Langmuir 2009; 25(3):1459-65.
- Cavalli R, Akhter AK, Bisazza A, Giustetto P, Trotta F, Vavia P. Nanosponge formulations as oxygen delivery systems. Int J Pharm 2010; 402(1-2): 254-257.
- Ansari KA, Torne S, Vavia PR, Trotta F, Cavalli R. Cyclodextrin based nanosponges for delivery of resveratrol: *In vitro* characterization, stability, cytotoxicity and permeation study. AAPS Pharm Sci Tech 2011; 12: (1), 279-86.<https://doi.org/10.1208/s12249-011-9584-3>
- Yadav G, Panchory H. Nanosponges: a boon to the targeted drug delivery system. J Drug Deliv Therap 2013; 3(4): 151-155.<http://dx.doi.org/10.22270/jddt.v3i4.564>
- Rosalba M, Roberta C, Roberto F, Chiara D, Piergiorgio P, Leigh E, Li S, Roberto P. Antitumor activity of nanosponge-encapsulated Camptothecin in human prostate tumors. Cancer Res 2011; 71:4431
<https://doi.org/10.1158/1538-7445.AM2011-4431>
- Renuka S, Roderick BW, Kamla P. Evaluation of the kinetics and mechanism of drug release from Econazole Nitrate nanosponge loaded carbapol hydrogel. Ind J Pharm Edu 2011; 45(1): 25-31.
- Renuka S, Kamla P. Polymeric nanosponges as an alternative carrier for improved retention of econazole nitrate onto the skin through topical hydrogel formulation Pharm Dev Technol 2011; 16(4):367-376.
<https://doi.org/10.3109/10837451003739289>
- Mamotra R, Bharti N, Bhandari N. Nanosponges as a potential carrier in novel drug delivery system. World J Pharm Sci 2016; 5 (6): 415-424.
<https://doi.org/10.18433/J3K308>
- Wong VN, Fernando G, Wagner AR, Zhang J, Kinsel GR, Zauscher S, Dyer DJ. Separation of peptides with polyionic nanosponges for MALDIMS analysis. Langmuir 2009; 25(3):1459-65.<https://doi.org/10.1021/la802723r>
- Cavalli R, Akhter AK, Bisazza A, Giustetto P, Trotta F, Vavia P. Nanosponge formulations as oxygen delivery systems. Int J Pharm 2010; 402(1-2): 254-257
<https://doi.org/10.22270/jddt.v6i3.1196>
- Sharma R, Roderick B, Pathak K. Evaluation of kinetics and mechanism of drug release from Econazole nitrate Nanosponges loaded carbopol hydrogel. Indian J Pharm Edu Res 2011; 45(1): 25-31.
<https://doi.org/10.1081/DDC-120016687>
- Lala R, Thorat A, Gargote C. Current trends in β -cyclodextrin based drug delivery systems. Int J Res Ayur Pharm 2011; 2(5): 1520-1526.
- Patil B S, Mohite SK. Formulation design and development of artesunate nanosponge. Eur J Pharm Med Res 2016; 3(5): 206-211.
- Wester R, Patel R, Natch S, Leyden J, Melendres J, Maibach H. Controlled release of benzoyl peroxide from a porous microsphere polymeric system can reduce topical irritancy. J American Acad Derm 1991; 24:720-726.
[https://doi.org/10.1016/0190-9622\(91\)70109-F](https://doi.org/10.1016/0190-9622(91)70109-F)
- Renuka S, Roderick BW, Kamala P. Evaluation of the kinetics and mechanism of drug release from econazole nitrate nanosponge loaded carbapol hydrogel. Ind J Pharm Edu Res 2011; 45:25-31.
<https://doi.org/10.1081/DDC-120016687>
- Rajeswari C, Alka A, Javed A, Khar RK. Cyclodextrins in drug delivery: an update review. AAPS pharm Sci Tech 2005; 6(2):E329-E357
<https://doi.org/10.1208/pt060243>
- Amber V, Shailendra S, Swarnalatha S. Cyclodextrin based novel drug delivery systems. J Incl Phenom Macrocycl Chem 2008; 62:23-42
<https://doi.org/10.1016/j.carbpol.2007.10.019>
- Shankar S, Linda P, Loredana S, Francesco T, Pradeep V, Dino A, Michele T, Gianpaolo Z, Roberta C. Cyclodextrin-based nanosponges encapsulating camptothecin: Physicochemical characterization, stability and cytotoxicity. Eur J Pharm Biopharm 2010; 74: 193-201. <https://doi.org/10.1016/j.ejpb.2009.11.003>
- Jenny A, Merima P, Alberto F, Francesco T. Role of β -cyclodextrin nanosponges in polypropylene photooxidation, Carbohydrate Polymers. 2011; 86: 127-135.<https://doi.org/10.2174/1385272823666191029115623>
- Ramnik S, Nitin B, Jyotsana M, Horemam SN. Characterization of cyclodextrin inclusion complexes – a review. J Pharm Sci Tech 2010; 2(3): 171-183.
<https://doi.org/10.3390/molecules23051204>
- Amber V, Shailendra S, Swarnalatha S, Cyclodextrin based novel drug delivery systems. J Incl Phenom Macrocycl Chem 2008; 62: 23-42
<https://doi.org/10.1016/j.carbpol.2007.10.019>
- Martin A, Swarbrick J, Cammarrata A. In: physical pharmacy-physical chemical principles in pharmaceutical sciences 1991; 3: 527
- Emanuele AD, Dinarvand R. Preparation, characterization and drug release from thermo responsive microspheres. Int J Pharm 1995; 237-242.