

**NANOTECHNOLOGY FOR DELIVERY OF HERBAL DRUGS****Kusum Sharma\*, Ruchi Shukla, Hitesh Jain and D. B. Meshram**

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**\*Corresponding Author****Kusum Sharma**Pioneer Pharmacy Degree  
College, Vadodara-390019,  
Gujarat, India.**ABSTRACT**

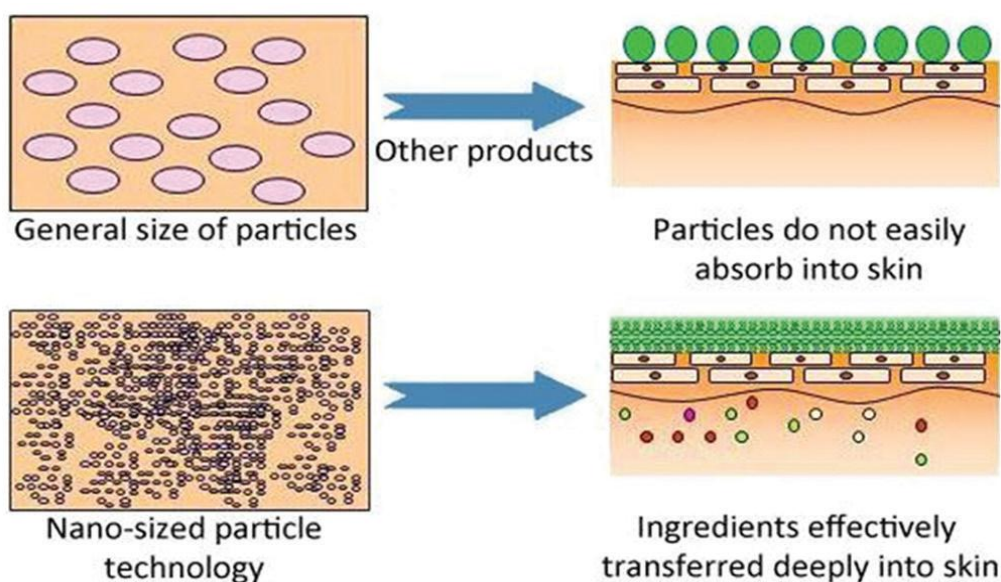
Herbal medicines are widely used in worldwide since ancient time. Now a day nanotechnology is a new prospective in all scientific and technological fields. There are different varieties of new herbal formulations in nanotechnology like nanoparticles, nanocapsules, nanoemulsions etc. The herbal drugs in form of nanoparticles give better effects than the allopathic medicines and can cure disease and reduce the adverse effects of the drugs. It releases drug in sustained manner and reduce the repeated administration to overcome patient compliance and increase the therapeutic effect of drug. Nano-sized drug delivery system enhance the activity of drug. Some extracts, such

as flavonoids, tannins and terpenoids are highly water soluble but poorly lipid soluble which can cause decrease in absorption by nanotechnology it is also overcome. Integration of the nanocarriers as NDDS is the traditional system is essential to conflict more chronic diseases like asthma, diabetes, cancer and other. Nanotechnology system can deliver desire concentration of drug at the specific site of action. Nano formulation is increasing the particle size and increase the surface area due to that bioavailability also increase and it is useful for the treatment, diagnosis, monitoring and control of biological system and have recently been referred to as nanomedicine. Nanotechnology can protect from toxicity, physical and chemical degradation of drug. This review is mainly focus on concept of nanotechnology in herbal drugs.

**KEYWORDS:** Nanotechnology, Novel drug delivery system, Herbal.**INTRODUCTION**

Herbal medicines have been widely used worldwide since ancient times and have been recognized by physicians and patients for their better therapeutic values as they have fewer adverse effects as compared to modern medicines.<sup>[1]</sup> The herbal treatment helps to increase

the therapeutic value by reducing the toxicity and side effects of drugs at the same time it also increases the bioavailability. The applications of nanotechnology for treatment, diagnosis, monitoring, and control of biological systems have recently been refer to as nanomedicine. The nanocarriers have been made of safe materials including synthetic biodegradable polymers, lipids and polysaccharides.<sup>[2]</sup> The known effects and no side effects have made natural products/herbal drugs a powerful therapeutic solution to the organisms. But the delivery of plant/herbal therapeutic molecules as drugs is problematic due to poor solubility, poor permeability, low bioavailability, instability in biological milieu and extensive first pass metabolism.<sup>[3]</sup>



**Fig.1: Transport of drug molecules through skin.**<sup>[4]</sup>

#### Needs of nanotechnology in herbal drug delivery<sup>[5-6]</sup>

- To target the herbal medicine to individual organ which improves the selectivity, drug delivery, effectiveness and safety and there by reduces dose and increase patient compliance.
- To increase the herbal drug solubility and help to localize drug in a specific site thus resulting in better efficacy.
- They are able to deliver high concentrations of drugs to disease sites because of their unique size in high loading capacities.
- Delivering the drug in small particle size enhance the entire surface area of the drugs therefore allocating faster dissolution in the blood.
- Shows enhance permeation and retention effect.

- Decreases the side effects.
- The concentration seems to persist at the sites for the longer periods.

### Types of nanoparticles<sup>[7]</sup>

**Table 1: Inorganic nanoparticles.**

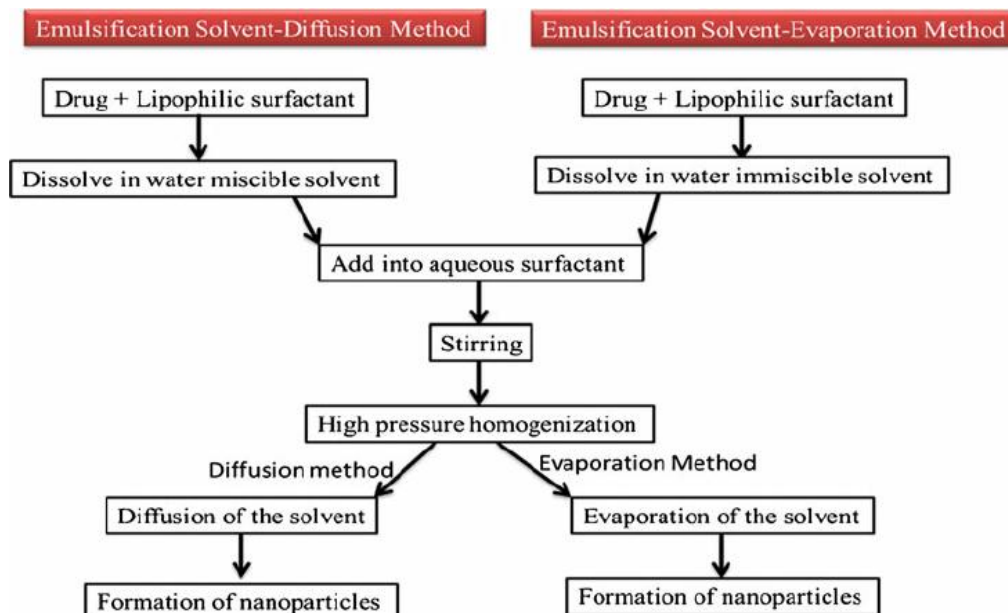
Inorganic compound	Description	Size range (nm)
<b>Metallic</b>	Gold and silver particles	<50
<b>Magnetic</b>	Super paramagnetic iron oxide particles	5–100
<b>Nanoshells</b>	Dielectric silica core in a thin gold metal shell	10–300
<b>Ceramics</b>	Inorganic porous biocompatible materials	<100

**Table 2: Organic nanoparticles.**

Organic compound	Description	Size range(nm)
<b>Carbon tubes</b>	Cylindrical graphite sheets	1.5–5000
<b>Quantum dots</b>	Semiconductor crystals with a cadmium core and metal shell	<10
<b>Dendrimers</b>	Highly branched macromolecules	5–20
<b>Liposomes</b>	Phospholipids	5–100
<b>Polymers</b>	Colloidal particles	10–1000

### Method of preparation

#### 1. High pressure homogenization method<sup>[8]</sup>

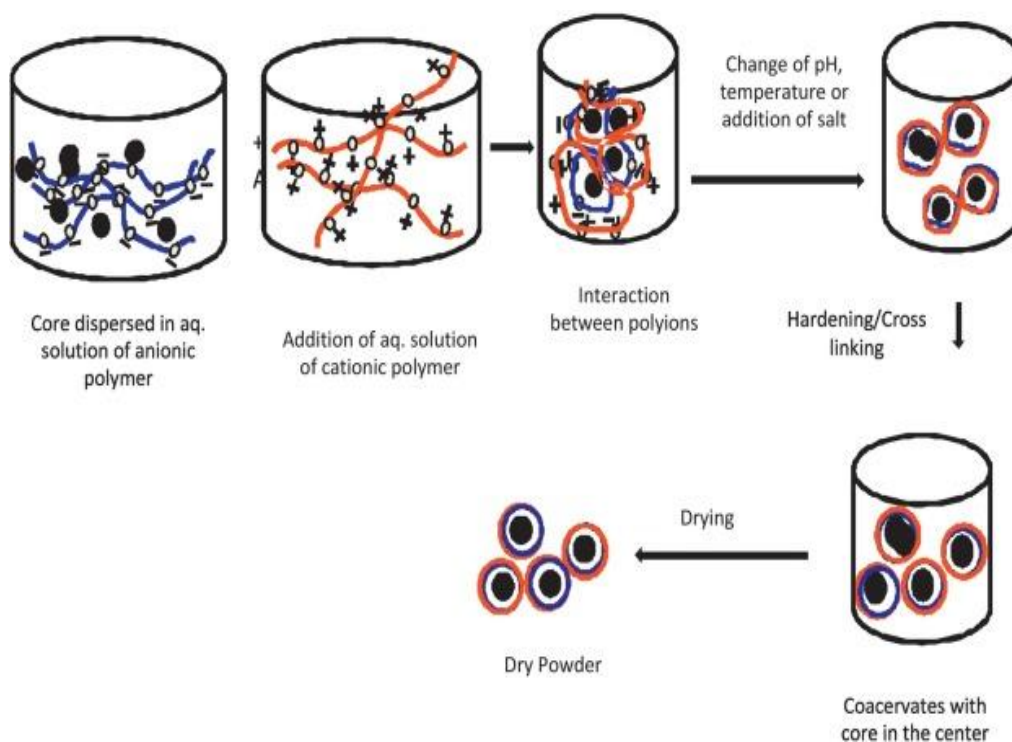


**Fig. 2: High pressure homogenization method.**

#### 2. Complex coacervation method<sup>[9-11]</sup>

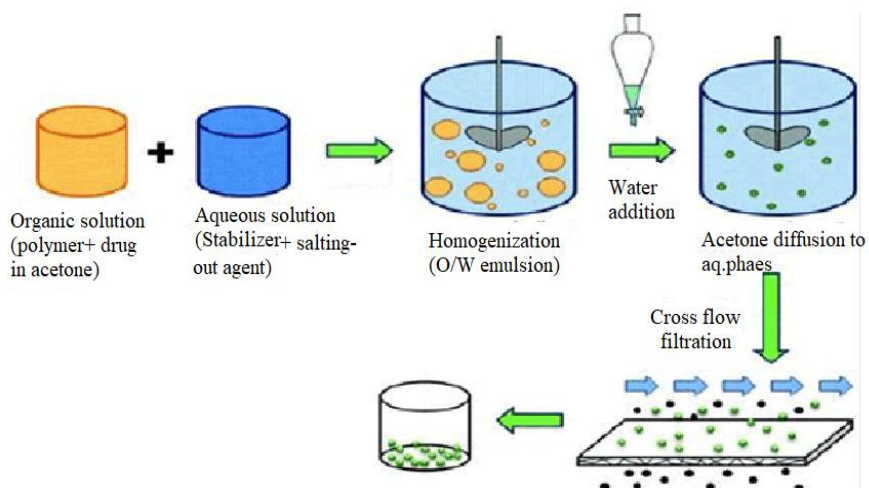
This is a spontaneous phase separation process of two liquid phases in colloidal system, which results by the interaction of two oppositely charge polyelectrolytes upon mixing in an

aqueous solution. This method has been reported to provide good dispersion stability to poorly water-soluble drugs.



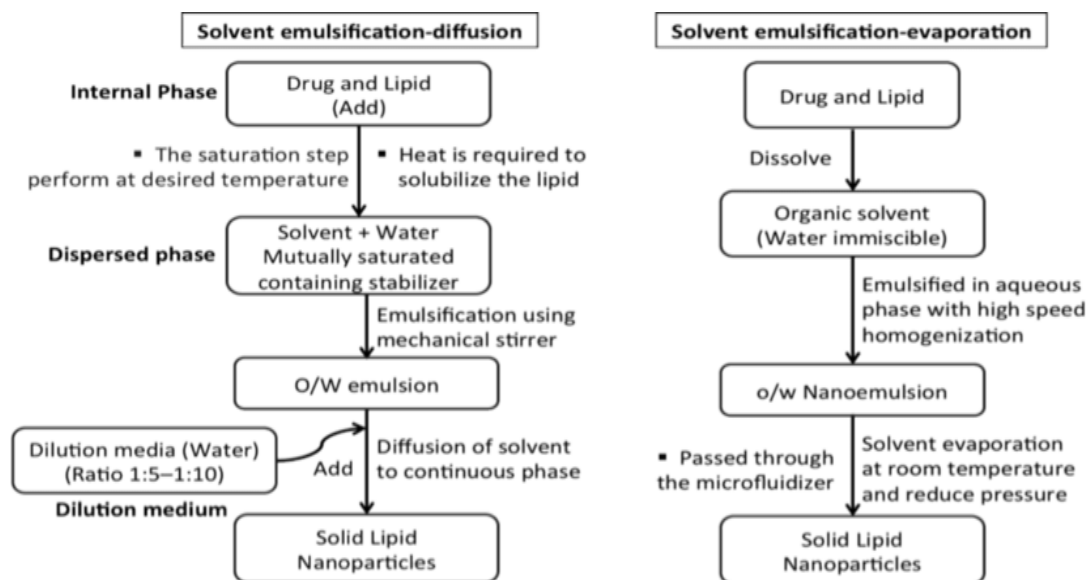
**Fig. 3: Complex coacervation method.**

**3. Salting out method<sup>[12]</sup>**



**Fig. 4: Salting out method.**

#### 4. Solvent emulsification-diffusion method<sup>[13]</sup>



**Fig. 5: Solvent emulsification–diffusion method.**

#### Characterization of synthesized nanoparticles

##### 1. Scanning electron microscope (SEM)

Scanning electron microscopy (SEM) is giving morphological examination with direct visualization. The technique is based on electron microscopy which can be used for morphological and sizing analysis; however, they provide limited information about the size distribution.<sup>[14]</sup>

##### 2. Transmission electron microscope (TEM)

Transmission electron microscope is complex and time consuming because of its requirement that sample should be ultra-thin for the electron transmittance.<sup>[15]</sup>

##### 3. Particle size analyzers

Particle size distribution and morphology of the nanoparticles are most important parameters for characterization of synthesized nanoparticles.<sup>[16]</sup>

##### 4. Dynamic light scattering (DLS)

Dynamic light scattering is widely used to determine the size of Brownian nanoparticles in colloidal suspensions in the range of nano and submicron.<sup>[17]</sup>

##### 5. Atomic force microscopy (AFM)

Atomic force microscopy offers ultra-high resolution in particle size measurement and is based on a physical scanning of samples at sub-micron level using a probe tip of atomic scale.<sup>[18]</sup>



### 6. Surface area analysis

Brunauer Emmett Teller (BET) analysis method is commonly used to determine the total surface area of nanoparticles.<sup>[19]</sup>

### Applications of nanotechnology in delivery of herbal drugs and nutraceuticals<sup>20</sup>

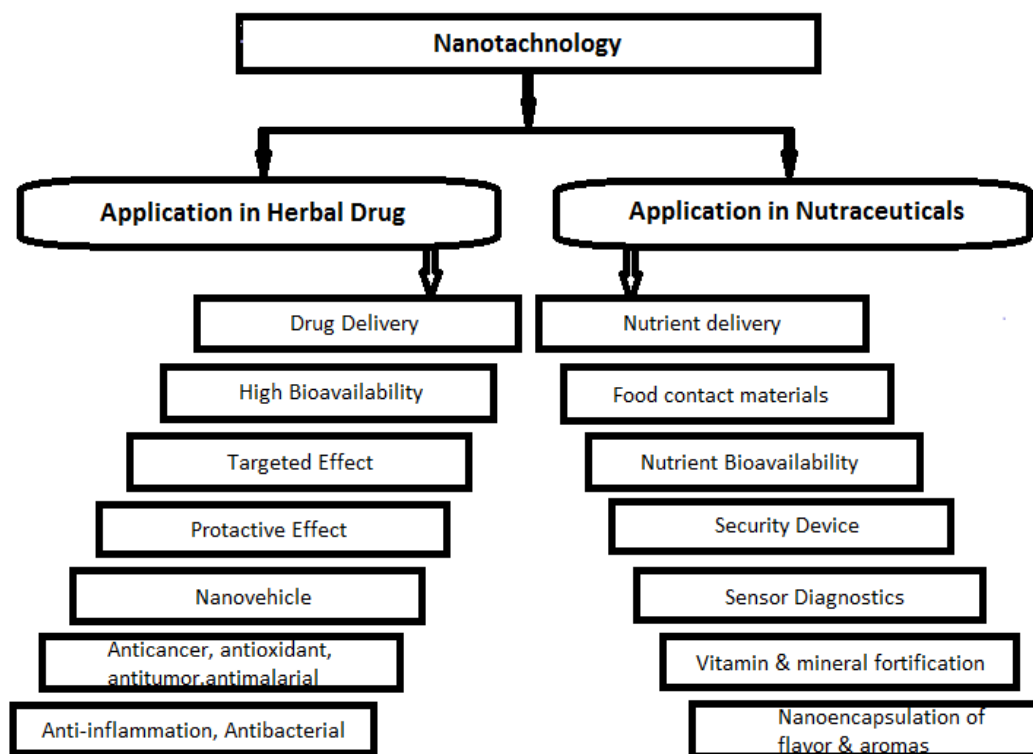


Fig.6: Different applications of herbal nanotechnology.

Examples of natural compounds extracted from higher plants used in nanomedicine aiming different approaches. Some of these extracts are as shown in Fig.7

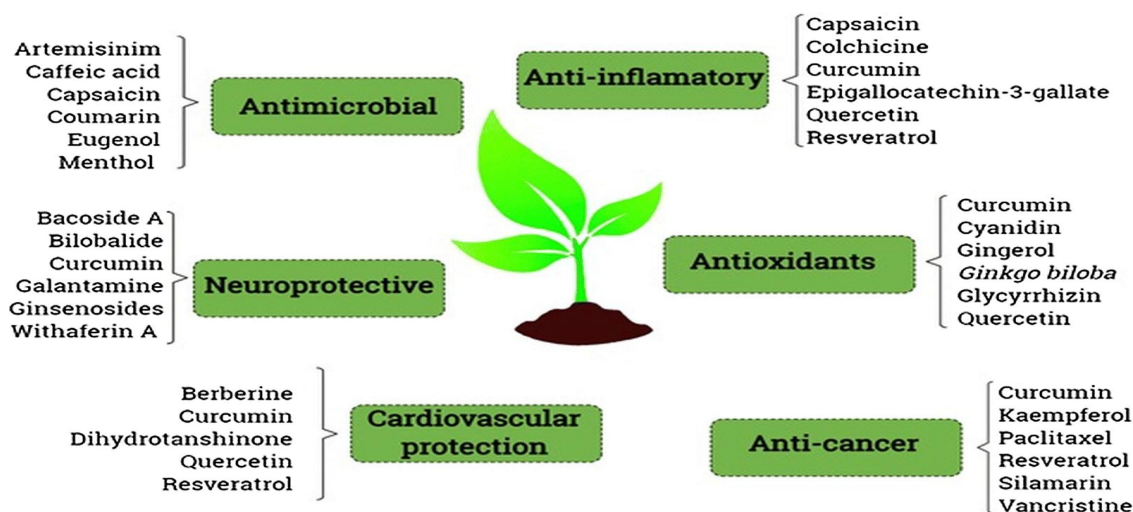


Fig.7: Examples of natural compounds use for different approaches.

**Table 3: Herbal nanoparticles formulations and their use.**<sup>[21-29]</sup>

Name of herb	Category	Uses
Curcumin	Anticancer	Potent anticancer and antitumor
Paclitaxel	Antineoplastic	Acts against several tumour, ovarian and breast cancer
Berberine	Anticancer	Inflammation and several cancers
Camptothecin	Anticancer	Potent anticancer
Ginkgo biloba	Alzheimer's dementia	Acts against loss of memory, thinking, language, behavior.
Triptolide	Anti-arthritis	Inflammatory and autoimmune diseases, especially for rheumatoid arthritis.
Salvia miltiorrhiza	Anti-hyperlipidemia	Cerebrovascular diseases, improve blood stasis.
Quercetin	Anti-oxidant	Potent anticancer
Breviscapine	Anti-cardiovascular	Cerebrovascular and cardiovascular diseases also against pulmonary fibrosis.
Naringenin	Antioxidant, anti-inflammatory	Treating metabolic syndrome, reducing oxidative damage to DNA
Dodder	Anti-inflammatory	Liver problems and spleen problems
Silymarins	Hepatoprotective	Several liver diseases, breast cancer.
Genistein	Antioxidant, Anticancer	Used in cardiovascular diseases, breast and uterine cancer also in osteoporosis.
Annual-mugwort	Antimalarial	In malaria, dysentery, diarrhoea, flu, colds.
Centella asiatica	Anxiolytic	Acts as anti-anxiety, also used in leprosy, cancer, syphilis & allergy.

## CONCLUSION

Overall, this review indicates that nanotechnology has great potential for delivering herbal drugs, and used in various health problems. The combination of nanotechnology with traditional herbal medicine may provide a very useful tool in designing future herbal medicine with improved bioavailability and less adverse effects. Nanotechnology is rapidly expanding and potentially beneficial field with tremendous, implication for industry, medicine, cosmetics.

## REFERENCES

1. Sachan AK, Gupta A. A review on nanoherbal drugs. International journal of pharmaceutical sciences and research, 2015; 6(3): 961-970.
2. Yadav D, Suri S, Hemant K. Novel approach: Herbal remedies and natural products in pharmaceutical science as Nano drug delivery systems. Int J Pharmtech, 2011; 3: 3092-116.
3. Ansari SH, Farha I, Sameem M. Influence of nanotechnology on herbal drug: a review. J Adv Pharm Technol Res., 2012; 3(3): 142-146.

4. Mamillapalli V, Malyada A, Khantamneni P. Nanoparticles for Herbal Extracts. *Asian Journal of Pharmaceutics*, 2016; 10(2): 1-7.
5. Arnlanandraj N, Dhivya S, Gopal V. A review on Herbal nanoparticles. *Pharma Tutor*, 2018; 6(5): 32-37.
6. Aher SS, Chaudhari DB, Saudagar RB. Novel nanotechnology in herbal and medicines. *World Journal of Pharmacy and Pharmaceutical Science*, 2018; 7(2): 515-525.
7. Alakh N. Nanotechnology in herbal medicine and cosmetics. *Int J Res Ayurveda Pharm.*, 2013; 4(3): 472-474.
8. Müller RH, Mäder K, Gohla S. Solid lipid nanoparticles (SLN) for controlled drug delivery – A review of the state of the art. *Eur J Pharm Biopharm*, 2000; 50: 161-77.
9. Fessi H, Puisieux F, Devissaguet JP, Ammoury N. Nanocapsule formation by interfacial Polymer deposition following solvent displacement. *Int J Pharm.*, 1989; 55: 1–4.
10. Sreeraj G, Thomas S, Augustine A. Introduction of nanotechnology in herbal drug and nutraceutical: a review. *Journal of Nanomedicines and Biotherapeutic Discovery*, 2016; 6(2): 2-8.
11. Yen FL, Wu TH, Lin LT. Nanoparticles formulation of *Cuscuta chinensis* prevents acetaminophen-induced hepatotoxicity in rats. *Food Chem Toxicol*, 2008; 46: 1771-7.
12. Tapadiya G, Kale MA, Saboo SS. Impact of nanotechnology on global trade of herbal drug: an overview. *International Journal of Green Pharmacy*, 2017; 11(3): 1
13. Chakraborty K, Shivakumar A, Ramachandra S. Nanotechnology of herbal medicines: a review. *International Journal of Herbal Medicines*, 2016; 4(3): 21-27.
14. Molpeceres J, Aberturas MR, Guzman M. Biodegradable nanoparticles as a delivery system for cyclosporine: preparation and characterization. *Journal of Microencapsulation*, 2000; 17: 599-614.
15. Shi HG, et. al. Characterization of crystalline drug nanoparticles using atomic force microscopy and complementary techniques. *Pharmaceutical Research*, 2003; 20: 479-484.
16. Shekunov BY, Chattopadhyay P, Tong HY, Chow HL. Particle size analysis in pharmaceuticals: principles, methods and applications. *Pharmaceutical Research*, 2007; 24(2): 203-227.
17. Polakovic M, Gorner T, Gref R, Dellacherie E. Lidocaine loaded biodegradable nanospheres-II Modelling of drug release. *Journal of Control Release*, 1999; 60: 169-177.
18. Pangi Z, Beletsi A, Evangelatos K. PEG-ylated Nanoparticles for biological and pharmaceutical application. *Advance Drug Delivery Reviews*, 2003; 24: 403-419. 21.



19. Bisht S, et.al. Polymeric nanoparticles-encapsulated curcumin (nanocurcumin): a novel strategy for human cancer therapy. *Journal of Nanobiotechnology*, 2007; 5(3): 2-18.
20. Jung J, Perrut M. Particle design using supercritical fluids: Literature and patent survey. *J. Supercrit. Fluids*, 2001; 20: 179–219.
21. Singla AK, Garg A, Aggarwal D. Paclitaxel and its formulation. *International Journal of Pharmaceutics*, 2002; 235: 179-192.
22. Fukuda K, Hibiya Y, Mutoh M. Inhibition of activation protein 1 activity by Berberine in human hepatoma cells. *Planta Med.*, 1999; 65: 381-383.
23. Chen KJ, et.al. The therapeutic efficacy of camptothecin-encapsulated super molecular nanoparticle, *Biomaterial*, 2012; 33: 1162-1169.
24. Shinji S, et.al. Analysis of brain cell activation by nano sized particles of Ginkgo biloba extract. *International Journal of Plant Physiology and Biochemistry*, 2011; 3(3): 28-33.
25. Ahmed S, Anuntiyo J, Malemud JC, Haqqi TM. Biological basis for the use of botanicals in osteoarthritis and rheumatoid arthritis: a review. *Evid Based Complement Altern Med.*, 2005; 2: 301-308.
26. Peng Q, et.al. Enhanced the oral bioavailability of saviianolic acid B by phospholipid complex loaded nanoparticles, *Die Pharmazie An International Journal of Pharmaceutical Science*, 2008; 63: 661-666.
27. Usui T. Pharmaceutical prospects of phytoestrogens. *Endocrine Journal*, 2006; 53: 7-20.
28. Raffa V, Vittorio O, Riggio C, Cuschieri A. Progress in nanotechnology for health care, In: *Minimally Invasive Therapy and Allied Technologies*, 2010; 19: 127-135.
29. Yen FY, Wu TH, Lin LT. Naringenin-loaded nanoparticles improve the physicochemical properties and the hepatoprotective effects of naringenin in orally-administered rats with CCl4-induced acute liver failure. *Pharm Res.*, 2009; 26: 893-902.