

Nanoparticles and Their Applications – A Review

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Abstract: *To increase public awareness of nanoparticles' many applications. In biological and medical sciences alike, nanoparticle production and the study of their dimensions and characteristics are crucial. Thus, knowledge of nanoparticles and their uses is essential. Particulate delivery systems have drawn a lot of attention from researchers over the past few decades. As a result, the physical methods of using particulate systems, such as nanoparticles, have been used to alter and enhance the pharmacokinetic and pharmacodynamic aspects of numerous kinds of pharmacological molecules. To spread information about nanoparticles and their applications in industry and medicine.*

Keywords: Medication Delivery Systems, Nanoparticles, Awareness, Particulate, and Nanotoxicity.

I. INTRODUCTION

Nanotechnology is related to things that are smaller than a nanometer¹. Cells are the basic unit of life. Yet, these cell components are quite small the main focus of nanotechnology is the creation, characterization, and design of Nano scale particles. Nanoparticles are essentially tiny objects that function as a single entity based on their attributes and modes of travel. Fine and ultrafine particles range in size from 100 to 2500 nm and 1 to 100 nm, respectively. [1-5] Moreover, they may be created to enhance the therapeutic and pharmacological benefits of the drugs. They also have a large surface area, which makes it possible for numerous functional groups to cling to them and bind to tumor cells [6]. They have shown to be a great alternative to chemotherapy and radiation since they can quickly assemble in the tumor's microenvironment. Many Nanos sized particles, including metals, semiconductors, and polymeric ones, have been produced recently and are used in molecular imaging and particulate delivery systems. Chitosan's, [7-11] silica nanoparticles, micelles, and polyethyleneimine liposomes all contribute to drug delivery with the least amount of side effects. They have also been used as cancer prevention tools [12]. Hence, nanotechnology primarily focuses on the creation of synthetic proteins, enzymes, and cells. In this review, we go through nanoparticle synthesis, types, uses, benefits, and drawbacks. [13-15]

II. HYBRIDIZATION OF NANOPARTICLES

They are created either chemically or biologically. Due to the presence of several toxic compounds absorbed on the surface, the chemical manufacturing process was linked to numerous negative effects. Microorganisms, enzymes, fungi, plants, or plant extracts are all used in the biological synthesis process. Materials like proteins, polysaccharides, and synthetic polymers can be used to make them. Furthermore, the extracellular creation of nanoparticles in fungi is caused by the presence of large secretory parts. The size of the nanoparticles, the inherent properties of the drug, its solubility and stability in water, charge, permeability, biodegradability, biocompatibility, toxicity, drug release, and antigenicity of the finished product are just a few of the variables that affect the choice of the matrix material'[16-18]

III. NANOPARTICLE TYPES

Nanoparticles of Silver

Silver ions are reduced by ethanol at 800 to 1000 degrees Celsius under ambient conditions to produce silver nanoparticles. These are the kind of nanoparticles that are most frequently employed. They are employed in the textile industry for water purification and sunscreen creams because of their potent antibacterial effectiveness. Research has

shown that plants like *Azadirachta indica*, *Capsicum annuum*, and *Carica papaya* are capable of producing beneficial amounts of silver nanoparticles through biosynthesis.[19]

Nanoparticles of Gold

Gold nanoparticles are created through a liquid chemical process called chloroauric acid (HAuCl_4) reduction. They are used to detect protein interactions and conduct immunochemical research. They are also employed to identify aminoglycoside drugs like gentamycin, streptomycin, and neomycin, as well as DNA in a fingerprint sample. Gold nanorods can also identify several bacterial types and cancer stem cells.[20]

Nanoparticles of Copper

A novel technique for producing copper nanoparticles under microwave irradiation is the reduction of copper sulphate with hydrazine in ethylene glycol. The concentration of polyvinylpyrrolidone has a significant impact on the size of copper nanoparticles, and higher concentrations result in particles with smaller dimensions. Basically, their size ranges from 1 to 100 nanometers. They can be used with electrochemical and biosensors [21]

IV. NANOPARTICLES APPLICATIONS: IN DRUGS DELIVERY

First, high stability, large carrier capacity, ease of accommodating both hydrophilic and hydrophobic compounds, and multiple modes of administration, including oral application and inhalation, are the most notable benefits of nanoparticles utilized on drug carriers. Several medications cannot survive the first pass of metabolism. This can be overcome by altering the nanoparticles, which also enable regulated sustained drug release from the matrix. These characteristics can increase the drug's absorption and therefore reduce using frequency³⁷. Nanometer sized quantum dots are tiny semiconductor particles. These manmade atoms with distinct electronic states are also known by that name.

They emit light with varying frequencies when exposed to light or electricity. By varying the dots' sizes, shapes, and materials, these frequencies can be changed, resulting in a variety of applications. ZnO Quantum dots are the most sophisticated form of quantum dots technology used in anticancer medication therapy. The key to this method is that the anticancer drugs are packed into the quantum dots, which are then enclosed in biocompatible polymers. One of the key uses of quantum dots technology is in the delivery of medications that target specific tumors. Verdun et al. demonstrated that doxorubicin concentrations in the liver, spleen, and lungs were higher in mice treated with iso hexyl cyanoacrylate Nano spheres integrated with doxorubicin than in animals treated with free doxorubicin⁴¹. The biggest problem with employing nanoparticles to target tumors is that the liver and spleen's mononuclear phagocytic systems hinder the uptake of the particles. The bio distribution and pharmacokinetics of a cyclic doxorubicin-nanoparticle formulation in tumor-bearing mice were used by Bobby et al. to establish this. [23-25]

These nanoparticles have been altered to serve as delivery systems for a variety of therapeutic medications, including drug conjugates with powerful anticancer properties, layered double hydroxides, and liposomal nanoparticles⁴²⁴⁶. Drug administration to the brain and central nervous system is more challenging, but nanoparticles can get around these challenges, increasing the likelihood of successful medication delivery in the brain.[26]

FOR FOOD

Encapsulation and emulsion creation, food contact materials, and sensor development are the main applications of nanotechnology in the food industry. Nano food is a concept that Garber uses to describe the cultivation, production, packing, and processing of food utilizing nanoparticles. Applications of nonfood have been identified by FSAI, including sensory enhancements (flavor, color, and texture modification), increased absorption, targeted delivery of nutrition bioactive compounds, stabilization of active ingredients like nutraceuticals in food sources, packaging and product improvement to increase shelf life, and sensors for food. Antimicrobials and food safety to eliminate harmful microbes. Hybrid nanoparticles called bionanocomposites have improved mechanical, thermal, and gas properties. They help extend the shelf life of food by being used in packaging. As a result, there will be less reliance on plastic for packaging, which is environmentally benign. Zein, a prolamin and a significant portion of corn protein, serves as an illustration.

Zein can be dissolved in ethanol or acetone to create a biodegradable seen film with increased tensile and water barrier properties⁴⁹. Omega3 fatty acids are added to white bread in Australia using Nano capsules. In Asia, work is being done to create nontoxic Nano scale pesticides that will interfere with weed seeds' seed coats and hinder germination. Hence, the use of nanotechnology in food is expanding quickly and touches every link in the food chain. [27-29]

MEDICAL

Nano medicine supports improved illness diagnosis, early detection, and preventative strategies. The development of nanotechnology has made gene sequencing easier, for example, by creating gold nanoparticles. When attached to the short DNA segments, they are also employed to detect genetic sequences. Nanotechnology can be used to replicate or mend damaged tissue. Nanotechnology has the potential to revolutionize prosthetic implantation and organ transplantation. It has been demonstrated that stem cells can be successfully isolated and grouped using magnetic nanoparticles. On the other hand, quantum dots have been employed for molecular imaging, stem cell tracing, etc. By using specially created nanoparticles, stem cell growth and differentiation may be controlled⁵¹.

The Central Nervous System's regeneration and neuroprotection is a further advantage of nanotechnology. One of the most well-known neurodegenerative diseases is Parkinson's disease. A great way to lessen the negative effects of Parkinson's disease treatment on the periphery is to use an intracranial Nano-enabled scaffold device (NESD) for site-specific dopamine delivery to the brain. The activation of signalling cues for controlled axon growth and the use of peptides and peptide nanoparticles as innovative treatments for diverse CNS illnesses are examples of unique approaches.

Also, they can restore injured neurons to full functionality in order to protect the brain and facilitate the passage of chemicals and drugs across the blood brain barrier. Alzheimer sufferers' brains are the primary locations of amyloid beta plaques. Due to their great affinity for these plaques, these nanoparticles may suppress them, thereby alleviating the symptoms of Alzheimer's disease. An infectious disease that is fatal is tuberculosis. Recent advancements in Nano based drug delivery technologies for the encapsulation and release of antiTB drugs⁵² have allowed for more effective and cost-efficient TB pharmacotherapy. Due to their exceptional luster retention and polish ability, Nano filled composite resin materials offer effective wear resistance, toughness, and great aesthetic values on par with operative dentistry. The use of spherical silicon dioxide Nano fillers in active dentistry offers the potential to change the amount of inorganic phase present. These Nano composites have excellent elasticity, low polymerization shrinkage, high bend strength, and extreme hardness. Treatment of oxidative stress, intraocular pressure measurement, therapy of choroid new vessels, avoidance of scar formation following glaucoma surgery, prosthetics, and other uses for nanotechnology in ophthalmology are a few examples. In recent years, nanotechnology dispersed eye ointment (NDEO) has been used to treat severe evaporative dry eye.

According to histological analysis, NDEO seems to have restored the normal corneal and conjunctiva morphology. Using zinc oxide nanoparticles can reduce antibiotic resistance, improving the antibacterial effectiveness of ciprofloxacin against pathogens. This is the result of these Nano particles interacting with the proteins that cause antibiotic resistance by blocking the release of histamine from mast cells into the blood and tissues, the Nano device Bucky balls can alter the immune response. Nano pharmaceuticals reduce the amount of hazardous systemic side effects, which improves patient compliance. They are essential in identifying the failure of conventional treatments that target active molecules at specific sites. Thrombolytic medicines based on nanoparticles have the ability to hasten the clot removing impact. In Nano dentistry, therapeutic options include using nanotechnology to denaturalize teeth, find a long-term cure for hypersensitivity, realign all of the teeth's orthodontics, etc. Nano robots under the supervision of a computer enable the removal of microorganisms that cause dental caries and the repair of tooth defects where decay has lodged. [30-35]

V. NANOTECHNOLOGY LIMITATIONS

The nanoparticles' potential for being undetectable after being released into the environment is a significant disadvantage, which can pose issues if remediation is required. In order to find nanoparticles in the environment, analytical techniques must be developed.

The link between surface area and chemistry and the operation and toxicity of nanoparticles requires sufficient knowledge. Novel nanoparticles also present a danger of exposure during production or use. Thus, thorough risk evaluations must be taken into account. An effective method for recycling and recovery is required when the rare material must be used for the creation of nanoparticles. As a result, more research is needed to close the vast information gap in the field of nontoxicity because doing so will help to enhance risk assessment. [36-41]

VI. CONCLUSION

Recent developments in nanotechnology have transformed physiologically active molecules that are unstable, poorly soluble, and poorly absorbed into potential medications that can be delivered. Information technology, cognitive science, and biotechnology are all undergoing rapid development thanks to nanotechnology's ability to manipulate matter at the tiniest scales. Nanotechnology research studies are important for every element of human life.

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REFERENCES

- [1]. Gurunathan S., Park J.H., Han J.W., Kim J.H. Comparative assessment of the apoptotic potential of silver nanoparticles synthesized by *Bacillus tequilensis* and *Calocybe indica* in MDA-MB-231 human breast cancer cells: Targeting p53 for anticancer therapy. *Int. J. Nanomed.* 2015; 10:4203–4222. doi: 10.2147/IJN.S83953. [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- [2]. Li W.R., Xie X.B., Shi Q.S., Zeng H.Y., Ou-Yang Y.S., Chen Y.B. Antibacterial activity and mechanism of silver nanoparticles on *Escherichia coli*. *Appl. Microbiol. Biotechnol.* 2010; 8:1115–1122. doi: 10.1007/s00253-009-2159-5. [PubMed] [CrossRef] [Google Scholar]
- [3]. Mukherjee P., Ahmad A., Mandal D., Senapati S., Sainkar S.R., Khan M.I., Renu P., Ajaykumar P.V., Alam M., Kumar R., et al. Fungus-mediated synthesis of silver nanoparticles and their immobilization in the mycelial matrix: A novel biological approach to nanoparticle synthesis. *Nano Lett.* 2001; 1:515–519. doi: 10.1021/nl0155274. [CrossRef] [Google Scholar]
- [4]. Chernousova S., Eppe M. Silver as antibacterial agent: Ion, nanoparticle, and metal. *Angew. Chem. Int. Ed.* 2013; 52:1636–1653. doi: 10.1002/anie.201205923. [PubMed] [CrossRef] [Google Scholar]
- [5]. Li C.Y., Zhang Y.J., Wang M., Zhang Y., Chen G., Li L., Wu D., Wang Q. In vivo real-time visualization of tissue blood flow and angiogenesis using Ag2S quantum dots in the NIR-II window. *Biomaterials.* 2014; 35:393–400. doi: 10.1016/j.biomaterials.2013.10.010. [PubMed] [CrossRef] [Google Scholar]
- [6]. Sondi I., Salopek-Sondi B. Silver nanoparticles as antimicrobial agent: A case study on *E. coli* as a model for Gram-negative bacteria. *J. Colloid Interface Sci.* 2004; 275:177–182. doi: 10.1016/j.jcis.2004.02.012. [PubMed] [CrossRef] [Google Scholar]
- [7]. Li L., Hu J., Yang W., Alivisatos A.P. Band gap variation of size- and shape-controlled colloidal CdSe quantum rods. *Nano Lett.* 2001; 1:349–351. doi: 10.1021/nl015559r. [CrossRef] [Google Scholar]
- [8]. Sharma V.K., Yngard R.A., Lin Y. Silver nanoparticles: green synthesis and their antimicrobial activities. *Adv. Colloid Interface.* 2009; 145:83–96. doi: 10.1016/j.cis.2008.09.002. [PubMed] [CrossRef] [Google Scholar]
- [9]. Gurunathan S., Kalishwaralal K., Vaidyanathan R., Venkataraman D., Pandian S.R., Muniyandi J., Hariharan N., Eom S.H. Biosynthesis, purification and characterization of silver nanoparticles using *Escherichia coli*. *Colloids Surf. B Biointerfaces.* 2009; 74:328–335. doi: 10.1016/j.colsurfb.2009.07.048. [PubMed] [CrossRef] [Google Scholar]
- [10]. Lin P.C., Lin S., Wang P.C., Sridhar R. Techniques for physicochemical characterization of nanomaterials. *Biotechnol. Adv.* 2014; 32:711–726. doi: 10.1016/j.biotechadv.2013.11.006. [PMC free article] [PubMed] [CrossRef] [Google Scholar]

- [11]. Pleus R. Nanotechnologies-Guidance on Physicochemical Characterization of Engineered Nanoscale Materials for Toxicologic Assessment. ISO; Geneva, Switzerland: 2012. [Google Scholar]
- [12]. Murdock R.C., Braydich-Stolle L., Schrand A.M., Schlager J.J., Hussain S.M. Characterization of nanomaterial dispersion in solution prior to in vitro exposure using dynamic light scattering technique. *Toxicol. Sci.* 2008; 101:239–253. doi: 10.1093/toxsci/kfm240. [PubMed] [CrossRef] [Google Scholar]
- [13]. Gurunathan S., Han J.W., Kim E.S., Park J.H., Kim J.H. Reduction of graphene oxide by resveratrol: A novel and simple biological method for the synthesis of an effective anticancer nanotherapeutic molecule. *Int. J. Nanomed.* 2015; 10:2951–2969. doi: 10.2147/IJN.S79879. [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- [14]. Dynashwar K. Mhaske, Dinanth D. Patil, Gurumeet C. wadhawa. Antimicrobial activity of methanolic extract from rhizome and roots of Valerianawallichii. *International Journal on Pharmaceutical and Biomedical Research*, 2011; 2(4):107- 111.
- [15]. Patil DD, Mhaske DK, Gurumeet MP, Wadhawa C. Antibacterial and antioxidant, anti-inflammatory study of leaves and bark of Cassia fistula. *Int J Pharm* 2012; 2(1):401-405.
- [16]. G. C. Wadhawa, M. A. Patare, D. D. Patil and D. K. Mhaske, Antibacterial, antioxidant and anti-inflammatory studies of leaves and roots of Anthocephalus kadamba. *Universal Journal of Pharmacy*, 2013.
- [17]. Shubhada S. Nayak, Nitin A. Mirgane, Vitthal S. Shivankar, Kisan B. Pathade, Gurumeet C. Wadhawa, Degradation of the industrial dye using the nanoparticles synthesized from flowers of plant Ceropogia attenuata, *Materials Today: Proceedings*, Volume 37, Part 2, 2021, Pages 2427-2431, ISSN 2214-7853, <https://doi.org/10.1016/j.matpr.2020.08.274>.
- [18]. Harborne JB., In: *Phytochemical Methods* (Chapman and Hall, London); 1973.
- [19]. Schulz H, Baranska M. Identification and quantification of valuable plant substances by IR and Raman spectroscopy. *Vibrational Spectroscopy*. 2007; 43: 13-25.
- [20]. Patil DD, Mhaske DK, Patare MA and Wadhawa GC: Antibacterial and antioxidant, anti-inflammatory study of leaves and root Delonix regia. *International Journal of chemical and pharmaceutical sciences* 2014; 3: 131-5
- [21]. Arun K. Valvi, Shubhada S. Nayak, Vitthal S. Shivankar, Gurumeet C. Wadhawa, Nano catalyst derived from plant Polyzygus Tuberosus: A suitable for efficient and green synthesis of benzimidazole derivatives, *Materials Today: Proceedings*, Volume 50, Part 5, 2022, Pages 1522-1526, ISSN 2214-7853, <https://doi.org/10.1016/j.matpr.2021.09.102>.
- [22]. Middleton, R.W.; Wimberley D. G. J. *Het. Chem.* 1980, 17, 1757.
- [23]. Kumar R, Joshi YC *E-journal of chemistry*, 2007, 4(4):606-610.
- [24]. Sukalovic V., Andric D., Roglic G., Sladjana K.R., Schratteholz A. and Soskis V., *Eur. J. Med. Chem.*, 2005, 40, 481.
- [25]. Hashem Sharghietal *Journal of Heterocyclic Chemistry* Volume 45, Issue 5 September/October 2008 Pages 1293–1298.
- [26]. J S Yadav, etal *Canadian Journal of Chemistry*, 2008, 86, 124-128.
- [27]. Elder field, R. C.; Kreysa, F. J. J. *Am. Chem. Soc.* 1948, 70, 44.
- [28]. B. S.Praveen, Reddy, P.; Sondhi, S. M.; Lown, J. W. *Pharmacol. Ther.* 1999, 84,(1) 1-111.
- [29]. Sluka, J.; Novak, J.; Budesinsky, Z. *Collect. Czech. Chem. Commun.* 1976, 41, 3628.
- [30]. Nitin A. Mirgane, Vitthal S. Shivankar, Sandip B. Kotwal, Gurumeet C. Wadhawa, Maryappa C. Sonawale, Degradation of dyes using biologically synthesized zinc oxide nanoparticles, *Materials Today: Proceedings*, Volume 37, Part 2021, 849-853, ISSN 2214-7853, <https://doi.org/10.1016/j.matpr.2020.06.037>.
- [31]. Patil, D.D., Deshmukh, A.K., and Wadhava, G.C., Diuretic activity of root and bark of erythrina indica lam. *International Journal of Pharmaceutical Sciences and Research*. 2011; 2 (7), 1811-1813
- [32]. Ranjita M., Sarbari A. and Sanjeeb K. S. (2010) *Cancer nanotechnology: application of nanotechnology in cancer therapy*. Volume 15.
- [33]. Wang, X. et al. (2008) Application of nanotechnology in cancer therapy and imaging. *CA Cancer J. Clin.* 58, 97–110.

- [34]. Vasir, J.K. and Labhasetwar, V. (2007) Biodegradable nanoparticles for cytosolic delivery of therapeutics. *Adv. Drug Deliv. Rev.* 59, 718–728.
- [35]. Fassas, A. and Anagnostopoulos, A. (2005) The use of liposomal daunorubicin (DaunoXome) in acute myeloid leukaemia. *Leuk. Lymphoma* 46.
- [36]. Charrois, G.J. and Allen, T.M. (2004) Drug release rate influences the pharmacokinetics, biodistribution, therapeutic activity, and toxicity of pegylated liposomal doxorubicin formulations in murine breast cancer. *Biochim. Biophys. Acta* 1663.
- [37]. Jiang, Y.H. et al. (2005) SPL7013 gel as a topical microbicide for prevention of vaginal transmission of SHIV89.6P in macaques. *AIDS Res. Hum. Retroviruses* 21, 207–213.
- [38]. Rao, K.S. et al. (2009) Targeting anti-HIV drugs to the CNS. *Expert Opin. Drug Deliv.* 6.
- [39]. Rawat, M. et al. (2006) Nanocarriers: promising vehicle for bioactive drugs. *Biol. Pharm. Bull.*
- [40]. Suresh N. & Digvir S. J. (2010), Nanotechnology for the Food and Bioprocessing Industries, *Food Bioprocess Technol* 4:39–47 cleavage of isopropylidene acetals in Tricky substrates. *Tetrahedron Lett.* 2008, 49, 1961–1964. [CrossRef]