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Methods and Strategies for the Synthesis of Diverse Nanoparticles and their Applications: A Comprehensive Overview

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ABSTRACT

Ongoing advances in nanotechnology research have established a variety of methods to synthesize nanoparticles (NPs) from a diverse range of materials, including metals, semiconductors, ceramics, metal oxides, polymers, etc. Depending upon their origin and synthesis methods, NPs possess unique physicochemical, structural and morphological characteristics, which are important in wide variety of applications concomitant to electronic, optoelectronic, optical, electrochemical, environment and biomedical fields. This review provides a comprehensive overview on various physical, chemical and bio-assisted methods largely employed to synthesize and fabricate NPs of varying size, surface characteristics, functionalities and physicochemical behavior. The key applications of nanoparticles have also been discussed.

Keywords: Nanoparticles, Synthesis, Nanocomposites, Nanostructures, Nanomaterials, Nanotubes, Nanofibers, Nanobelts
1. Introduction

‘There’s Plenty of Room at the Bottom’ a revolutionary lecture by Richard Feynman in 1959 has provided a new thought process to the scientific community to miniaturize and advance the prevailing technology. Inspired with this vision and substantiated with tremendous research efforts, a new branch of science called nanotechnology got originated. Nanotechnology is basically an amalgamation of two words “Nano” and “Technology” that deals with dimensions and tolerances of less than 100 nanometres, especially the engineering of individual atoms and molecules. With a promise to advance and benefit the society through its wide range of applications in medicines, energy, environment, information technology, aerospace science etc., nanotechnology has emerged as a cutting-edge science in this modern era. Owing to its capability to display improved characteristics at small dimensions, nanomaterial is considered as one of the crucial components of nanotechnology. Enriched with outstanding physical, chemical, biological, optical and electronic properties, nanomaterials helped in improving the existing technologies and also opened new avenues to develop novel scientific/technological fields. Depending upon the desired application, the nanomaterials can be synthesized in diverse shapes and dimensions as described in Table 1.

Table 1 Brief overview on different types of nanostructures, their synthesis methods and related applications

<table>
<thead>
<tr>
<th>Type of Nanomaterials</th>
<th>Synthesis</th>
<th>Applications</th>
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<tbody>
<tr>
<td><strong>Nanocage:</strong> First reported in 2002, this class of nanomaterials has hollow interior and porous walls containing metallic nanoparticles (MNPs) inside. Their size ranges between 10-150 nm.</td>
<td>Nanocages can be synthesized by simple template engaged galvanic replacement reaction among the metallic precursor and the nanotemplates of more reactive metals.</td>
<td>As tracers for tracking using multi-photon luminescence, contrasting agents for photoacoustic and multimodal imaging, photothermal agents for the selective destruction of cancerous tissue and Drug delivery vehicles for smart release in response to external stimuli such as NIR radiation or high-intensity focused ultrasound.</td>
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<td><strong>Nanocrystal:</strong> Nanocrystal is single or multi-phase polycrystalline solids with grain size of less than 100 nm.</td>
<td>Murray et al. have reported high temperature colloidal method to synthesize semiconducting magnetic nanocrystals that are uniform in size to ± one atomic layer, composition, shape, internal structure, and surface chemistry. Li et al. have developed an unified liquid–solid–solution phase transfer synthetic strategy to fabricate monodisperse</td>
<td>Memory devices, solar cells, solid-state displays, photodetectors and field-effect transistor (FET) detectors.</td>
</tr>
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</table>
nanocrystals with different chemistries and properties using versatile materials including noble metal, magnetic/dielectric, semiconducting, rare-earth fluorescent, biomedical, organic optoelectronic semiconducting and conducting polymers NPs. Hyeon et al. have nicely reviewed the synthesis strategies to design monodisperse nanocrystals of metals, semiconductors and metal oxides.\(^\text{12}\)

<table>
<thead>
<tr>
<th>Nanobelt: Nanobelt is thin and flat sheets of ribbon-like structures that are typically 30-300 nm in size.(^\text{15}) Nanobelts with rectangular cross-section and well-defined crystalline facets enable to attain unique optical confinement, microcavity, catalysis and piezoelectric effect.(^\text{16})</th>
</tr>
</thead>
<tbody>
<tr>
<td>Wang et al. have reported the synthesis of 1D ultra-long nanobelts of lanthanum hydroxide using composite-hydroxide-mediated synthesis method.(^\text{16})</td>
</tr>
<tr>
<td>Nanobelts have profound impact in the field of self-powered nanodevices and nanosystems. They have applications in FET devices, nanometer-sized ultrasensitive gas and biosensors, resonators, cantilevers, etc.(^\text{17})</td>
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<tr>
<th>Nanofiber (NF): These are 2D fibre structures having diameter less than 100 nm.</th>
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<tr>
<td>Electrospinning is the most explored method for large scale production of NFs of versatile materials including polymers, metal oxides, carbon based materials, supramolecular dipeptides, etc.(^\text{18-20}) Other method to design NFs are emulsion polymerization, self-assembly, melt blowing and phase separation etc.(^\text{21-23})</td>
</tr>
<tr>
<td>Water filtration systems, surgical implants, biosensors, drug delivery systems, electronic devices, tissue engineering, etc.(^\text{24-26})</td>
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<tr>
<th>Nanoparticle (NP): According to IUPAC, particle of any shape with dimensions in the (1 \times 10^{-9}) and (1 \times 10^{-7}) m range is known as nanoparticle (NP).</th>
</tr>
</thead>
<tbody>
<tr>
<td>Different chemical, physical and biological strategies are available to design NPs of various material types (polymers, inorganic oxides, metals, etc.) of diverse sizes, surface characteristics and topography.(^\text{27}) Our review is an attempt to compile all the synthesis methods for the NP design.</td>
</tr>
<tr>
<td>NPs are having a wide range of applications that includes their role in biomedical devices (biosensors, tissue engineering, drug delivery, bioactuators, bio-imaging devices, etc.), electronics and optoelectronic devices, food industry, construction industries, etc.(^\text{28})</td>
</tr>
<tr>
<td>Nanotube (NT) and Nanorod (NR):</td>
</tr>
<tr>
<td>Nanowire (NW): Nanowire is smart 1D nanostructural material with dimensions of the order of 10^-9 meters.</td>
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<td>Quantum dot (QD): Quantum dots is nanocrystal of semiconducting material, small enough to exhibit quantum mechanical properties and their excitons are confined in all three dimensions. These exhibits strong size dependent optical and electronic properties. QD can contain as few as 100 to 100,000 atoms within the quantum dot volume, with a diameter of 10 to 50 atoms.</td>
</tr>
<tr>
<td>Nanocomposites: It is multiphase material where at least one of the constituent phases has one dimension less than 100 nm. The promise of nanocomposites lies in their multifunctionality, the possibility of realizing unique combinations of properties unachievable with traditional pristine materials.</td>
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Due to its small dimension and exceptional surface properties, NPs have shown great potential for diverse applications. In general, NPs are the most fundamental element of nanomaterials and act as bridging link between the atomic/molecular structures and nanomaterials. The NPs demonstrate size and shape dependent properties, which can be tailored over wide range. Dendrimers, Liposomes, polymer micelles, quantum dots are the other structural analogues of NPs, used for various applications. Depending upon the type of the material, NPs are used for broad spectrum applications including drug delivery, biosensors, bioimaging, molecular tagging, food technology, textile manufacturing, antimicrobial coatings, quantum computers, quantum lasers, energy and environment, etc (Discussed in section 2.4).

To enfold such a broad range of applications, one of the most essential aspects is to understand various possible synthesis methods to design desired NPs. Selection of the synthesis method to realize required NPs largely depends on the desired size, appropriate surface properties and the type of concerned material such as metals, semiconductors, ceramics, polymers, etc. In present article, we understand and review the key synthesis methods for the preparation of various types of NPs. Synthesis methods are broadly divided into three parts viz., physical methods, chemical methods and bio-assisted methods. In addition, attempts have also been made to provide a brief discussion on the key applications of different types of nanoparticles.

2. Methods for the synthesis of nanoparticles
High-throughput NPs with good and controlled quality are desirable for their commercialization in various fields of applications. There are two basic approaches commonly employed to prepare NPs; 1) Top-down approach, where synthesis is initialized with the bulk counterpart that leaches out systematically bit-after-bit leading to the generation of fine NPs. Photolithography, electron beam lithography, milling techniques, anodization, ion and plasma etching are some of the commonly used top-down methods for the mass production of NPs; 2) Bottom-up approach, which involves the coalescence or assembling of atoms and molecules to generate diverse range of NPs. Examples of bottom-up approach include self-assembly of monomer/polymer molecules, chemical or electrochemical nanostructural precipitation, sol–gel processing, laser pyrolysis, chemical vapour deposition (CVD), plasma or flame spraying synthesis and bio-assisted synthesis. In general, NP synthesis methods can be divided in three groups - 1) physical methods, 2) chemical methods, and 3) bio-assisted methods (Fig. 1).
**Fig. 1** Overview on different type of synthesis methods to produce variety of nanoparticles.

### 2.1 Physical methods for the synthesis of nanoparticles

Physical methods apply mechanical pressure, high energy radiations, thermal energy or electrical energy to cause material abrasion, melting, evaporation or condensation to generate NPs. These methods mainly operate on top-down strategy and are advantageous as they are free of solvent contamination and produce uniform monodisperse NPs. At the same time, the abundant waste produced during the synthesis makes physical processes less economical.

High energy ball milling, laser ablation, electrospraying, inert gas condensation, physical vapour deposition, laser pyrolysis, flash spray pyrolysis, melt mixing are some of the most commonly used physical methods to generate NPs.
2.1.1 High energy ball milling (HEBM): High energy ball milling, first developed by John Benjamin in 1970 to synthesize oxide dispersion strengthened alloys capable of withstanding high temperature and pressure, is a robust and energy efficient synthesis method to generate NPs with varying shapes and dimensionalities.\textsuperscript{62} In HEBM process, the moving balls transfer their kinetic energy to the milled material. This results in the breaking of their chemical bonds and rupturing of the milled materials into smaller particles with newly created surfaces (Fig. 2a). Milling media, milling speed, ball-to-powder weight ratio, type of milling (dry or wet), type of high energy ball mill (vibrator mill, planetary mill, attritor mill, tumbler ball mill, etc.), milling atmosphere and duration of milling regulate the amount of energy transfer between the balls and the material during the process, and thus affect the physical and morphological properties of the resultant nanomaterials. The HEBM process sometimes involve very high local temperature (>1000 °C) and pressure (several GPa) conditions and thus also considered as a mechanochemical synthesis process.

Using metallic iron powder as the precursor material, Carvalho \textit{et al.} synthesized superparamagnetic magnetite NPs with size ranging from 12 nm to 20 nm by HEBM process.\textsuperscript{63} Bartwal and co-workers optimized various milling parameters (milling time, rotation per minute; rpm, ball size, etc.) to prepare uniform LiNbO\textsubscript{3} NPs with particle size ranging from ~30 to 60 nm.\textsuperscript{64} Salah and co-workers synthesized ZnO NPs of ~30 nm size from ZnO microcrystalline powder using HEBM process.\textsuperscript{65} Synthesized ZnO NPs showed antibacterial activities due to their desirable lattice constant. Chen \textit{et al.} utilized microwave assisted HEBM method to prepare pure and well-crystallized cobalt ferrite NPs with mean size of 20 nm, and high saturation magnetization (82.9 emu/g), at low temperature (≤ 100°C) without subsequent calcination.\textsuperscript{66} Cobalt oxalate hydrate and Fe powder were used as the raw materials for this synthesis and stainless steel or pure iron balls with diameter of 1.5 mm were used for milling. Xing \textit{et al.} recently established HEBM as the powerful green synthesis method for large-scale production of nitrogen doped carbon NPs for catalytic applications.\textsuperscript{62}
Particle Aggregation due to high surface energy generated on the milled particles during HEBM resulting in generation of larger particles

Surfactant molecules adsorbed on the surface of the generated particles and helps in lowering their surface energy by creating an organic layer on them and thus preventing them from aggregation. It leads to the production of nanoparticles with smaller size range with desired surface properties.

**Fig. 2** (a) High energy ball milling (HEBM) system and (b) Schematic representation of the NPs synthesis using HEBM method with and without surfactant.

Currently, surfactant assisted high energy ball milling is used as an efficient strategy for the synthesis of NPs with precise size and specific surface characteristics (Fig. 2b). Surfactants are the surface active agents containing both hydrophobic and hydrophilic properties and can be classified as anionic, cationic, zwitterionic and nonionic depending upon the surface charge characteristics of their hydrophilic group. Following adsorption on the material surface, the surfactant molecules generate electrostatic/steric forces which stabilize the milling particles, and thus minimize the uncontrolled fracturing of particles. Surfactant can also lower the surface energy of the freshly generated fine particles by forming thin organic layer and introducing long range capillary forces that lower the energy for crack propagation. This prevents the particles from agglomeration and cold welding that may lead to enhancement of particle size. Nature and amount of the surfactant used during the HEBM tremendously affect the physical characteristics of the NPs. Normally, increasing the surfactant volume can decrease the particle size by second to third order of magnitude. Zheng
et al. designed rare-earth-based magnetically hard single-crystal NPs of Sm$_2$Co$_{17}$ with average diameter of 9 nm and 26 nm by HEBM using oleic acid and oleylamine, respectively, as the surfactant with SmCo$_5$ as precursor.\textsuperscript{68} Islam et al. used this surfactant assisted HEBM method to prepare pure CaCO$_3$ NPs with diameter of 30 nm from cockle shells using dodecyl dimethy betain (BSL-12).\textsuperscript{69} Surfactant assisted HEBM has also been used to synthesize graphite NPs of the size ranging from 1-30 nm,\textsuperscript{70} where the carbon black was used as a precursor and phosphate ester as a surfactant in aqueous media.

2.1.2 Inert gas condensation (IGC): Inert gas condensation is one of the primitive methods for the NP synthesis that employ inert gases (e.g. He or Ar) and liquid nitrogen cooled substrate holder for the preparation of NPs. The evaporated materials are transported with inert gases and condensed onto the substrate attached with liquid nitrogen. Ward et al. used this method for the synthesis of Mn NPs.\textsuperscript{71} Perez-Tijerina et al. fabricated AuPd NPs using IGC process and studied the effect of annealing on the structure of the deposited NPs.\textsuperscript{72} IGC was found to be highly efficient method for the synthesis of good quality silver and platinum NPs.\textsuperscript{73} Sputtered deposition unit was used as the source for the generation of these NPs. Raffi et al. also reported the synthesis of Ag NPs by IGC method and found a strong dependence of the evaporation temperature (varied between 1123K and 1423K) and inert gas (He) pressure (varied between 0.5 and 100 Torr) on the morphology, crystallinity and size distribution.\textsuperscript{74} The mean particle size was found to increase with increasing evaporation temperature and He gas pressure and was in the range from 9 nm to 32 nm. Fabrication of size controlled Cu NPs (1-5 nm) was performed using IGC method combined with glow discharge sputtering as a source of Cu NPs.\textsuperscript{75}

Benelmekki et al. reported the synthesis of metallic-dielectric multi-core-shell NPs by IGC method.\textsuperscript{76} The FeAg NPs were deposited in Si shell (Fig. 3a-c), where Si shell protected the NPs against oxidation and restrained their aggregation. The basic principle for the synthesis of such hybrid nanoparticles (HNPs) was based on the condensation of atomic vapor produced from sputtering of multiple targets (Fe, Ag and Si) simultaneously at high pressure. Argon working pressure was kept at 10 mbar during the deposition. Using DC sputtering, vapours of metal atoms were produced near the target surface and condensed into nanoclusters along its propagation path through aggregation zone as shown in Fig. 3a. These nanoclusters were then extracted and accelerated due to difference in pressure between aggregation zone (10 mbar) and deposition unit (10$^{-5}$ mbar) and then deposited onto the
substrate (Fig. 3b and 3c). The mechanism responsible for the fabrication of multi-core-shell HNPs is illustrated in Fig. 3d. The mechanism was divided into three steps. **Step 1** was associated with the nucleation and growth of the NPs. Plasma density was found to be an important parameter, which influences the nucleation, growth and crystallinity of the NPs. The plasma density and sputtering yield (~1.2) were maximum in case of Ag. This led to the nucleation and growth of the highly crystalline Ag NPs before leaving the plasma zone, which then quenched into mono-crystalline NPs in aggregation zone. Since the plasma density and sputtering yield (0.45) was lower in case of Fe than Ag, their nucleation and growth occurred when the atoms reached the aggregation zone. This transportation process caused loss of energy of the atoms resulting in the creation of amorphous nanoclusters/NPs. The plasma density and sputtering yield (0.29) was lowest in case of Si leading to the formation of amorphous Si nanoclusters/NPs. **Step 2** involve cluster-cluster collisions, where the nanoclusters collide and coalescence with each other in aggregation zone, leading to the formation of larger NPs. However, formation of core/shell or dumbbell-like structures was most probable due to the large positive free energy of Ag and Fe mixing. In other words, during collision in aggregation zone, Ag and Fe nanoclusters could not completely coalescence due to lack of energy. This led to the formation of dumbbell-like structures in amorphous Si matrix. Finally, during their movement through aggregation zone, AgFe/Si core-shell NPs were found to collide and partially coalescence with each other resulting in the formation of multicore-shell HNPs in **step 3**. Fabricated HNPs were found to be the promising candidates for magneto-optic bio-applications.
2.1.3 Physical vapor deposition (PVD): Physical vapor deposition is a collective set of processes commonly used to produce NPs and to deposit thin layers of material, typically in the range of few nanometers to several micrometers. PVD is an environment friendly vacuum deposition technique consisting of three fundamental steps: 1) vaporization of the material from a solid source, 2) transportation of the vaporized material, 3) nucleation and growth to generate thin films and NPs. Most commonly used PVD methods for NPs synthesis are:

i) Sputtering
ii) Electron beam evaporation
iii) Pulsed laser deposition
iv) Vacuum arc

Fig. 4 shows the schematics representations of different PVD techniques.
2.1.3.1 Sputtering: Sputtering is a vacuum-based PVD process, which is often used to deposit films and NPs (Fig. 4a). Sputtering works on the principle of momentum transfer in which the atoms from the target (which is made up of material to be deposited) are ejected by the ion bombardment. The deposition of material by sputtering can be achieved using DC, pulsed DC and radio frequency (RF) powers. The sputter deposition takes place in following steps:

i) The plasma of neutral gases, commonly Ar, is generated between the two electrodes by the collisions of electrons to gaseous molecules.

ii) Ions present in the plasma are then accelerated toward the target by applying potential between the two electrodes.

iii) These ions with appropriate energy thus hit the target, leading to ejection of the material from the target.

iv) The ejected material are then transported and deposited onto the substrate.
Simple sputtering and magnetron sputtering are two types of sputtering processes. The difference between the two processes arises due to the fact that simple sputtering does not contain magnets behind the target, while magnets are always present in case of magnetron sputtering. The magnetron sputtering is upgraded version of simple sputtering and has following two advantages over former process:

- Higher deposition rate
- Prevent target overheating and damage

Higher deposition rate in magnetron sputtering is obtained due to the fact that under the influence of magnetic field the path of electrons become curved i.e. they follow helical path. This leads to more ionization of gases, causing more ions hit to the target, and hence results in higher deposition rate. In sputtering, sputtering yield is an important parameter, which describes the number of atoms ejected per incident ions. It should be noted that DC sputtering can be used to deposit nanostructures of conducting materials, while the deposition of insulating material by DC sputtering is difficult. However, RF sputtering can be used for the deposition of insulating materials also. Further, when reactive gases are introduced into the sputtering process chamber then sputtering process can be called as reactive sputtering. Reactive sputtering is used to deposit variety of compound/hybrid materials.

In all types of sputter deposition processes the NPs of ideal size can be synthesized by selecting the appropriate deposition conditions. For example, Hatakeyama and Nishikawa have synthesized Au NPs by analytically varying the sputtering conditions that influences the particle size and its distributions.\textsuperscript{78} Veith \textit{et al.} used magnetron sputtering for the deposition of Au NPs on WO\textsubscript{3} and activated carbon surfaces.\textsuperscript{79} Using magnetron sputtering and selecting the appropriate deposition parameters, Bouchat \textit{et al.} reported the synthesis of variety of metallic and non-metallic NPs including titanium oxide (TiO\textsubscript{2}), silver (Ag), gold (Au), yttrium (Y), carbon (C), cobalt (Co) and iron (Fe).\textsuperscript{80} Asanithi \textit{et al.} optimized the process conditions for the deposition of uniform sized, < 10 nm, silver NPs by DC magnetron sputtering.\textsuperscript{81} While varying target-to-substrate (TTS) distance from 10 to 20 cm, they demonstrated significant tailoring of particle size with the formation of highly uniform small grains (3.8±0.7 nm) at optimized TTS distance of 20 cm. Moreover, TTS was also reported to influence the shape and distribution of the Ag NPs. They further tuned the sputtering current (50 mA) at optimized TTS to design monodisperse Ag NPs. However, at higher sputtering current worm-like morphology with increased grain size was obtained. Ichida \textit{et al.}
synthesized Germanium (Ge) NPs and Ge quantum dots (Ge QDs), both used in electronic and optoelectronic industry, by high pressure RF magnetron sputtering.  

![Fig. 5](image)

**Fig. 5** High resolution transmission electron microscopy images of Pt NPs synthesized by sputtering process. (a)-(f) Pt NPs grown at constant target angle of 23.8° but varied growth time of (a) 5s, (b) 10s, (c) 20s, (d) 30s, (e) 45s and (f) 55s. (g)-(i) Pt NPs grown at constant growth time of 2min but varied target angle of (g) 16.2°, (h) 23.8° and (i) 38.8°. Reproduced with permission from ref. 85.

Ghosh et al. fabricated Mn doped ZnS NPs (ZnS:Mn) by RF magnetron sputtering and studied their photoluminescence and field emission properties. At an optimum Mn concentration, the excellent photoluminescence and field emission properties in ZnS:Mn NPs were realized. In another work, Veith et al. fabricated 2 nm size Au NPs on γ-Al₂O₃ using one step magnetron sputtering process which was comparable to those obtained by traditional
chemical methods. Ramalingam et al. employed tilted-target sputtering in order to grow sub-2 nm sizes and density tuneable Pt NPs at room temperature. They showed that tilting of target angle and variation of growth time significantly altered the mean diameter and density of Pt NPs (Fig. 5).

Sputtering process is also widely used for the synthesis of magnetic NPs for numerous applications. For example, Zhang et al. demonstrated the synthesis of L1₀ ordered FePt NPs on MgO substrate by DC magnetron sputtering. The FePt NPs were grown at base pressure of 5×10⁻⁷ Torr and low argon pressure of 1.5×10⁻³ Torr at 750 °C. During the deposition of FePt NPs the sputtering rate was 1.2 Å/s. Takahashi et al. synthesized FePt NPs by sputtering method and studied the effect of size on ordering of the NPs. They suggested that there is a critical value of size of NPs for ordering i.e. FCC FePt NPs with size < 4nm diameters do not display L1₀ ordering. Further, Dmitrieva et al. fabricated FePt NPs by sputtering process and suggested that inclusion of nitrogen during NP deposition can enhance the L1₀ phase of FePt NPs.

2.1.3.2. Electron beam evaporation (EBE): Electron beam evaporation is a vacuum-based PVD process which is used to fabricate thin films and NPs. EBE system consists of vacuum unit, electron beam source and target materials (Fig. 4b). Electron beam source consists of a filament, which is heated by passing current through it resulting in the generation of electron beam. The generated electron beam is focused and then directed onto the target material using magnets. The deposition of films by EBE takes place in the following steps:

i) Electron beam hits the target and heat the target material
ii) Target material (atoms) evaporates when temperature reaches above its boiling point
iii) Evaporated material then transported and
iv) Condensed onto the substrate

Advantage of EBE methods are

i) High deposition rate
ii) Can be used to deposit materials ranging from conducting to insulating
iii) Unlike thermal evaporation, EBE can be used to deposit materials of higher boiling point

By selecting optimum process parameters and conditions during EBE process, NPs of different sizes and shapes can be deposited. Perry et al. have reported the use of EBE in the
development of NPs for both 2D and 3D metal patterning. Hsieh et al. synthesized Au NPs and Pt NPs on multiwall carbon nanotubes (MWCNTs) to form composite electrodes for sensors and energy storage applications. They observed that the distribution of Au NPs was better on MWCNTs than Pt NPs. Uhm et al. demonstrated the fabrication of antibacterial Ag NPs on diameter-controlled TiO$_2$ nanotubes by facile EBE process. Further, Castaldi et al. prepared CoPt NPs of size between 5 nm and 20 nm on thermally oxidized silicon using EBE process and found that the size of NPs was dependent on the thickness of the CoPt layer.

Using EBE process, Bello et al. fabricated Ag NPs decorated three dimensional graphene (GR) scaffold for electrochemical application. In comparison to pure GR electrode, the GR/Ag NPs electrode showed higher capacitive performance and better cyclability for supercapacitor applications.

2.1.3.3. Laser ablation (LA) and pulse laser deposition (PLD): Laser ablation method utilizes a high power laser beam that evaporates particles from a solid source. In normal LA process, the laser can be either continuous laser or pulsed laser. LA offers a flexible approach in the production of micro- and nano-structures of polymeric materials. Kris et al. reported a micro-lens assembly with the utilization of LA technique by scanning the polymer surface to achieve a lens shape with the optimal focal distance and diameter. Gopal et al. have synthesized colloidal zinc metallic NPs using the LA technique.

PLD is another vacuum-based PVD process that employs laser energy to remove the material from the target. The high power laser pulses hit the surfaces of target, leading to melting, evaporation and ionization of the material (Fig. 4c). Finally the ablated materials deposit onto the substrate. PLD utilize pulsed laser beam, mainly coming from either excimer laser or Nd:YAG laser, for the ablation of the material. PLD is used for the preparation of variety of materials including polymers, oxides, metallic systems, fullerenes, carbides, nitrides, etc.

In context to NPs production, Lee et al. reported the synthesis of FeCo NPs in an inert gas atmosphere by PLD. Ong et al. examined the role of process parameters such as number of pulses, ambient gas pressure and temperature gradient on the morphology and size of FeCo NPs synthesized by PLD. The morphology of FeCo was changed from linear interconnected chains to fibrous when the number of pulses were increased. At low pressure, NPs tend to form flocules-like nano-networks while chain-like network was observed at higher pressure. In addition, the average size of NPs was increased with increasing gas pressure. Andrea et al. used PLD for the synthesis Ag NPs arrays for surface enhanced Raman
scattering (SERS).\textsuperscript{99} They controlled the size and morphology of Ag NPs by optimizing the Ar pressure and number of pulses. In another work, Jing \textit{et al.} deposited Ag NPs on nickel hydroxide nanosheet arrays using PLD for application to SERS.\textsuperscript{100} They found that deposition time influenced the size and inter-particle gap of Ag NPs and hence could control the SERS enhancement. Kumar \textit{et al.} synthesized self-assembled magnetic NPs by PLD and demonstrated the embedment of Fe NPs and Ni NPs of sizes 5-10 nm in amorphous alumina and crystalline TiN materials.\textsuperscript{101} Quintana \textit{et al.} performed an interesting work involving the fabrication of selenium (Se) NPs on three different substrates: 1) metallic Au film, 2) Si wafers and 3) glass using PLD.\textsuperscript{102} While analyzing the morphology of the deposited NPs, they found that size, shape and population density of Se NPs were strongly dependent on the experimental process parameters and underlying base such as energy density, number of pulses and substrate. Lin \textit{et al.} studied the effect of target-substrate geometry and ambient gas pressure on the synthesis of FePt NPs by PLD.\textsuperscript{103} They suggested that backward plume, which is a novel target-substrate geometry can produce FePt NPs with high uniformity, better phase formation than conventional PLD process. Amoruso \textit{et al.} fabricated Ni NPs by femtosecond PLD, which was confirmed by atomic force microscopy (AFM).\textsuperscript{104} Dhlamini \textit{et al.} reported the synthesis of lead sulphide (PbS) NPs in amorphous SiO\textsubscript{2} matrix on Si substrate using PLD and realized excellent photoluminescence properties.\textsuperscript{105} Table 2 covers different NPs synthesized by the PLD method with the process details.

\textit{Table 2} Various nanoparticles synthesized using PLD technique.

<table>
<thead>
<tr>
<th>Type of Nanoparticles</th>
<th>Deposition Medium</th>
<th>Deposition Parameters</th>
<th>Size of Nanoparticles</th>
<th>Laser Type</th>
<th>References</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ag NPs</td>
<td>Aqueous SDS solution</td>
<td>LI: 120 mJ/pulse RR: 10 Hz PW: 5 ns TTS Distance: 4.5 mm</td>
<td>4.2 ± 1.9 nm in SDS (Anionic Surfactant)</td>
<td>Nd:YAG Laser, 532 nm</td>
<td>94</td>
</tr>
<tr>
<td></td>
<td>Aqueous CTAB solution</td>
<td>LI: 120 mJ/pulse RR: 10 Hz PW: 5 ns TTS Distance: 4.5 mm</td>
<td>7.8± 4.5 nm in CTAB (cationic Surfactant)</td>
<td>Nd:YAG Laser, 532 nm</td>
<td>94</td>
</tr>
<tr>
<td></td>
<td>Aqueous SDS solution</td>
<td>LI: 60 mJ/pulse RR: 10 Hz PW: 5 ns TTS Distance: 4.5 mm</td>
<td>6.8 ± 2.7 nm in SDS</td>
<td>Nd:YAG Laser, 532 nm</td>
<td>94</td>
</tr>
<tr>
<td></td>
<td>Aqueous CTAB solution</td>
<td>LI: 60 mJ/pulse RR: 10 Hz PW: 5 ns TTS Distance: 4.5 mm</td>
<td>9.4 ± 5.9 nm in CTAB</td>
<td>Nd:YAG Laser, 532 nm</td>
<td>94</td>
</tr>
<tr>
<td>Ag NPs</td>
<td>Ar gas, 70</td>
<td>BP: 2 × 10\textsuperscript{-7} Pa</td>
<td>1.5-4.5 nm</td>
<td>248 nm KrF</td>
<td>99</td>
</tr>
<tr>
<td>Material</td>
<td>Pressure/Environment</td>
<td>Specifications</td>
<td>Target Substrate Distance</td>
<td>Notes</td>
<td></td>
</tr>
<tr>
<td>----------</td>
<td>----------------------</td>
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<td></td>
</tr>
<tr>
<td>Ni NPs</td>
<td>Vacuum</td>
<td>RP: 10⁻³ Pa, LI: 1 mJ/pulse, LIA: 45°, TTS: 30 mm, NLS: 200, LED: 0.3 J cm⁻²</td>
<td>5-100 nm MS ≈ 40 nm</td>
<td>1 kHz Ti:sapphire oscillator–amplifier system emitting at 780 nm</td>
<td></td>
</tr>
<tr>
<td>Fe NPs</td>
<td>High Vacuum</td>
<td>BP: 8 × 10⁻³ Torr, RR: 10 Hz, PW: 10 ns, LED: 2 J cm⁻², ST: 500° C</td>
<td>~10 nm</td>
<td>-</td>
<td></td>
</tr>
<tr>
<td>Se NPs</td>
<td>High Vacuum</td>
<td>BP: 8 × 10⁻³ Torr, PD: 108 W cm⁻², PW: 10 ns, LIA: 45°, TTS: 35 mm</td>
<td>75-100 nm</td>
<td>Nd:YAG Laser, 532 nm</td>
<td></td>
</tr>
<tr>
<td>ZnO NPs</td>
<td>Aqueous SDS solution</td>
<td>RR: 10 Hz, LI: 100 mJ/pulse, IT: 60 mins</td>
<td>9.4-32.1 nm, AD: 14.7 nm</td>
<td>Nd:YAG Laser, 532 nm</td>
<td></td>
</tr>
<tr>
<td>PbS</td>
<td>O₂ Gas, 300 mTorr</td>
<td>RR: 10 Hz, PW: 30 ns, TTS Distance ~ 50 mm, LI: 183 mJ/pulse</td>
<td>10-50 nm AD: 17 nm</td>
<td>248 nm KrF laser</td>
<td></td>
</tr>
<tr>
<td>FeCo NPs</td>
<td>Ar gas, Pressure</td>
<td>BP: 10⁻³ Pa, RR: 10 Hz, PW: 10 ns, ST: 150° C</td>
<td>4.8-8.6 nm</td>
<td>Nd:YAG Laser, 532 nm</td>
<td></td>
</tr>
<tr>
<td>FePt NPs</td>
<td>Ar gas</td>
<td>BP: 5 × 10⁻³ mbar, LI: 75 mJ/pulse, LED: 955 J cm⁻², TTS: 6 mm</td>
<td>4.8-22 nm</td>
<td>Nd:YAG Laser, 532 nm</td>
<td></td>
</tr>
</tbody>
</table>

**Abbreviations:** AD- Average Diameter, BP- Basic Pressure, CTAB- Cetyltrimethylammonium bromide, IT- Incubation Time, LED- Laser energy density, LI- Laser Intensity, LIA- Laser Incident Angle, MS- Mean Size, NLS- Number of Laser Shots, PW- Pulse Width, RP- Residual Pressure, RR- Repetition Time, SDS- Dodecylsulfuric acid sodium salt, ST- Substrate Temperature, TTS Distance- Target-Substrate Distance
2.1.3.4 Vacuum arc (VA): Vacuum arc is a vacuum-based PVD process in which the arc is used to vaporize the material for the synthesis of metallic, ceramic and composite NPs and films (Fig. 4d). Akbari et al. synthesized the Mg-Al alloy NPs by varying Mg and Al concentrations using plasma arc process. Lei et al. used vacuum arc discharge process for the synthesis of inter-metallic Fe-Sn NPs. Amaratunga demonstrated the synthesis of thin film of linked carbon NPs using localized high pressure arc discharge process and realized excellent mechanical properties in deposited material. Chhowalla and Amaratunga used a similar approach of localized high pressure arc discharge for the synthesis of thin film of fullerene-like MoS$_2$ NPs, which showed extremely low friction. Fig. 6a and 6b show the high resolution electron microscope images of thin films of cross-linked carbon NPs and fullerene-like MoS$_2$ NPs, respectively.

![Fig. 6](image)

**Fig. 6** (a) High resolution electron microscopic image of thin film of carbon cross-linked NPs and (b) High resolution transmission electron microscopic image of thin film of fullerene-like MoS$_2$ NPs. Reproduced with permission from ref. 108 and ref 109.

2.1.4 Laser pyrolysis: The CO$_2$ laser pyrolysis technique, first developed by Haggerty et al. at the beginning of 1980s, is a vapour phase synthesis process. This method can be used to synthesize NPs of large variety of oxide (TiO$_2$, SiO$_2$, Al$_2$O$_3$, Fe$_2$O$_3$), non-oxide (Si, SiC, Si$_3$N$_3$, MoS$_2$) and ternary composites like Si/C/N and Si/Ti/C. In this process, the resultant condensable products are generated from the laser-induced chemical reactions at the interface of the laser beam and the molecular flow of gaseous/vapour phase reactants (Fig. 7a). One of the criteria to induce energy coupling into the reaction system is that one of the reactant/precursor should be able to absorb through resonant vibrational mode of Infra-red (IR) CO$_2$ laser radiations. Alternatively, additional chemicals like ammonia (NH$_3$), sulphur hexafluoride (SF$_6$), ethylene gas (C$_2$H$_4$), etc. are introduced among the reactants as inert photo-sensitizers to accomplish the energy transfer process between the laser light and
precursors. The high power of the CO$_2$ laser initiates the sequential absorption of several IR photons in the same molecule, followed by collision assisted energy pooling, leading to a rapid increase of the average temperature in the gas through vibration-translation energy transfer processes, often accompanied by the appearance of a flame in the interaction volume. If molecules are excited above the dissociation threshold, molecular decomposition, eventually followed by chemical reactions, occurs with the formation of condensable and/or volatile products. Compared to other vapour phase methods, the laser pyrolysis permits highly localized and faster heating (leads to rapid nucleation) with faster quenching of the particle growth (in few ms). Thus, this methodology is promising in generating NPs of the average diameter ranging between 5 to 60 nm with a narrow size distribution in the hot zone. However, as soon as the NPs escape the hot chamber, they try to aggregate resulting in the formation of NPs chains. High energy ball milling and high-power ultrasonic treatment are the effective mechanical processes that help in generation of well-separated NPs using laser pyrolysis method.

Rosaria D’Amato et al. utilized laser pyrolysis technique to synthesize variety of ceramic NPs (e.g. SiC, SiO$_2$ and TiO$_2$) for energy applications and cultural heritage preservation using silane (SiH$_4$), tetraethyl orthosilicate (Si(OC$_2$H$_5$)$_4$) and titanium(IV) isopropoxide respectively as precursor materials.\textsuperscript{110} Owing to the advantage of single step doping in this method, Scarisoreanu et al. synthesized carbon coated TiO$_2$ NPs for photocatalysis applications.\textsuperscript{111} Marino et al. synthesized carbon nitride NPs with varying nitrogen incorporation between 2% and 20% using laser pyrolysis.\textsuperscript{112} Jansa et al. fabricated carbon NPs using pulsed laser pyrolysis of hydrocarbons and found that at same experimental parameters, the carbon NPs produced using benzene showed higher conductivity than those produced using either acetylene or ethylene.\textsuperscript{113} They also found the variation of size of NPs with varying laser power.

Silicon carbide (SiC), silicon and molybdenum disulphide (MoS$_2$) NPs have also been produced by the laser pyrolysis technique using the same technique.\textsuperscript{114} Veintemillas-Verdaguer et al. synthesized magnetic composites Fe-based NPs encapsulated in carbon/silica (C/SiO$_2$@Fe) or carbon (C@Fe) with average diameter of 29±3 nm and 56±3 nm, respectively.\textsuperscript{115} Laser Pyrolysis of the carbonyl precursors is emerging as an excellent technique for the direct synthesis of iron NPs with smaller particle size, narrow size distribution without much noticeable aggregation. Following the same lines, Bomat et al. have prepared the iron–iron oxide core–shell NPs (10-30 nm) by laser pyrolysis of Fe(CO)$_5$.
vapours and in-situ passivation by controlled oxidation process.\textsuperscript{116} Erogbogbo \textit{et al.} have reported the synthesis of biocompatible Si QD by laser-driven pyrolysis of silane using phospholipid micelles for the imaging of cancer cells.\textsuperscript{117}

\textbf{Fig. 7} Schematic Illustration of (a) laser pyrolysis, (b) flame spray pyrolysis and (c) electrospraying technique for the synthesis of NPs.
2.1.5 Flame spray pyrolysis (FSP): FSP is the latest of all the flame aerosol technologies.\textsuperscript{118} It is a one-step combustion process where the precursor is in liquid form, with significantly higher combustion enthalpy (>50% of total energy of combustion), usually in an organic solvent. The important technological elements of this process include self-sustaining flame, usage of liquid feeds and less volatile precursors, proven scalability, high temperature flames and large temperature gradients (Fig. 7b). It is one of the most exploited techniques for the production of complex and functional NPs. For the formation of particles, the liquid precursor can follow the droplet-to-particle route or gas-to-particle route; however, later results in more homogenous morphologies and size. For NPs fabrication, precursor spray has to follow the sequential steps viz., 1) precursors evaporate/decompose forming metal vapors; 2) nucleation as a result of supersaturation; 3) growth by coalescence and sintering; and 4) particle aggregation (by chemical bonds) and agglomeration (by physical interactions) (Fig. 7b). Fig. 8 summarise all the particle configurations (and combinations) obtained by FSP. FSP of single metal component precursors in open ambient (air) condition produces simple oxides or carbonates in the case of alkaline metals through reaction with the CO\textsubscript{2} laden flame and follow Route I (Fig. 8) for the NPs synthesis. NPs of wide range of elements across the periodic table, from MgO to Yb\textsubscript{2}O\textsubscript{3}, have been synthesized via this route. FSP of metallic NPs including Co, Bi and Cu (following Route 2 in Fig. 8) have been reported under the anaerobic conditions that helps preventing rapid oxidation of Bi to Bi\textsubscript{2}O\textsubscript{3} and resulting in the formation of the protective surface oxide or carbon layer on the Co and Cu NPs. For the multi-component systems, the particle formation pathways are much more complicated and follow different routes (Route III-VI, Fig. 8).
Fig. 8 (a) Formation of different particle configurations via the gas-to-particle mechanism for single and multicomponent systems, (b) as-prepared metal oxides NPs of different morphologies made by FSP, varying from hexagonal/octagonal platelet g-Fe$_2$O$_3$, lightly oblongated ZnO, spherical TiO$_2$, and rhomboid-shaped CeO$_2$ with sharp edges. Reproduced with permission from ref. 118.

Sokolowski et al. first utilized this strategy to synthesize Al$_2$O$_3$ NPs by combusting an ultrasonically-dispersed spray of aluminium acetylacetonate in benzene–ethanol solvent mixture.\textsuperscript{119} Later, FSP process was used tremendously to design various oxide and non-oxide ceramic NPs e.g. CeO$_2$, SiO$_2$, Al$_2$O$_3$, MgO, C-Cu, BaCO$_3$, CaO, CaCO$_3$, CaSO$_4$, etc. Madler et al. adequately reviewed the research articles concerning the design and fabrication of NPs by FSP method and also tabulated very nicely all different oxide and non-oxide ceramic NPs synthesized using this method.\textsuperscript{118} Some of the most notable applications of the FSP derived NPs are in the areas of catalysis optics and photonics, sensors, health care, magnetic materials, etc.\textsuperscript{120-123} For example, Arutanti et al. fabricated WO$_3$/TiO$_2$ NPs by flame-assisted spray pyrolysis and explored them for photocatalysis.\textsuperscript{124} They studied the role of amount of WO$_3$ and TiO$_2$ in the range 0–100 wt% on size, shape, crystallinity, and photocatalytic performance. While analyzing the results they found that photocatalytic performance of composite WO$_3$/TiO$_2$ NPs was superior compared to 100 wt% TiO$_2$ catalyst. Channei et al. synthesized Fe-doped CeO$_2$ NPs with varying amount of Fe using flame-assisted spray pyrolysis for photocatalysis application and found that photocatalytic efficiency was 100% when 2 mol% Fe was introduced in CeO$_2$.\textsuperscript{125} Similarly Siriwong et al. synthesized Pt-doped Zn$_2$TiO$_4$ NPs with varying amount of Pt from 0.25-1.0 mol% using flame-assisted spray pyrolysis and suggested promising use of this process for the synthesis of high quality and controlled sized NPs.\textsuperscript{126}

2.1.6. Electrospraying technique: Electrospraying is a technique similar to electrospinning that differs in the type of materials to be produced. The former is done to synthesize NPs while the latter is done for the fabrication of nanofibers.\textsuperscript{127} The electrospraying method is based on an electromechanical device in which a mixture of solution containing the selected polymer and the solvent is taken up by the syringe, and a high voltage is applied to the capillary tip that results in the production of charged droplets (Fig. 7c). The solvent is evaporated on its way to the counter electrode and particles or fibers are then collected as the end product. The electrospraying technique provides both flexibility and control of the surface parameters.\textsuperscript{128} Deppert et al. have reported distinct NPs deposition and the formation of gold NPs for nanostructure growth by utilizing the electro-spraying technique.\textsuperscript{129} Sridhar et
al. reported a comprehensive review over the use of electrospaying technique to produce lipid-based delivery systems and biodegradable polymer NPs as carriers for drug delivery in application to a wide variety of therapeutic area such as cancer therapy, anti-inflammatory, antibacterial agents, etc.\textsuperscript{130} Maedeh et al. reported the assembly of core-shell PLGA NPs encapsulating the protein, bovine serum albumin, by ‘coaxial’ electrospaying method that can be applied in the management of various severe diseases.\textsuperscript{131} Advantages of utilizing electrospaying technique include the ability to generate uniform NP size, fast preparation method and the capability for bulk assembly of NPs. However, this method can also lead to degradation of some macromolecule as a result of thermal or shear stresses in drying and in the syringe.\textsuperscript{130}

The electrospaying technique can also be utilized to transform water vapour present in the atmosphere into engineered water nano-structures (EWNS). The EWNS has a size of 25 nm that is equipped with an extraordinary set of physical, chemical, biological and morphological properties with low toxicity level that possess a remarkable mechanism of action as EWNS is loaded with reactive oxygen species (ROS) that is able to interact with and to stop the activity of harmful airborne bacteria.\textsuperscript{132} With the electrospaying method, NPs of various type and sizes can be synthesized based on the use of a variety of solvents (Table 3).

**Table 3** Different types of NPs synthesized using electrospaying method and their field of applications.

<table>
<thead>
<tr>
<th>Type of Nanoparticles</th>
<th>Solvent</th>
<th>Size</th>
<th>Application</th>
<th>References</th>
</tr>
</thead>
<tbody>
<tr>
<td>Au NPs</td>
<td>Water + methanol</td>
<td>20nm-100nm</td>
<td>Biomedical field, cancer nanotechnology, molecular imaging, molecular diagnosis, targeted therapy and bioinformatics.</td>
<td>133,134</td>
</tr>
<tr>
<td>Ag NPs</td>
<td>Toluene</td>
<td>3-7nm</td>
<td>Biosensor, anti-reflection coatings, artificial joint replacement, cancer diagnosis, anti-bacterial activity.</td>
<td>135,136</td>
</tr>
<tr>
<td>Sn NPs</td>
<td>Isopropyl alcohol</td>
<td>3 nm</td>
<td>Humidity sensor, microelectronics and batteries.</td>
<td>137,138</td>
</tr>
<tr>
<td>SiO\textsubscript{2} NPs</td>
<td>Ethylene glycol</td>
<td>20 nm</td>
<td>Photodynamic therapy, magnetic resonance imaging, optical imaging, drug delivery, gene delivery and protein delivery.</td>
<td>139,140</td>
</tr>
<tr>
<td>Si NPs</td>
<td>I-Octanol</td>
<td>3 nm</td>
<td>Biological interface devices, anti-cancer drug delivery and cancer detection.</td>
<td>139,141</td>
</tr>
<tr>
<td>Al\textsubscript{2}O\textsubscript{3} NPs</td>
<td>Ethanol</td>
<td>500 nm</td>
<td>Catalysis, waste water treatment, heat transfer fluidics, biosensors,</td>
<td>139,142</td>
</tr>
<tr>
<td>NPs</td>
<td>Composition</td>
<td>Diameter</td>
<td>Application</td>
<td>Reference</td>
</tr>
<tr>
<td>--------------</td>
<td>-----------------------------------------------------------------------------</td>
<td>----------</td>
<td>-------------------------------------------------------------------------------------------------------</td>
<td>-----------</td>
</tr>
<tr>
<td>Ni NPs</td>
<td>Ethylene glycol monooethyl ether acetate+alkynl phthalene+polyamine</td>
<td>2-3 nm</td>
<td>Magnetism, energy technology and biomedicine.</td>
<td>139,143</td>
</tr>
<tr>
<td>TiO2 NPs</td>
<td>I-Butanol</td>
<td>10-40 nm</td>
<td>Drug delivery, cosmetics, antibacterial materials, sun-screens and electronics.</td>
<td>139,144</td>
</tr>
<tr>
<td>ZrO2 NPs</td>
<td>I-Butanol</td>
<td>10–40 nm</td>
<td>Catalysis, anti-corrosion coating in surgical appliances, explosive primers, dental implant material, cell and tissue engineering.</td>
<td>139,145,146</td>
</tr>
<tr>
<td>Pt NPs</td>
<td>Ethanol</td>
<td>8 nm</td>
<td>Chemothrapy, cellular imaging, MRI, sun-screen protection and anti-septic products.</td>
<td>147,148</td>
</tr>
<tr>
<td>PLGA NPs</td>
<td>Acetonitrile and dimethyl sulfoxide</td>
<td>165nm – 1.2µm</td>
<td>Nanocarriers for drug delivery, cancer diagnosis and imaging.</td>
<td>131,149</td>
</tr>
<tr>
<td>PLA NPs</td>
<td>Butanol/methylene chloride mixture</td>
<td>100 nm</td>
<td>Drug encapsulation, thin film deposition and biomedical engineering.</td>
<td>150,151</td>
</tr>
<tr>
<td>PMSQ NPs</td>
<td>Ethanol</td>
<td>62 ± 20 nm</td>
<td>Organic thin film insulator and printed electronics.</td>
<td>152,153</td>
</tr>
<tr>
<td>PMMA NPs</td>
<td>Acetone</td>
<td>300 nm</td>
<td>Drug/SiRNA systems, artificial enzymes and photostable bio-imaging agents.</td>
<td>154,155</td>
</tr>
<tr>
<td>PCL NPs</td>
<td>Chlorofoam</td>
<td>4.8±0.2 - 17.7±0.5 µm</td>
<td>Pharmaceutics, drug delivery, cancer treatment and cell culture.</td>
<td>156,158</td>
</tr>
<tr>
<td>COLL/NaCl NPs</td>
<td>Acetic Acid (COLL/NaCl) or 630nm (COLL/CaCl2)</td>
<td>900nm</td>
<td>Biomedical and therapeutics, nanomedicine.</td>
<td>159,160</td>
</tr>
<tr>
<td>COLL/CaCl2 NPs</td>
<td>Acetic acid</td>
<td>520nm</td>
<td>Anti-cancer treatment, drug delivery, gene delivery and growth factor carriers.</td>
<td>161,162</td>
</tr>
</tbody>
</table>

**Abbreviations:** PLGA- Poly(lactic-co-glycolic acid), PLA- Poly-DL-lactic-acid, PMSQ- Polymethylsilsesquioxane, PMMA- Poly(methyl methacrylate), PCL- Polycaprolactone, COLL- Collagen

### 2.1.7 Melt mixing:
This method involves the mechanical mixing of a polymer with modified nanofillers by extrusion or kneading, and less commonly by injection moulding technique. This is one of the oldest methods to design polymer composites with NPs as the fillers to achieve desired material characteristics. This is the most commonly used mechanical process because it is environment friendly and it is also well-suited for current industrial practices.

Zuhal et al. have utilized the melt mixing technique to prepare nanocomposite NPs of conducting polymer polypyrrole with polypropylene and the resultant NPs have been then
processed further using injection moulding technique. Weis 
et al. have reported in-situ formation of hybrid NPs during melt blending of hydroxybenzoate (HBA) and 27% hydroxynaphthanoate liquid polymer polyesters with zinc salts of sulfonated polystyrene ionomers at elevated temperatures due to chemical reaction that involved the liquid crystalline polyester and residual zinc acetate from the neutralization of the ionomer.

2.2. Chemical methods for the synthesis of nanoparticles

Sol-gel method, microemulsion technique, hydrothermal synthesis, polyol synthesis, chemical vapour synthesis and plasma enhanced chemical vapour deposition technique are some of the most commonly used chemical methods for the NPs synthesis.

2.2.1. Sol-gel method: In the sol-gel processing method, there are two types of components viz., ‘sol’ which is a colloidal suspension of solid particles in a liquid and ‘gel’ which are polymers containing liquid. Thus, this process includes the creation of ‘sols’ in the liquid that lead to the formation of a network of discrete particles or network polymers by the connection of sol particles. Hydrolysis and condensation are the typical steps of sol-gel process, in which the former uses water to disintegrate the bonds of the precursor that is also the first step in formation of the gel phase. This process is then followed by condensation that leads to the formation of nanomaterials after which excess water is removed to determine the final structure of the material. For instance, ZnO NPs were synthesized by this method using zinc acetate dehydrate and triethanolamine (TEA) as the precursors. Since experimental conditions are very important for the synthesis of NPs by sol-gel method, Behnajady et al. optimized the synthesis parameters such as precursors, solvent percentage, water percentage, reflux temperature, reflux time, calcination temperature, sol drying method for the fabrication of TiO$_2$ NPs by this method. High photocatalytic activity from TiO$_2$ NPs was observed when they synthesized considering isopropoxide as a precursor, methanol as a solvent and titanium (IV) under 3 h reflux at 80 °C with sol thermal drying and calcination temperature of 450 °C. Further, the synthesis of size controlled SnO$_2$ NPs was demonstrated by optimizing the calcination temperature and molecular weight of polyethylene glycol (PEG, this is serving as chemical modifier reagent) using sol-gel method. Results suggest that the size of SnO$_2$ NPs decreases with increasing molecular weight of PEG from 300 to 4000 and decreasing the calcination temperature from 600 °C to 450 °C. Goncalves et al. demonstrated the one-pot synthesis of CdSe NPs using sol-gel method and realized quantum size effect in fabricated
NPs, which is important for developing advanced optoelectronic devices.\textsuperscript{170} Apart from them sol-gel is also a promising method for the synthesis of variety of NPs such as metal aluminate NPs, magnetic Fe-Co NPs, ZnO and Fe$_2$O$_3$ NPs, ZrB$_2$ NPs, GdVO$_4$ NPs, Ta$_2$O$_5$ NPs, etc.\textsuperscript{171-176}

2.2.2. Microemulsion technique: The term microemulsion was first assigned by Schulman et al. in 1959.\textsuperscript{177} Microemulsions can be defined as the thermally stable, macroscopically homogenous, optically transparent and isotropic dispersions constituting minimum of three components i.e., polar phase (generally water), non-polar phase (generally hydrocarbon liquid or oil) and surfactant. Surfactant molecules creates the interfacial layer separating the aqueous and the organic phases, reduces the interfacial tension between the microemulsion and the excess phase and act as a steric barrier preventing the coalescence of the droplets.\textsuperscript{177} Microemulsion system consists of monodispersed spherical droplets (diameter ranging from 600 nm to 8000 nm) of water-in-oil (w/o) or oil-in-water (o/w) depending on the surfactant used. The w/o reverse micellar system acts as an excellent reaction site for the NPs synthesis. Reverse micelle is water-in-oil microemulsion where the polar head groups of the surfactant creating the aqueous core and resides towards inside whereas the organic tails of the surfactant molecules directed towards outside as shown in Fig. 9a.

**Fig. 9** shows (a) typical Reverse Micelle System, (b) various steps involved in one microemulsion process and (c) reaction sequence involved in the two microemulsion nanoparticles synthesis.
In general there are two microemulsion routes to synthesize the NPs namely 1) one microemulsion method and the 2) two microemulsion method (Fig. 9b and 9c). One microemulsion method can be further divided into two types i.e., energy triggering method that needs a triggering agent to initiate the nucleation reaction within the single microemulsion containing the precursor and other is one microemulsion plus reactant method which is initiated by adding one of the reactant directly into microemulsion already carrying the second reactant (Fig. 9b). Kurihara et al. have reported the laser and radiolytically induced colloidal Au NPs formation in w/o microemulsion system with diameter ranging from 2.5-10 nm. One microemulsion processes are diffusion controlled since the second trigger/reactant has to diffuse through the interfacial wall of the microemulsion encapsulating the first reactant to accomplish the NPs synthesis. In two microemulsion method, the two microemulsions carrying the separate reactants are mixed together in appropriate ratios (Fig. 9c). Brownian motion of the micelles helps them to approach each other resulting in intermicellar collisions and sufficiently energetic collisions leads to the mixing of the micellar components. Once both the reactant comes in a same micellar compartment, the chemical reaction takes place in this nanoreactor. As the critical number of molecules (N_{crit}) attained inside the micelle, it initiates the nucleation process and results in NPs formation. Numerous intermicellar collisions are needed for the sufficient reactant exchange, their mixing and finally their reaction to terminate at the end product.

Microemulsion technique was used most commonly for the synthesis of the inorganic nanomaterials including metal NPs (Au, Pt, Pd), semiconducting metal sulphite NPs (CdS, PbS, CuS, Cu_2S and CdSe), metal salt NPs (BaCO_3, CaCO_3 and SrCO_3), metal oxide NPs (ZrO_2, TiO_2, SiO_2, GeO_2 and Fe_2O_3), magnetic NPs ((Mn,Zn)Fe_2O_4, (Ni,Zn)Fe_2O_4, ZnFe_2O_4 and BaFe_{12}O_{19}) and composite NPs (CdS-TiO_2, CdS-ZnS, CdS-SnO_2). Metal NPs were easily prepared using microemulsion technique using the reduction strategy. Water-to-surfactant molar ratio (ω), type of continuous phase, metal ion concentration, type and concentration of the reducing agent, structure and amount of the surfactant used are some of the key parameters that control the metal NPs synthesis effectively. Boutonnet et al. have used the reverse micelle microemulsion strategy to synthesize Pt, Pd, Rh and Ir NPs employing (H_2PtCl_6, Pd(NH_2)_4Cl_2, RhCl_2 and IrCl_3, respectively as the precursors with hydrazine (N_2H_4) and active hydrogen as the reducing agent. Solla-Gullon et al. have recently reported fast, easy and scalable production of Pt cubic NPs by reducing H_2PtCl_6 with sodium borohydride using w/o microemulsion process. They revealed significant effect of introducing varying
amount of HCl in the H$_2$PtCl$_6$/NaBH$_4$ microemulsion system in controlling the shape/surface of the Pt NPs. Zhang et al. have studied the synthesis of Ag NPs and also investigated the effect of $\omega$ (water/surfactant ratio) on the particle size and their size distribution. Results revealed that there is continuous decrease in the particle size from ~5 nm to ~1.5 nm with reduction in the $\omega$ value from 15 to 2.5. They also investigated the effect of reducing agent concentration on the Ag NPs synthesis and leads to the fact that both less and excessive amount of reducing agent can hinder the synthesis process by affecting the stability of the colloids. The same group have investigated the effect of increasing surfactant (sodium bis(2-ethylhexyl) sulfosuccinate; AOT) concentration on the Ag NP synthesis and observed reduction in the particle size with increasing surfactant amount. Pileni et al. have introduced the synthesis of Cu NPs by reverse micelles using hydrazine as the reducing agent and revealed the effect of increasing water content in controlling the NPs size and the oxide content within the NPs structure. CdS NPs were synthesized in AOT and triton reverse micelles with cadmium lauryl sulphate and cadmium AOT as the functional surfactant. The diameter of the NPs is reported to be dependent on the relative composition of Cd$^{2+}$ and S$^{2-}$ ions. Chen et al. have accounted the synthesis of Ni NPs with average diameter of 4.7 nm using cationic w/o microemulsion of water/CTAB (cetyltrimethylammonium bromide)/n-hexanol system by the reduction of nickel chloride with hydrazine at an elevated temperature. Subrata Kundu reported a novel methodology to synthesize shape-selective Au NPs by reduction of Au (III) ions with alkaline 2,7-Dihydroxynaphthalene (DNP) in CTAB micellar media under 30 min continuous UV-irradiation. The size and shape of the NPs were reported to be tuned by varying the Au (III): CTAB molar ratio. Similar synthetic pathway was accounted to synthesize catalytically enhanced shape selective rhodium nanoparticles (Rh NPs) in the presence of DNP (reducing agent) and CTAB (surfactant) using 6h UV-irradiation. The growth of the NPs with different shapes (spherical, flower-like or cubic) was found to be directed by the surfactant-to-metal ion molar ratio and the concentration of DHN. Praharaj et al. also reported the size selective synthesis of Au NPs with average diameter from 9 nm to 15 nm in toluene employing cationic surfactant of variable chain length. Asher and co-workers developed and demonstrated a novel strategy to synthesize silica-CdS nanocomposites that allocate tailoring the size and morphology of the resultant NPs. Gold–silver bimetallic NPs with varying mole fractions were synthesized in water-in-oil microemulsions of water/Aerosol OT/isooctane system by co-reduction of HAuCl$_4$ and AgNO$_3$ with hydrazine. Voucher et al. reported the preparation of...
crystalline NPs of hexacyanoferrate, cobalt Pentacyanonitrosylferrate and chromium hexacyanochromate coordination polymers in water-in-oil microemulsions.  \(^{191}\) Mandal et al. reported the synthesis of bimetallic core-shell Pd-Au and Pd-Ag NPs of the size range of 10-30 nm using TX-100 micellar system by seed-mediated method.  \(^{192}\) Microemulsion technique was not progressed much in term of designing organic NPs due to phase separation constraint and thus only few reports are available.  \(^{178}\) This method is known to be microemulsion polymerization when it comes to organic structures. Atkin and Thomas reported first successful microemulsion polymerization in 1981 of latex NPs (average diameter: 20 nm-35 nm) using CTAB/styrene/hexanol/water oil-water microemulsion coordination.  \(^{193}\) Three component microemulsion method was described to synthesize polystyrene NPs within size range of 20-30 nm using dodecyl trimethyl ammonium bromide (DTAB) and potassium persulphate (KPS) initiator.  \(^{194}\) Holdcroft and Guillet synthesized polystyrene NPs using pulse UV-triggered microemulsion polymerization.  \(^{195}\) Palani et al. studied the polymerization of MMA using MMA/ethylene glycol dimethacrylate/water system with acrylamide as amphiphile.  \(^{196}\) Organic NPs of cholesterol, Retinol, Rhodiarome, Rhovanil were also been reported using different microemulsion systems including AOT/heptane/water, Triton/decanol/water, and CTABr/hexanol/water.  \(^{197}\) Microemulsion method was also been used to synthesize NPs of conducting polymers including polypyrrole, polyaniline, polythiophene, etc.  \(^{198-200}\) 

2.2.3. Hydrothermal synthesis: This method is used to fabricate NPs of metal oxide, iron oxide and lithium iron phosphate keeping control over the characteristics of particles by varying the properties of near or supercritical water by using different pressure and temperature conditions. It can be performed in two types of systems, the batch hydrothermal or continuous hydrothermal process. The former is able to carry out a system with the desired ratio phases while the latter allows a higher rate of reaction to be achieved at a shorter period of time.  \(^{201}\) In a chemical solution, NPs are produced from a colloidal system that consists of two or more phases (solid, liquid or gas states) of matter (e.g. gels and foams) mixed together under controlled pressure and temperature. The advantage of using this method includes the capability to synthesize a huge amount of NPs with an optimized size, morphology, composition and surface chemistry that is rationally inexpensive.  \(^{202}\) Du et al. reported the facile one-pot synthesis of Pt NPs using hydrothermal method and observed excellent
electrocatalytic activity. Ma et al. fabricated hydrophilic NaYF$_4$:Yb,Er NPs using this method and observed good bright upconversion luminescence. They suggested that these hydrophilic NaYF$_4$:Yb,Er NPs have great potential as luminescent labelling materials for biological applications. This method was also used for the synthesis of NPs of oxides of iron, nickel and copper such as Fe$_3$O$_4$, α-Fe$_2$O$_3$, NiO, CuO, etc. Hydrothermal is a facile and fast process for the synthesis of NPs of various other materials such as CoFe$_2$O$_4$, Ag, FeWO$_4$, La$_{1-x}$Sr$_x$CrO$_3$, CdS, Zr, ZnO, etc.

2.2.4. Polyol synthesis: Polyol process is the synthesis of metal-containing compounds using poly(ethylene glycol)s as the reaction medium, that plays a role of solvent, reducing agent and complexing agent at the same time, with dissolved stabilizing/protecting agents. This chemical process was used to synthesize a wide range of metal based NPs (Ag, Pt, Pd, Pr, Cu), metal oxide NPs (ZnO, Indium-tin-oxide; ITO, Gd$_2$O$_3$, Cu$_2$O), magnetic NPs and metal hybrid NPs. Xia et al. reported the synthesis of Pt NPs using polyol process at varying molar ratio of NaNO$_3$ and H$_2$PtCl$_6$. They also reported the tailoring of size, morphology and crystallinity of NPs by slowing down the reduction of Pt(II) and Pt(IV) species (achieved by ethylene glycol) mediated by the varying concentration of nitrate ions in the solution which have a strong tendency to form stable complex with Pt(II) and Pt(IV). Moon et al. described the synthesis of spherical Cu NPs (Diameter ~ 45±8 nm) in the presence of poly(vinylpyrrolidone) as the protecting agent via polyol method. Synthesis of spherical Ag NPs with controlled size (17 nm ± 2 nm) and high monodispersity was established using the polyol process. They analyzed the effect of two varied polyol synthesis strategies, one dealing with the heating of the reaction medium having the precursor dissolved in it and other with heating the reaction medium receiving the precursor slowly using the injection method, on the NPs characteristics. Smaller silver NPs with more uniform size were obtained using the precursor injection method as compared to the precursor heating method.

Xia et al. reported the synthesis of polycrystalline NPs of Cu$_2$O using copper nitrate, ethylene glycol and poly(vinyl pyrrolidone) as the precursor, reducing agent and the stabilizing agent, respectively. Morphological transformation of these spherical Cu$_2$O NPs to nanocubes was observed by introducing NaCl in the reaction mixture which is explained by the fact that the added Cl$^-$ ions play a role in retarding the reduction kinetics, stabilizing the {100} plane of the Cu$_2$O crystals that inducing the formation of single-crystal nanocubes, which can grow in size via Oswald ripening. Low temperature synthesis of ITO NPs was carried out by Jeyadevan and coworkers in polyols and alcohols including ethylene glycol, trimethylene
glycol and 1-heptanol. The NPs characteristics were found to be influenced by the type of polyol used as the reaction medium. In another work, Moon et al. reported the preparation of ZnO NPs by polyol method and showed that the amount of water and way of precursor addition affects the characteristic of the NPs immensely. Similarly, the polyol synthesis method was also used to synthesize NPs of Gd$_2$O$_3$ and Pr$_6$O$_{11}$.

![Diagram](image)

**Fig. 10** The morphology evolution of iron oxide NPs in polyol processes using two different polyols. Adapted from ref. 225.

Polyol process was also employed for developing low temperature strategies to synthesize magnetic NPs of magnetite, Fe$_3$O$_4$, etc. Cheng *et al.* have very nicely described the effect of two different polyols (ethylene glycol; EG and 1,2-propylene glycol; PG) having different reduction capability on the morphology evolution of the Fe$_3$O$_4$ magnetic NPs (Fig. 10). Due to the higher reductive ability of PG, there is faster formation and growth rate for the NPs leads to Fe$_3$O$_4$ aggregation and generation of more misaligned mesocrystals. In the mesocrystal, the neighbouring NPs merge together and finally integrate into a single crystal with a porous structure during the Ostwald ripening process. However, due to relatively lower NP formation rate in EG, most of the particles got enough time to self-assemble along
the same orientation in the cluster. Each cluster, with its highly ordered arrangement of NPs, can be regarded as a single mesocrystal. And this mesocrystal can retain its secondary structure owing to the lower surface energy. This one-pot facile chemical process can also be used to design hybrid metal NPs including CoSb$_3$, FeCo, etc.\textsuperscript{226,227}

2.2.5. Chemical vapour deposition (CVD) & chemical vapour synthesis (CVS): CVD is a process, which is often used for the deposition of solid films from vapour phase via chemical reactions occurring at very high temperature condition. Thin films produced by CVD process at certain conditions also contain ultrafine particles. Hence, the synthesis of NPs can also be possible by this method if the CVD system is modified/optimized for the conditions given below:

i) High temperatures (in hot wall reactors)

ii) High supersaturations (high partial pressure of monomers at a low vapor pressure of the bulk solid)

iii) Long residence times (low gas flows or long reactors)

iv) Small substrates.

Hence, when the process conditions adjusted in such a way that CVD process produces NPs instead of thin solid films, then the modified process is called as CVS. The process CVS also termed as chemical vapor reaction (CVR), chemical vapor precipitation (CVP) and chemical vapor condensation (CVC). In this synthesis method, the precursors that exist in the three different states (solid, liquid, gas) are produced in the form of vapour in the reactor under an environment that requires particles to undergo the process of nucleation. In addition, CVS forms multi-component or doped NPs by the use of a variety of precursors. For instance, erbium (Er) had been incorporated into Si NPs by the use of precursors such as the organometallic erbium and disilane compound, and composite NPs are produced by encapsulating a material within the other (e.g. silicon tetrachloride in reaction with sodium chloride that result in the formation of sodium chloride-encapsulated silicon particles).\textsuperscript{228} CVS method has been extensively utilized for the synthesis of NPs using a variety of materials including ZnO, WS$_2$, iron oxide, copper and silicon oxides, Al doped ZnO, fluorine doped tin oxide, Cr doped ZnO, etc.\textsuperscript{229-236} Polarz \textit{et al.} synthesized size selected ZnO NPs (6-30 nm) by CVS process using the tetrameric alkyl-alkoxy zinc compound [CH$_3$ZnOCH(CH$_3$)$_2$]$_4$ as the precursor, which was chemically transformed into ZnO through gas-phase reactions.\textsuperscript{229} Hartner \textit{et al.} synthesized Al doped ZnO (AZO) NPs by CVS process using diethylzinc (DEZn) and triethylaluminum (TEAl) as precursors for Zn and Al,
respectively, while He gas was chosen as carrier gas.\textsuperscript{230} Oxygen gas with constant flow rate of 1000 sccm was inserted in alumina tube with an inner diameter of 19 mm, and temperature and pressure of the system was kept at 1073 K and 20 mbar, respectively. With these conditions, high quality AZO NPs were synthesized and their particle size was found to decrease with increasing the Al concentration. In comparison to other processes, CVS process has following advantages for the synthesis of AZO NPs:

i) In contrast to wet synthesis, CVS process allows high percentage of Al doping in ZnO.

ii) CVS process can produce NPs at higher growth rate, leading to reduction of the cost of the material.

iii) CVS produced AZO NPs with high crystallinity.

iv) Unlike other chemical process, CVS process avoids the use of additional step which is the removal of organic contaminant after the synthesis.

Jin \textit{et al.} reported the synthesis of Cr doped ZnO (CZO) NPs using CVS process.\textsuperscript{231} Their CVS system was splitted into two zones for evaporation and subsequently pyrolysis of the precursors. Zinc acetylacetonate (Zn(acac)\textsubscript{2}) and chromium acetylacetonate (Cr(acac)\textsubscript{3}) were used as precursors for the synthesis of CZO NPs. These precursors were evaporated from alumina boat at low temperature first zone and then transported through He carrier gas into high temperature second zone for pyrolysis and oxidizing at 1173 K. It should be noted that the temperature in first zone was kept at 493 K for pure ZnO NPs, and changed to 423 K for CZO NPs. The crystallographic and microscopic analysis revealed that the size of pure ZnO NPs was ~ 19 nm, which was decreased to ~ 6 nm in case of CZO NPs with Cr concentration of 6 at.%. Suffner \textit{et al.} demonstrated the synthesis of fluorine doped SnO\textsubscript{2} (FTO) NPs using CVS method.\textsuperscript{232} Tetramethyltin (TMT) and Difluoromethane (DFM) were used as the precursors, while oxygen and helium were used as the reactive and carrier gases, respectively, for the synthesis of FTO NPs. The flow rate of TMT, oxygen and helium was kept constant at 25 sccm, 1500 sccm and 150 sccm, respectively, while it was changed from 0 to 1.5 sccm for DFM in order to synthesize pure tin oxide (TO) NPs and FTO NPs. The chemical reactions for the synthesis of these NPs occurred at temperature and pressure of 1200 °C and 15 mbar, respectively. When synthesized at DFM flow rate of 1.5 sccm, size of FTO NPs were found to be in the range of 3 to 10 nm. The synthesized ZnO, AZO, CZO and FTO NPs can find wide electronic and optoelectronic applications.

Using atmospheric pressure CVS (APCVS) process, Lahde \textit{et al.} reported the synthesis of Cu NP composite in an amorphous silicon dioxide (a-SiO\textsubscript{2}), where Cu NPs were either coated with a-SiO\textsubscript{2} or embedded in a-SiO\textsubscript{2} matrix.\textsuperscript{233} In order to prepare NP composite structure
using APCVS, first step was to synthesize metal-organic precursor. They synthesized metal-organic precursor \( \text{[CuN(SiMe}_3)_2]_4 \), also called as Cu(I), using standard process. The ARCVS system, which was used for synthesis of NP composite, included different zones such as:

i) Precursor feeding zone

ii) Heating zone

iii) Dilution and cooling zone

iv) Sampling and particle collection zone

Using a heat bath, the synthesized Cu(I) precursor was vaporized at temperature between 100 and 250 °C. The pre-heated (100-250 °C) pure nitrogen or a mixture of hydrogen (10%v) and nitrogen was used as carriers, while the flow rate was kept to be 0.3 L/min. After this, precursor vapor was transported to heating zone of system and mixed with nitrogen at flow rate of 1.8 L/min at temperature 800 °C. However, at the exit point of the heating zone the aerosol was diluted with nitrogen at the flow rate of 10.4 L/min to quench the particle. Finally, the particle sampling and collection was done from the gas phase. Inclusion of hydrogen in nitrogen gas significantly affected the structure of NP composite. In pure nitrogen atmosphere the Si/SiO\(_2\) protective coating was formed around the Cu NPs. While in mixed hydrogen/nitrogen environment, the Cu NPs were embedded in SiO\(_2\) matrix. Schematic showing the mechanism of formation of NP composite in pure nitrogen and mixed nitrogen/hydrogen atmosphere is depicted in Fig. 11.

Further Lee et al. reported the fabrication of fullerene-like WS\(_2\) NPs using CVS process.\(^{234}\) The W(CO)\(_6\) was used as precursor for tungsten (W), which was decomposed in the temperature between 420 and 1000 °C. The evaporation temperature was varied from 80 to 110 °C at flowing He gas with varied flow rate from 2000 cc/min up to 4000 cc/min. In order to form WS\(_2\) NPs, the decomposition of tungsten hexacarbonyl W(CO)\(_6\) was performed in the presence of sulphur vapour with their partial pressure varied from 0.02 to 0.04 atm. Finally, at optimized conditions, the CVC process produces fullerene-like WS\(_2\) NPs having mean particle size in the range from 20 to 70 nm.

CVS process was also used widely for the synthesis of iron oxide NPs of mean particle size from 80-90 nm and various other metal oxide NPs.\(^{235,236}\)
**Fig. 11** Schematic representation of formation of Cu-Si/SiO\(_2\) NPs by APCVS process. Reproduced with permission from ref. 233.

### 2.2.6. Plasma enhanced chemical vapour deposition (PECVD):

Plasma enhanced chemical vapour deposition, also called as plasma assisted chemical vapour deposition (PACVD), is a popular CVD process which is widely used for the deposition of thin films. PECVD process can also be used for the synthesis of NPs. As the name suggests, plasma enhances the chemical reactions for the synthesis of thin films and NPs. PECVD unit mainly includes: 1) vacuum processing system, 2) gaseous precursors, 3) power supply (AC or DC) and 4) heater. Unlike usual CVD process, the synthesis of thin films and NPs by PECVD takes place at comparatively lower temperature. As the plasma is partially ionized gas, ionized species and radicals participate in growth of thin films and NPs. Synthesis of NPs of several types of materials was demonstrated by PECVD. Shimada *et al.* reported the synthesis of gallium nitride (GaN) NPs by microwave plasma enhanced chemical vapour deposition (MPECVD).\(^{237}\) Schematic of the MPECVD system is shown in Fig. 12a. For the synthesis of GaN NPs, trimethylgallium (TMG) was used as a precursor for Ga, while nitrogen and ammonia (NH\(_3\)) were used for nitridation. Various parameters such as antenna length, power
and operating pressure were varied to optimize the input parameters. Using $5 \times 10^{-4}$ vol. % TMG at operating pressure of 20 kPa, the decrease in size of GaN NPs with increase in microwave power was demonstrated. Average particle size of 7.3, 5.6 and 5.2 nm was achieved using microwave powers of 800, 1000, 1200W, respectively.

**Fig. 12** (a) Schematic of MPECVD system and (b) possible mechanism of growth of NPs at low and high powers. Reproduced with permission from ref. 237.

The mechanism involved in the synthesis of NPs of varied sizes is depicted in Fig. 12b. At low input power, the plasma reactivity remains low causing slow chemical reaction rate and hence, results in the formation of monomers. Because of slow reaction rate, relatively small number of nuclei are formed and at the same time significant amount of unreacted precursor molecules exit. When large numbers of unreacted species condense and react with relative small number of nuclei, the size of the resulting particles becomes larger. However, at higher power the plasma reactivity and chemical reaction got enhanced, leading to formation of more nuclei and disappearance of unreacted species. Under this condition the coagulational growth can be prevented and formation of small size NPs would be possible. Thus, the size of GaN NPs can be efficiently controlled by the input power. Dwivedi et al. used PECVD method to synthesize hard carbon nanoparticle based films by tuning the self-bias, introducing different doping strategies and using metal/carbon bi- and multilayer designs; where nitrogen was used as foreign atoms while Ti, Cu and Ag were used as metal layer in
Plasma Pretreatment

Pretreatment Time

Designed carbon nanoparticles films showed excellent electrical, field emission, optical and mechanical properties with potential applications in the field of emission devices, protective coatings, solar cells, etc.

The synthesis of carbon NPs and hydrogenated Si NPs was also reported by MPECVD and radio frequency PECVD (RF-PECVD) techniques, respectively. Yun et al. demonstrated the synthesis of silica NP array on flexible polymer substrate using PECVD process involving following two steps:

i) Pre-plasma treatment of polymer surface followed by

ii) Coating of silica NPs array

Fig. 13 Plasma pre-treatment for creation of polymer protrusions. (a) Schematic diagram and (b–d) cross section as well as (e–g) plane view FE-SEM images showing the effect of treatment time on aspect ratio of formed polymer protrusions. (h) AFM line profiles and (i) the corresponding quantitative data of the protrusions (h and d) with plasma treatment times. Reproduced with permission from ref. 247.

To perform plasma pre-treatment, the polymer substrate (polyethylene terephthalate; PET) was placed on electrode housed in vacuum system (base pressure of 6.7 Pa). The plasma pre-treatment was performed on PET at constant RF power of 200 W (1.1 W cm$^{-2}$), working pressure of 22.7 Pa and Ar flow rate of 50 sccm, while the plasma treatment time was varied. The Ar plasma pre-treatment of the PET surfaces led to the formation of polymer protrusions...
due to morphological modifications of polymer chains as shown in Fig. 13. Height of protrusions (h) and the distance between neighbouring protrusions (d) was found to increase continuously with increasing pre-treatment time from 1 min to 7 min (Fig. 13). Subsequently silica NPs are deposited on pre-treated polymer surfaces by plasma polymerization of hexamethyldisiloxane (HMDSO). First in evaporator, the vapors of HMDSO are formed by thermal evaporation of precursor liquid HMDSO at a heating temperature of 130 °C and then they are mixed with O₂ and Ar gases at a flow rate ratio of HMDSO: O₂: Ar = 1.8: 10: 100 before entering into the synthesis chamber. Finally silica NPs are grown having O: Si atomic ratio of 1.9-2.0. Fig. 14 shows the growth of silica NPs on pre-treated polymer surfaces.

![Fig. 14](image)

**Fig. 14** (a) Schematic showing the role of pre-treated polymer surfaces on the synthesis of silica NPs. (b–d) Cross-sectional and (e–g) plane view FE-SEM images of silica NPs formed on pre-treated polymer surfaces. (h–j) High magnification cross-sectional FESEM images of silica NPs formed on pre-treated polymer surfaces for different treatment times of (h) 1 min, (i) 3 min, and (j) 7 min. Reproduced with permission from ref. 247.

Of note, plasma pre-treatment time of polymer surfaces significantly affects the growth of NPs. As the polymer protrusions reduce with increasing plasma pre-treatment time, the density of the formed silica NPs also reduces with increasing plasma-pre-treatment time. The globular silica NPs synthesized by this method showed excellent antireflective property.
2.2.5. Chemical methods for the synthesis of graphene oxide nanoparticles: Recently, graphene oxide nanostructures received wider attention owing to their multiple applications including drug delivery, catalysis, cell therapy, environmental monitoring, biosensors, field emission transistors, clean energy devices, etc.\textsuperscript{248} Considering their broad range applicability, this section on the synthesis of graphene oxide nanoparticles (GON) has been included here to enhance the comprehensiveness of the present review. In most of the reports available, modified Hummers methods have been used to synthesize GON of varied sizes.\textsuperscript{249} Zhao et al. adopted modified Hummers method to synthesize nearly spherical GON with nearly monodisperse size of 43.36 ± 8.42 nm from graphite powder.\textsuperscript{250} Authors have further functionalized the GON with cytamine modified PEGylated alginate to design drug delivery platform for tumour microenvironment-responsive triggered release of doxorubicin. Jana et al. synthesized Tris (nitrilo Tris-acetic acid), Tris-NTA, functionalized GON using modified Hummers method with average particle size of 213 nm which can reconstitute kinesin mediated intracellular cargo transport and can deliver multiple proteins and therapeutic antimitotic dodecapeptide peptides into the cancer cell.\textsuperscript{251} Authors used similar strategy to fabricate dual functionalized GON employing Tris-NTA and biotin for cellular delivery of oligohistidine- and biotin-tagged biomolecules.\textsuperscript{252} Lee et al. reported the synthesis of GON with the size of 30 nm by first exfoliating graphite into graphene oxide using modified Hummer method followed by their ultrasonication.\textsuperscript{253} These tiny GON have been shown to be applicable as non-bleaching optical probe for two photon luminescence imaging and cell therapy.

2.3. Bio-assisted methods for the synthesis of nanoparticles

Bio-assisted methods, biosynthesis or green synthesis provides an environmentally benign, low-toxic, cost-effective and efficient protocol to synthesize and fabricate NPs. These methods employ biological systems like bacteria, fungi, viruses, yeast, actinomycetes, plant extracts, etc. for the synthesis of metal and metal oxide NPs. Bio-assisted methods can be broadly divided into three categories:

i) Biogenic synthesis using microorganisms

ii) Biogenic synthesis using biomolecules as the templates

iii) Biogenic synthesis using plant extracts
i) Biogenic synthesis using microorganisms

Prokaryotic bacteria, actinomycetes, fungi, algae and yeast are extensively used as bioreactors for the synthesis of NPs. Enormous scientific efforts were made to develop this strategy of producing a variety of NPs (Ag, Au, Pd, TiO$_2$, CdS, etc.). Microorganisms grab target ions from their environment and then turn the metal ions into the element metal through enzymes generated by cellular activities. This synthesis can be classified into intracellular and extracellular depending upon the location of NP synthesis. The intracellular method involves transporting of metal ions into the microbial cell to form NPs in the presence of enzymes. The extracellular synthesis of NPs involves trapping of metal ions on the surface of the cells and reducing ions in the presence of enzymes. Bacteria utilize a number of anionic functional groups, proteins and enzymes, reducing sugars, etc. in bacterial biomass to reduce interacting metal ions. Bacterium *S. aureus* was used for the extracellular synthesis of bioactive Ag NPs with antimicrobial activities. Boon *et al.* described non-enzymatic production of Ag NPs using biomatrix of the bacteria both as the reducing and the capping agent. The presence of anionic functional group rich cell wall around the gram-positive bacteria constituting of peptidoglycan, teichoic acids, lipoteichoic acids, proteins, and polysaccharides acts as sites for the biosorption and subsequent reduction of silver cations. Lactobacillus bacterium was used as the biofactories for the synthesis of TiO$_2$ NPs with size ranging from 40-60 nm. Gunasekaran *et al.* reported the isolation of new hyper metal resistant *Bacillus cereus* PGNI bacterial strain that form intracellular crystalline Ag NPs. Bacterial system was also employed to synthesize semiconducting CdS nanocrystals (wurtzite crystal structure). In another interesting report, authors explored the potential of marine bacteria for the production of AuNPs. Fungal-mediated green chemistry approach for the synthesis of NPs has many advantages like higher bioaccumulation, economic viability, easily scaled up synthesis method due to simple downstream processing and biomass handling. In this context, Mukherjee *et al.* demonstrated a fungal-assisted biological method to synthesize silver NPs using fungus *Verticillium*. Exposure of the fungal biomass to the aqueous Ag$^+$ ions leads to their intracellular reduction to silver NPs of dimension 25 ± 12 nm. Microscopic investigation reveal mycelia surface as the predominant site for the NPs synthesis which is mediated by the enzymes available on the mycelia cell wall. Biosynthesis of Ag NPs was also reported using *Aspergillus terreus* fungus and herein the speculated mechanism is reported to be NADH dependant enzyme catalyzed extracellular reaction process. There are number of other
reports available on the biosynthesis of Ag NPs using versatile fungal species including *Pleurotus Ostreatus*, *Aspergillus flavus*, *Bryophilous Rhizoctoni*, etc. Owaiais and co-workers synthesized Au NPs (20-40 nm) employing *candida albicans* mediated bioreduction strategy. The mechanism of fungal biosynthesis of Au NPs is very well explained by Kitching et al. Bansal et al. demonstrated the biotransformation of amorphous silica in cheap agro-based rice husk to nanocrystalline silica (2-6 nm). There are several reports on the NPs biosynthesis of meganitite, TiO$_2$ and ZrO$_2$ using various fungal species.

Yeast and actinomycetes also established their applicability for the bio-inspired synthesis of NPs. Yeast is an eukaryotic microorganisms belonging to the fungi kingdom. Kowshik et al. reported bulk extracellular synthesis of silver NPs (2-5 nm) using silver tolerant yeast strain MKY3. Kandasamy et al. shown the applicability of *Hanensula anomala* species to reduce gold salt generating AuNPs. Yarrowia lipolytica was also reported as efficient reducing agent for AuCl$_4$ to produce gold NPs. Carboxyl, hydroxyl and amide groups on the cell surface are anticipated to be responsible for AuNPs synthesis. Yeast was also reported to be useful in the synthesis of Cd and PbS NPs using *Candida glabrata* and *Rhodosporidium diobovatum* species, respectively. Enriched with various enzymes, actinomycetes have been for the reduction of AuCl$_4^-$ to generate Au NPs (8 nm) by *Thermomonospora sp*. Torres-Chavolla et al. reported the synthesis of comparatively larger monodispersed Au NPs (30-60 nm) in the presence of *Thermomonospora curvata*, *Thermomonospora fusca* and *Thermomonospora chromogena* species. Table 4 lists various NPs synthesized using a variety of microorganisms.

Table 4 Various microorganisms reported to be involved in the synthesis of different types of NPs.

<table>
<thead>
<tr>
<th>Microorganism</th>
<th>Type of Nanoparticles</th>
<th>Size of Nanoparticles</th>
<th>Mode of Synthesis</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>BACTERIA</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td><em>Staphylococcus aureus</em></td>
<td>Ag NPs</td>
<td>160-180 nm</td>
<td>Extracellular</td>
<td>255</td>
</tr>
<tr>
<td><em>Bacillus cereus</em></td>
<td>Ag NPs</td>
<td>4-5 nm</td>
<td>Extracellular</td>
<td>258</td>
</tr>
<tr>
<td><em>Marinobacter pelagius</em></td>
<td>Au NPs</td>
<td>2-10 nm</td>
<td>Extracellular</td>
<td>260</td>
</tr>
<tr>
<td><em>Actinobacter sp.</em></td>
<td>Au NPs</td>
<td>50-500 nm</td>
<td>Extracellular</td>
<td>279</td>
</tr>
<tr>
<td><em>Enterobacter sp.</em></td>
<td>Hg NPs</td>
<td>2-5 nm</td>
<td>Intracellular</td>
<td>280</td>
</tr>
<tr>
<td><em>Lactobacillus</em></td>
<td>TiO$_2$ NPs</td>
<td>40-60 nm</td>
<td>Extracellular</td>
<td>257</td>
</tr>
<tr>
<td><em>Escherichia coli</em></td>
<td>CdS QDs</td>
<td>2-5 nm</td>
<td>Intracellular</td>
<td>259</td>
</tr>
<tr>
<td><em>Escherichia coli</em></td>
<td>CdTe QDs</td>
<td>2-3.2 nm</td>
<td>Extracellular</td>
<td>281</td>
</tr>
<tr>
<td><strong>FUNGI</strong></td>
<td></td>
<td></td>
<td></td>
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</tr>
<tr>
<td><em>Verticillium</em></td>
<td>Ag NPs</td>
<td>25±12nm</td>
<td>Intracellular</td>
<td>261</td>
</tr>
<tr>
<td><em>Aspergillus terreus</em></td>
<td>Ag NPs</td>
<td>1-20 nm</td>
<td>Extracellular</td>
<td>262</td>
</tr>
<tr>
<td><em>Bryophilous Rhizoctoni</em></td>
<td>Ag NPs</td>
<td>20-50 nm</td>
<td>Extracellular</td>
<td>263</td>
</tr>
<tr>
<td><em>Aspergillus flavus</em></td>
<td>Ag NPs</td>
<td>8.92 ± 1.61 nm</td>
<td>Extracellular</td>
<td>264</td>
</tr>
<tr>
<td><em>Pleurotus ostreatus</em></td>
<td>Ag NPs</td>
<td>8-50 nm</td>
<td>Extracellular</td>
<td>265</td>
</tr>
</tbody>
</table>
2.3.2. Biomolecules as templates to design nanoparticles

Various biomolecules like nucleic acids, membranes, viruses and diatoms were used as templates to synthesize NPs. DNA is widely known as an excellent biomolecular template that has a strong attraction with transition metal ions. It was shown that DNA hydrogel could be made and crosslinked before incorporating transition metal ions (e.g. gold, Au(III) metal ions) to DNA macromolecules that eventually lead to the formation of Au NPs. The process involves a reduction of Au(III) leading to the formation of Au atoms and metal clusters that develop into Au NPs on the chain of DNA.\textsuperscript{284} Kundu et al. reported the synthesis of wide variety of NPs and nanoparticles assemblies using DNA as the templates or organic scaffolds.\textsuperscript{285-300} They accounted a DNA-mediated strategy to synthesize highly stable wire-like clusters of Ag NPs with average particle size of the 17 ± 3 nm and inter-particle gap of ~1.7 ± 0.2 nm for their potential applicability as ultrasensitive SERS substrates.\textsuperscript{285} These Ag NPs assemblies grown over DNA chains have also been shown to possess good catalytic activity towards the reduction of aromatic nitro compounds.\textsuperscript{286} An electroless, photolytic, DNA-mediated method was described to synthesize gold nanoclusters (10-40 nm) and continuous long nanostructures (Diameter: 40-70 nm) with the nanostructures exhibiting resistivity equivalent to the pure metals.\textsuperscript{287} In another report, authors demonstrated synthesis strategy to design shape-sensitive ZnO nanoparticles with various morphologies (wire-like, ~150 ± 15 nm; flake like, ~80 ± 10 nm; flower like, ~350 ± 50 nm) using DNA as bio-template for dye sensitized solar cell applications.\textsuperscript{288} Attempts were also made to design organosols of Os NPs and β-MnO\textsubscript{2} NPs using DNA as scaffolds via homogenous reduction route with their promising applications in catalytic hydrogenation and oxidative
polymerization of pyrrole, respectively.\textsuperscript{289,290} DNA was also been used as scaffolds to grow self assembled NiWO$_4$, ZnWO$_4$ and MnWO$_4$ nanoparticles with different morphologies.\textsuperscript{291-293} Fast, electroless, UV-irradiation-/microwave-assisted methods with DNA as educing as well capping agent was also been reported to synthesize continuous, electrically conductive nanowires of Au, Pd and CdS which can be used as the building blocks for functional nanodevices, miniaturized computers, sensors and optoelectronic applications.\textsuperscript{36,294-297} Table 5 compiled self-assembled nanoclusters of versatile nanoparticles synthesized by using DNA as bio-template or scaffold.

**Table 5** Self-assembled nanoclusters of different nanoparticles synthesized using DNA as template or scaffold.

<table>
<thead>
<tr>
<th>Self Assembled Nanocluster of</th>
<th>Shape of Nanocluster</th>
<th>Dimensions</th>
<th>Applications</th>
<th>References</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ag NPs</td>
<td>Wire Like</td>
<td>$\phi_P = 17 \pm 3$ nm, $\text{IPD} = 1.7 \pm 0.2$ nm</td>
<td>As Ultrasensitive SERS substrate</td>
<td>285</td>
</tr>
<tr>
<td>Os NPs</td>
<td>Wire Like</td>
<td>$\phi_P \sim 2 \pm 0.5$ nm, $\phi_w \sim 290 \pm 20$ nm</td>
<td>Catalysis and SERS</td>
<td>298</td>
</tr>
<tr>
<td></td>
<td>Honeycomb Like</td>
<td>$\phi_P = 1.5 \pm 0.2$ nm, $\phi_w = 400$ nm</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Os NPs (Organosol)</td>
<td>Wire like</td>
<td>$\phi_P = 2.6 \pm 0.2$ nm, $L = 0.54 \pm 0.03$ µm</td>
<td>Catalysis and SERS</td>
<td>289</td>
</tr>
<tr>
<td></td>
<td>aggregated wires</td>
<td>$\phi_P = 1.2 \pm 0.2$ nm, $L = 8 \sim 10$ micron</td>
<td></td>
<td></td>
</tr>
<tr>
<td>ZnO NPs</td>
<td>Wire-Like</td>
<td>$\phi_P = 150 \pm 15$ nm, $L = 1 \sim 2$ µm</td>
<td>Catalysis and Dye-Sensitized Solar Cells</td>
<td>288</td>
</tr>
<tr>
<td></td>
<td>Flower-Like</td>
<td>$\phi_P = 350 \pm 50$</td>
<td></td>
<td></td>
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<td></td>
<td>Flake-Like</td>
<td>$\phi_P = 80 \pm 10$ nm</td>
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<tr>
<td>$\beta$-MnO$_2$ NPs</td>
<td>Wire-Like</td>
<td>$L = 1.9 \pm 0.2$ nm, $\phi_P = 35 \pm 5$ nm</td>
<td>Catalysis and supercapacitor</td>
<td>299</td>
</tr>
<tr>
<td></td>
<td>Flake-Like</td>
<td>$L = 275 \pm 25$ nm, $\phi_P = 25 \pm 5$ nm</td>
<td></td>
<td></td>
</tr>
<tr>
<td>TiO$_2$ NPs</td>
<td>Wire Like Cluster (Large)</td>
<td>$\phi_P = 15 \pm 5$ nm, $\phi_C = 180 \pm 20$ nm</td>
<td>Supercapacitor and dye sensitized solar cells</td>
<td>300</td>
</tr>
<tr>
<td></td>
<td>Wire Like Cluster (Small)</td>
<td>$\phi_P = 10 \pm 2$ nm, $\phi_C = 40 \pm 5$ nm</td>
<td></td>
<td></td>
</tr>
<tr>
<td>NiWO$_4$ NPs</td>
<td>Chain Like (Small)</td>
<td>$L = 2 \pm 0.2$ µm, $\phi_C = 175 \pm 15$ nm, $\phi_P = 20 \pm 5$ nm</td>
<td>Catalysis and Supercapacitor</td>
<td>291</td>
</tr>
<tr>
<td></td>
<td>Chain Like (Large)</td>
<td>$L = 3.4 \pm 0.2$ µm, $\phi_C = 245 \pm 15$ nm, $\phi_P = 26 \pm 4$ nm</td>
<td></td>
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</tbody>
</table>
Likewise, owing to the presence of ultra-fine pores in their structure, biological membranes were also utilized as templates to design NPs. Rubber membrane made from the *Hevea brasiliensis* trees was used as a preservative in the production of Au NPs synthesized at the surface of membranes through the reduction of Au(III) in a solution at 80°C for various reaction times. Alternatively, a uniform size and morphology of NPs could be obtained by the use of viruses as they contain hollow spaces in the center that can be used as the template for production of particles. Diatoms such as *Amphora-46* was also reported to synthesize polycrystalline Ag NPs of size ranging 20 to 25 nm under the exposure to light. This process is done by the utilization of its extract and silver nitrate (AgNO₃), in which the presence of a pigment known as fucoxanthin reduces the Ag ion that leads to further production of Ag NPs.

### 2.3.3. Plant extracts for nanoparticles synthesis

Biosynthesis of NPs using plant extracts or plant biomass is one of the very effective, rapid, clean, non-toxic and eco-friendly methods. This method has been utilized predominantly to synthesize NPs of noble metals, metal oxides, bi-metallic alloys, etc. Yun *et al.* have adequately demarcated various plant biometabolites that could help in the preparation of NPs based on their valuable role as reducing agents and capping agents. The mechanism for the phytosynthesis of metal NPs is given in Fig. 15.
Reducing Agents

Amino acids, citric acid, dehydrogenase, flavenoids, functional groups (alcohols, amines, carboxyl acids, ketones and sulfhydryl), heterocyclic compounds, hydrogenase, intracellular CO2, membrane proteins, NADH reductase, peptide, phenolics, polyphenols, polyols, protein, quercetin, reducing sugars, saponin, secondary metabolites, sequiterpenes, tannic acid, tartaric acid, terpenoids

Capping Agents

Citric acid, extracellular proteins, enzymes, functional groups (alcohols, aldehydes, amines, carboxylic acid, ketones, and sulfhydryl), peptides, phyllanthin, prolines, tartaric acid, tannic acid

Fig. 15

Mechanism for the biogenic synthesis of metal NPs using plant extract. Reproduced with permission from ref. 305.

Kinetics of phytosynthesis of NPs is comparatively much higher to that of other biosynthesis methods and sometime equivalent to the rate of chemical routes. Shanker et al. reported the preparation of gold nanotriangles by treating the lemongrass leaf extract with the aqueous AuCl ions. Similarly the leaf extract of different plants like Tamarindus indica, Aloe vera, Emblica Officinalis, etc. were reported to be useful for designing Au NPs. Few nanometer sized Pd NPs and Pt NPs were prepared using the extract taken out from various parts of different plants. Shanker et al. also reported highly concentrated Ag NPs obtained from the leaf extract of Azadirachta indica and from the fruit extract of Embalica officinalis.

Li et al. and Leela et al. used the leaf extract of Aloe vera, Capsicum annum and Helianthus annus for the efficient synthesis of Ag NPs. Extracellular production of Cu NPs was carried out using stem latex of Euphorbia nivulia (medicinal plant) which were found to be stabilized by the peptides and terpenoids present within the latex. In a very interesting report, Seraphin et al. demonstrated the synthesis of In2O3 NPs (5L50 nm) using plant extract from Aloe vera.

Furthermore, Hou et al., synthesized wurtzite ZnO NPs from...
Zn-hyper-accumulator plant *Sedum alfredii* with mean size of 53.7 nm. Ascencio *et al.* reported the bio-assisted synthesis of iron-oxide NPs by using the biomass of *Medicago sativa* (alfalfa) plant. Glutathione, an antioxidant tripeptide in plants, animals, fungi and archaea, was reported to design well-ordered aggregates of Au NPs by favouring the inter-particle interactions. Kundu and Nithiyanantham also demonstrated the in-situ fabrication of shape-selective silver nanostructures using curcumin as stabilizing and reducing agent.

### 2.4. Key Applications of Different Nanoparticles

NPs have wide range of applications from electronic, optoelectronic, magnetic, data storage, energy and energy storage, nanomedicines, bioimaging, etc. We compiled the key applications of commonly used NPs in Table 6.

**Table 6** Key Nanoparticles and their important Applications in various scientific fields

<table>
<thead>
<tr>
<th>Nanoparticles</th>
<th>Possible Key Applications</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ag</td>
<td>Nanomedicines: antibacterial agent (healthcare, food technology, textile coating, environmental cleaning, water disinfection)(^{321}), wound healing,(^{322,323}) cancer therapeutics(^{324}). Diagnosis: designing highly sensitive biosensor electrodes/chips(^{325}), biological tagging for quantitative detection(^{326}). Solar cells(^{327}), electroluminescent displays(^{328}), optical sensors, surface enhanced Raman spectroscopy(^{329,99,285}). To generate eco-friendly electrical contacts for electrical devices(^{330}). Supercapacitors(^93). Catalysis(^{331,286}).</td>
</tr>
<tr>
<td>Au</td>
<td>Nanomedicines: drug and gene delivery(^{322}), hyperthermia-assisted cancer phototherapy(^{333,334}), photodynamic therapy(^{335}). Biosensors: highly sensitive optical(^{336}), electrochemical(^{337}) and plasmon resonance biosensors(^{338}). Sensors(^90). <em>Ex-vivo</em> and <em>In-vivo</em> bio-imaging(^{339}): two photon luminescence imaging, magnetic resonance imaging, photoacoustic tomography, Raman imaging, etc. Antibacterial agents(^340). Solar Cells(^341), Energy Storage Devices(^90).</td>
</tr>
<tr>
<td>Fe</td>
<td>Biomedical applications: Biosensors,(^{42}) MRI contrast enhancement,(^{343}) hyperthermia(^{344}). Magnetic and electrical applications: Magnetic recording media, soft magnetic materials, motors. Catalytic applications</td>
</tr>
<tr>
<td>Pt</td>
<td>Biomedical applications: affinity probe for the detection of small biomolecules,(^{445}) Catalytic nanomedicines,(^{346}) biosensors(^347). Catalysis(^348). Solar Cells(^349).</td>
</tr>
<tr>
<td>Pd</td>
<td>Surface enhanced Raman scattering and electrocatalysis&lt;sup&gt;356&lt;/sup&gt;&lt;br&gt;Catalysis&lt;sup&gt;351&lt;/sup&gt;&lt;br&gt;Antibacterial Activity&lt;sup&gt;352&lt;/sup&gt;&lt;br&gt;Fuel Cells&lt;sup&gt;353&lt;/sup&gt;&lt;br&gt;Biosensors&lt;sup&gt;354&lt;/sup&gt;</td>
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<tr>
<td>------</td>
<td>----------------------------------------------------------------------------------</td>
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<tr>
<td>Ge</td>
<td>Electronic/Optoelectronic&lt;sup&gt;32&lt;/sup&gt;</td>
</tr>
<tr>
<td>CeO&lt;sub&gt;2&lt;/sub&gt;</td>
<td>Potential regenerative antioxidant&lt;sup&gt;355&lt;/sup&gt;&lt;br&gt;Therapeutic applications for reactive oxygen species (ROS)-related diseases e.g. Cancer, diabetes, arthritis, infertility, macular degeneration&lt;sup&gt;356,357&lt;/sup&gt;&lt;br&gt;Catalytic convertor for removing toxic gases&lt;sup&gt;355&lt;/sup&gt;&lt;br&gt;Solid oxide fuel cells&lt;sup&gt;355&lt;/sup&gt;&lt;br&gt;Biosensors&lt;sup&gt;358&lt;/sup&gt;&lt;br&gt;Photocatalysis&lt;sup&gt;125&lt;/sup&gt;&lt;br&gt;Antibacterial activity&lt;sup&gt;359&lt;/sup&gt;</td>
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<tr>
<td>TiO&lt;sub&gt;2&lt;/sub&gt;</td>
<td>Drug delivery&lt;sup&gt;321,368&lt;/sup&gt;&lt;br&gt;For cell delivery&lt;sup&gt;361&lt;/sup&gt;&lt;br&gt;Antimicrobial coatings&lt;br&gt;As UV filters in sunscreens, toothpaste, cosmetics, etc.&lt;br&gt;Biosensors&lt;sup&gt;363&lt;/sup&gt;</td>
</tr>
<tr>
<td>ZnO</td>
<td>Solar cells&lt;sup&gt;364&lt;/sup&gt;&lt;br&gt;Photocatalytic and hygienic coatings&lt;sup&gt;365&lt;/sup&gt;&lt;br&gt;Biosensors&lt;sup&gt;366&lt;/sup&gt;&lt;br&gt;Important additive for different cosmetics, ointments and food products.&lt;br&gt;Cure activator for rubbers of different kinds&lt;sup&gt;367&lt;/sup&gt;&lt;br&gt;Antibacterial&lt;sup&gt;65&lt;/sup&gt;</td>
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<tr>
<td>Fe&lt;sub&gt;2&lt;/sub&gt;O&lt;sub&gt;3&lt;/sub&gt;</td>
<td>Due to its excellent magnetic properties, they are being used in magnetic seals and inks, magnetic recording media, catalysts, and ferrofluids.&lt;sup&gt;368&lt;/sup&gt;&lt;br&gt;Development of immunoassays, magnetic resonance imaging contrast agents, and targeted drug delivery vehicles, as well as in magnetic hyperthermia.&lt;sup&gt;369&lt;/sup&gt;</td>
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<tr>
<td>CdS</td>
<td>As fluorescent probes&lt;sup&gt;370&lt;/sup&gt;&lt;br&gt;Optical, electrochemical and photoelectrochemical biosensors&lt;sup&gt;371,372&lt;/sup&gt;&lt;br&gt;Solar cells&lt;sup&gt;373&lt;/sup&gt;&lt;br&gt;Photoluminescence devices&lt;sup&gt;374&lt;/sup&gt;</td>
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<td>Magnetic Nanoparticles</td>
<td>Biomedical applications: Gene delivery&lt;sup&gt;377&lt;/sup&gt;, Magnetic resonance imaging,&lt;sup&gt;378&lt;/sup&gt; Magnetic carriers for bio-separation&lt;sup&gt;379&lt;/sup&gt;&lt;br&gt;Catalyst supports&lt;sup&gt;380&lt;/sup&gt;</td>
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<tr>
<td>Upconversion (Lanthanides) Nanoparticles</td>
<td>&lt;i&gt;In-Vivo&lt;/i&gt; Cell Imaging&lt;sup&gt;381&lt;/sup&gt;&lt;br&gt;Near-infrared photodynamic therapy&lt;sup&gt;382&lt;/sup&gt;&lt;br&gt;Light Nanotransducers for photoactivation applications&lt;sup&gt;383&lt;/sup&gt;&lt;br&gt;Solar Cells&lt;sup&gt;384&lt;/sup&gt;</td>
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<td>Carbon Nanoparticles</td>
<td>Cancer diagnostic and therapy&lt;sup&gt;385,386&lt;/sup&gt;&lt;br&gt;Antibacterial&lt;sup&gt;387&lt;/sup&gt;&lt;br&gt;Lithium sulfur batteries&lt;sup&gt;388&lt;/sup&gt;&lt;br&gt;Designing of fluorescent imaging probes&lt;sup&gt;389&lt;/sup&gt;&lt;br&gt;Catalysis&lt;sup&gt;62&lt;/sup&gt;&lt;br&gt;Solar Cells, Protective Coatings and Emission devices&lt;sup&gt;238-244&lt;/sup&gt;</td>
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</table>
3. Conclusions and Future Prospects

Since the inception of NPs over half a century ago, scientists are continuously exploring advanced novel methods of synthesizing NPs with the optimal size and morphology that would be beneficial for various disciplines. To engineer and fabricate an ideal size and morphology of NPs, depending upon their application/use, diverse range of physical, chemical or biological methods are already available. In this review, we provided a comprehensive overview of the commonly employed/explored strategies for the synthesis of NPs in term of their working principle, NPs designing stratagem and relevant literature. It is anticipated that this review will act as a guide article for the researchers to selectively choose particular synthesis strategy for designing the desired NPs with particular material type, size, surface properties, targeted application, etc.

Owing to its enormous technological potential, nanoparticles research is at the forefront. Being a leading nanomaterial in targeted drug delivery and nanomedicines, precise size with specified surface characteristics are the key desirables during nanoparticle synthesis. Cell toxicity, potential target, drug release profile, cost-effectiveness, etc. are the key concerns which are needed to be taken care of by the optimization of size and surface properties of nanocarriers by selecting appropriate synthesis methods. Electrospraying is growing enormously as a dry technique to design polymeric nanoparticles without using any hazardous chemicals. It provides prospects to develop nanoformulations incorporating drugs/growth factors/biomolecules with versatile structural designs like core shell NPs, hybrid NPs, composite NPs, etc., that will be useful in drug delivery, tissue engineering, biosensors, etc. Furthermore, electrospraying is still almost unexplored for the production of metal and ceramic nanoparticles. Plasmonics, optoelectronics, information and data storage, etc. are also the key areas where the synthesis strategies for the scalable production of
nanoparticles needs to be further explored and optimized which can help to develop economically viable technologies.

**Acknowledgement**

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References


130. R. Sridhar and S. Ramakrishna, *Biomatter*, 2013, **3**, e24281.


Table of Content

Various methods to synthesize diverse nanoparticles with their different applications

Methods of Nanoparticles Synthesis

- Physical Methods
  - High Energy Ball Milling
  - Inert Gas Condensation
  - Pulse Vapor Deposition
  - Laser Pyrolysis
  - Flash Spray Pyrolysis
  - Electrospraying
  - Melt Mixing

- Biological Methods
  - Microorganisms assisted Biogenesis
  - Bio-Template assisted Biogenesis
  - Plant Extracts assisted Biogenesis

- Chemical Methods
  - Sol Gel Synthesis
  - Microemulsion Technique
  - Hydrothermal Synthesis
  - Plasma Enhanced Chemical Vapor Deposition
  - Polylol Synthesis
  - Chemical Vapor Synthesis

Applications

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<tr>
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<th>Solar Cells</th>
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