



Recent Developments and Applications of Nanotechnology to Develop Novel Medical Devices, Infectious Disease Diagnosis, and its Treatment

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ABSTRACT

Increasing rate of emergency infections and novel diseases is a serious threat to livelihood and health. Pathogens in the form of virus, bacteria, fungi, and molds are responsible for antibiotic resistance which give birth to the novel diseases/ infections like COVID-19 (Coronavirus disease of 2019). Scientific research, drug discoveries, and specific treatments are not enough to mitigate these challenges. This article reviews the distinct aspects of nanotechnology against emergency infections and novel diseases, development of nanoparticles, usage, and applications for various support functions of human, animal, and plant life. This article also covers the classification, preparation method, development of silver nanomaterials and its promising antimicrobial properties as solution of current infectious environment. It also covers the synthesis of various nanomaterials, properties, challenges, and applications in various domains like biomedical, food, textile, environment, and water. It represents various gaps and associate challenges with medical devices and consumables. Coating nanomaterials are part of this review which provides a hope to develop the novel antimicrobial medical devices as an immediate need of current infectious environment. It includes recent clinical advancements, applications, various challenges and future prospectives of nanotechnology.

Keywords: Medical devices, Disinfectant, Antimicrobial(s), Nanotechnology, Silver nanoparticles, Applications.



INTRODUCTION

Emerging infections and novel diseases are badly impacting the world's health and livelihood. Over three hundred contagious diseases were reported till 2004 caused by different strains like bacteria, fungi, viruses, and protozoa. Specific ecological and environmental factors influenced the viral spread of infectious diseases¹. Recently, some novel disinfecting agents and techniques have been discovered to control some typical diseases, such as cholera, tuberculosis, and pneumonia². Reusable medical devices undergo multiple usage and reprocessing steps, causing the accumulation of organic residues, which enhance microbial growth. Removal of these resistive contaminants is a challenging process³. Several techniques are available to control this spread of microorganisms but still need to develop some fast, cost-effective, and consistent approaches⁴. To overcome these challenges, nanotechnology may be an alternate fast-growing science approach.

Nanotechnology is a science which works with a defined scale of 1 to 100 nanometers, inclusive of various domains like biology, chemistry, physics, and material science⁵. Nanotechnology includes silver nanoparticles, surface coating, nanocomposites, and Quantum dots, among others. Due to silver nanoparticles' antimicrobial properties, it acts as an effective disinfectant used for the development of silver-enabled air and liquid filters, textiles, food packaging and biomedical⁶. Surface coatings have a high potential to better the life of reusable medical devices. Improvements in coatings can help in reprocessing, disinfection, and sterilization by minimizing the corrosion, discoloration, bubble formation, and erosion of the outer surface^{7,8}. In current scenario, there are limited materials and consumables which have the potential to resolve the issue of antibacterial resistance.

As per current challenges, it is required to develop a sustainable solution to reduce the bacterial adherence probability over the medical devices and different type of consumables. Development of novel coating material by using silver nanocomposite may be a sustainable solution for these challenges. To prevent infections and diagnosis of bacterial disease, it is required to develop novel nanocomposites

as a powerful weapon in healthcare sector. This review article highlights the advancement of nanotechnology and applications in medical, water, air, and food organizations.

Various infectious challenges

There is an extensive list of diseases caused by high concentrations of microbes, for example, biofilm treatment is more resistant to "traditional biofilm" and develops in wet environments⁹. Endoscope is one recent example of infection due to Enterobacteriaceae¹⁰. Health care units are currently facing the issue of the resistant nature of some specific microorganisms against multiple antibiotics. Due to that, health care experts are unable to provide proper support to patients¹¹.

Due to limited resources and dense population, contaminants in water in microbial form, along with excess contents of chemicals which, may cause severe disease¹². In present scenario, airborne particles are encouraging the growth of microorganisms like viruses, fungi, and other bacterial strains which cause multiple diseases, for example, anthrax, SARS (severe acute respiratory system), and asthma¹³. Hospital environment is always prone to spread infection and responsible for spreading severe infections to the environment^{14,15}. Textile industries are struggling for stable, durable, sweating free and safe one^{16,17}. Silver release from food packages system needs more research to evaluate its impacts for consumers¹⁸. Initial source of contamination in food industry is the processing plant having biofilm. These biofilms sometime become resistant to the disinfection and cleaning detergents¹⁹. In hatchery rooms viruses, bacteria, bacterial toxins, fungi, spores, pesticides are the contaminants which are produced due to feed residues, feathers, feces, and epiderm²⁰.

Now a days food industries are struggling due to unavoidable bacterial, fungal, parasites and viral infections²¹. Hospital acquired infections now a days a major challenge which is causing multiple death²². There are some cases which reported lapses in reprocessing (cleaning, disinfection/sterilization) even fulfilled the requirements of applicable guidelines. Effectiveness of disinfectant may be increased based on these specific comparisons²³. These challenges can be taken care

of by following standard precautions, transmission-based precautions, and monitoring²⁴. Usage of nanosilver and nanocomposites are playing a key role in infection control²⁵. It may be a mistake by regulatory authorities if nano silver does not declare as a new chemical with its multiple properties^{26, 27}.

Proposed practices for safe usage of medical devices, infection diagnosis, and controls.

Simulated Reprocessing, cleaning validation, and sterilization.

To control microbial worldwide spread surgeons and researcher proposed few innovative techniques against infectious strains. Disinfection and sterilization technique reduced the infections significantly for different areas like hospitals, clinical areas, and biological research²⁸. Disinfection process eliminates a variety of pathogenic microorganism but are limited to bacterial spores. Multiple disinfectants were developed from time to time for specific applications. Disinfectants were used as liquid chemicals or wet pasteurization to inhibit the spread of microorganism over a surface²⁹. Cleaning and disinfection of environmental surfaces plays a significant role in effective infection prevention program for example pulsed-xenon device having mercury bulb to produce UV (ultraviolet) light of range 200-320 nm range is also effective at patient rooms as it significantly decreases the pathogen concentration³⁰. Along with these innovations, there are some opportunities which can be adopted for further improvements, for example nanotechnology^{31, 32}.

Nanotechnology overview and its advantage over other infection diagnosis and control.

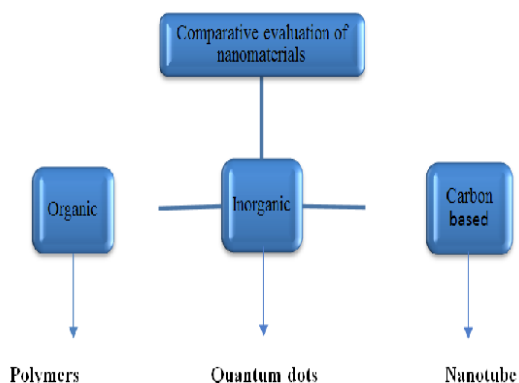


Fig. 1. Comparative evaluation of Nanomaterials

Types of nanomaterials:

In combination with above Fig. 1, there are below several types of exemplary nanomaterials which are being used in numerous applications.

Non-metallic inorganic nanomaterials (TiO_2 , SiO_2 , ZnO , $\text{Al}(\text{OH})_3$, Fe_2O_3 , Fe_3O_3 , ZrO_2 , CaO , ITO, ATO, CeO_2).

Metal alloys and metals (Au, Ag, Pt, Pd, Cu, Fe, Ni, Co, Al, Mn, Mo).

Nanomaterials enabled with carbon like fullerenes, carbon nanotubes, carbon nanofibers, graphene.

Dendrimers and nano polymers (polymeric nanoparticles, polymer nanotubes, nanowires and nanorods, nanocellulose, nanostructured polymer films).

Quantum dots (cadmium telluride, cadmium selenide, quantum dots free of cadmium).

Out of these nanoparticles, silver nanoparticles have various advantages due to antimicrobial potential. Other nanomaterials require more research and experiments to prove their credibility. Except silver nanomaterial, limited data is available for other nanomaterials. Refer below synthesis methods of silver nanoparticles:

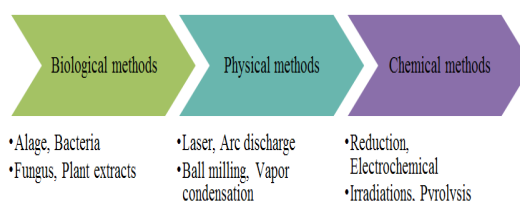


Fig. 2. Synthesis methods of silver nanoparticles

It was also found that some novel nanoparticles formed by placing silver sources in contact with surfaces. Same scenario is also applicable for copper silver nanoparticles. As there are multiple nanoscale objects which remain in contact with human also the source of incidental nanoparticles in the environment³³. Apart from the methods explained in Fig. 2, there may be some other innovative methods to develop silver nanoparticles which require more scientific research.

Antibacterial functioning of silver nanoparticles

As per Fig. below Fig. 3, silver ions are responsible for damaging cell wall of the gram positive and gram-negative bacteria responsible

for various infectious diseases. Oxidative stress of silver ions generates disruption in metabolic activities

which denature the ribosome in cytoplasm and cause cell death.

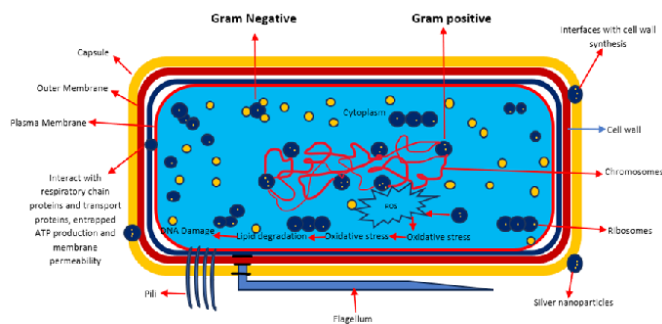


Fig 3. Antibacterial functioning of silver nanoparticles

Various methods for “silver nanoparticle” synthesis, relative benefits, and limitations³⁴

Table 1: Various methods used for silver nanoparticles’ synthesis

Method	Nano Size (nm)	Benefits	Drawbacks
Starch: Reducing and stabilizing agent, synthesis through AgNO_3 reduction to NS.	10 to 34	- Green synthesis - Stable for >90 days - Quite simple - Reproducible - Starch Biocompatibility	- Weak binding interactions - Reversible at higher temperatures
Enterobacteria: Bacterial enzymes, Synthesis through AgNO_3 to NPs.	28 to 122	- Environmentally safe - Used natural strain - Chemical free synthesis	- Less control over NP size - Unstable size distribution
Chitosan: AgNO_3 mix dropwise to chitosan in the presence of acetic acid, Ag^+ reduced to NS	4 to 5	- Small NS - Narrow size allocation - Inhibit NP accumulation	- Modify the NS properties.
$[\text{Ag}(\text{NH}_3)_2]^+$: N vinyl- 2-pyrrolidone with UV light, synthesized by reacting ammonia and silver oxide	4 to 6	- Stable synthesis - Narrow size distribution - No accumulation - Stabilizing sense	- Hard to increase size - PVP impact biological things
Electrical current: applied among two silver wires in presence of deionized water, it helps surface Ag atoms to concentrate aqueous NS	5 to 35	- Simple - Chemical free - Narrow size distribution - Kill <i>S. aureus</i>	- NP agglomeration - Less term stability
Penicillium Brevicompectum: Reduce AgNO_3 to NPs	23 to 105	- Environmentally friendly - No accumulation	- Wide size distribution - Lengthy process (>72 hours)
UV: Reduce AgNO_3 by using sodium dodecyl sulfate (SDS) and ethanol	23 to 67	- Kill <i>P. aeruginosa</i> - Reduced SDS-capped	- Superior antibacterial activity - mammalian cytotoxicity
Peptide: AgNO_3 stirred with peptide solution at different pH values for 12 hours	19	- Effective capping agents - Decrease NS toxicity	- Risk in medical applications

Silver and its salts are also beneficial in drug manufacturing, imaging and drug delivery systems³⁵ restorative dentistry, ophthalmology, skin/tissue repair, and nano biosensors³⁶. It required more studies to understand the broad application of silver nanoparticles for various medical devices.

Different methods of nanoparticle's "Surface coating," relative benefits and limitations:

For any of the metallic device, it is important to understand the type, structure, and its coating process. For example, in food industry, engineered nanoparticles like photocatalyst metal oxides may

be the promising agent to manage these resistive contaminants through surface coatings. It includes the development methods of antimicrobial surface coatings, associated characterization methods, evaluation of physical stability of coating along with antimicrobial efficiency³⁷. There are few exemplary techniques below which can be used for coating metal oxide. Due to metal-to-metal compatibility, surface coating has more potential to increase shelf life of coated materials. Now a days, industries have advanced techniques to develop surface coating, for example coating of metal oxide.

Table 2: Various techniques for coatings of metal oxide

Coating Technique	Description of process / technique	Used References
Sputter accumulation	Electron beam irradiation on metal mold steel surface to develop ceramic film	[38]
Atomic layer deposition	TMA- H ₂ O -process	[39]
Vapor deposition (chemicals)	Pressure vapor deposition process for the deposition. of MA3Bi2I9 perovskite film using co-sublimation of methylammonium	[40]
Sol-gel synthesis	Semicrystalline system (liquid and solid phases) with morphology for convert discrete particles to polymer networks	[41]
Coating with binders	Arc ion plating helps coating of TiCN-based cermets	[42]
Melt-blending	Nanotube/polymer composites diluted through pure polymer with the help of melt mixing process	[43]
Plasma-spraying	Additive manufacturing based on velocity and enthalpy of particle stream to strengthen the adhesion by using some specific mathematical models	[44]
Anodic oxidation	Plasma Electrolytic Oxidation (PEO) ionized Al alloy	[45]
Electrophoretic deposition	By this technique, movement of charged particles through liquid in the presence of electric field which deposited on conductive substrate (opposite charged)	[46]

Metal oxides demonstrated improved coating stability for the food packaging system along with other necessary cleaning and sanitization protocols. Ventilation, walls, floors, machinery, cause infections in hospitals⁴⁷. To overcome some critical surface infections, coating is best alternative. Shelf life of coating material and surface leaching are main challenges which need to be overcome for a stable coated product.

Different methods for "Nanocomposites" development, relative benefits, and limitations:

There are some studies which stated novel development of nanocomposites, for example, 3% Ag/TiO₂ nanocomposites developed with UV. It was the better alternative for antimicrobial activity for Gram-negative and gram-positive bacteria². Now a days, nanocomposite membranes are developed to increase the bacterial removal efficiency⁴⁸. Also, antimicrobial experiment proved that efficiency of Nanocomposite membrane is greater than 98%⁴⁹. Another example is Titanium dioxide which is synthesized by photocatalytic, can easily perform at ambient conditions, also it is eco-

friendly⁵⁰. Few models have capability for treatment of diabetic wound healing process^{51,52}. Plasma at low temperature activates the fiber surface to help strong binding of silver nanoparticles onto polyester fabrics⁵³. Effective composites form can be potential antibacterial agent for sanitation and disinfection industries^{54,55}. Based on iodine antibacterial nature and its effects against drug resistant organisms, iodine nanocomposite was developed by iodine immobilization through metal-organic frameworks can be beneficial for orthopedic implant⁵⁶. If there are alternative approaches to treat microbial infections, usage of silver-based products should be avoided. Overall, it is always a choice to use silver products based on their antimicrobial efficiency over cellular toxicity. Due to limited scope of few materials, nanocomposites have potential to tackle antibacterial resistance. Now a days in industries, advanced techniques are available for alloys combination or development of nanocomposites.

Different methods for “Quantum dots” development, relative benefits, and limitations:

Carbon Quantum Dots as assembled Spermidine are found active against resistant bacteria strain. This synthesis used ammonium citrate through pyrolysis and modified spermidine by simple heat treatment. In vitro wound healing studies were performed in mice to show quick healing, quick formation of collagen and better epithelialization when Spd/ CQDs used as a material for dressing⁵⁷. Biocompatibility results demonstrated no cytotoxic impact on mouse embryonic fibroblast cell lines but showed cytotoxicity against epithelial cell line of human. Few synthesis mechanisms can be used to develop disinfection material⁵⁸. Minor effect as hemolytic against red blood cells were seen in some cases⁵⁹.

Different methods for “Other nanotechnology approaches,” relative benefits and limitations

Other nanotechnology approaches are more related to air, water, textile, and food applications.

Nanotechnology approaches in “aerosol technology”

To control the aerosol contents, one of the studies was performed in which silver nanoparticles of approximate size 11 nm in diameter were coated onto a medium filter as antiviral agent. This coating

was performed by spark discharge generation system. Required parameters like filtration efficiency, anti-viral ability and pressure drop were evaluated with dust loading. Below Fig. 4 states the mechanism of this process. Results show that the purification capability and drop of pressure increase with dust, but the antiviral capability decreases. This mechanism can be useful to predict the filter life, anti-viral abilities of filter⁶⁰.

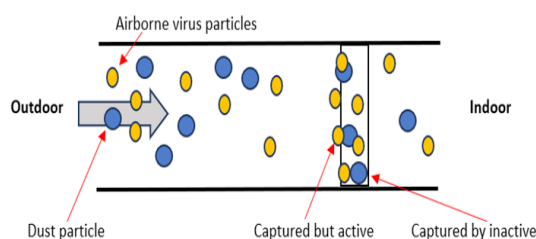


Fig. 4. Anti-viral action with dust loading

Silver ions also play vital role against SARS coronavirus due to its bacteriostatic and active ingredients properties. Based on these properties it is using for disinfection of medical equipment, wound therapy, and water purification. Recent studies also demonstrated effectiveness of iodophor iodine against non-enveloped virus⁶¹.

Nanotechnology approaches in “Textile technology”

Textile industries have been using nanotechnology for a long time. There are few assumptions about nanotoxicity that nanomaterial release during washing, but recent studies showed it is safe to human health as per regulatory requirements. Main concerns for this developing market are to keep the environment safe from residues developed through climate change⁶⁴. Functionalized technologies are being used to release the total silver and nanoparticles from textiles in the form of artificial sweat. Results of textiles functionalization lab prepared textiles showed the presence of embedded silver nanoparticles⁶⁵. Also, some other lab prepared textiles that showed presence of silver on coated fiber surface. This silver release from composites was found less as compared to surface coated textiles⁶⁶. Synthetic fibers can be used in cloth textiles for manufacturing comfortable clothes like sports clothes. Antimicrobial effect can be evaluated on woven and non-woven fabric with defined methods like padding or exhaust

methods. Addition of antimicrobial agent in spinning mass is an effective and durable method to treat the synthetic fibers⁶⁷.

Nanotechnology approaches in “Food technology”

There is a variety of consumer products developed by nanotechnology for children for example, sippy cups, cleaning goods, breast milk storing cups, cleaning products as well as humidified

fitment. Ionic silver release was noticed as 1:50 product to liquid mass ratio with synthetic saliva, sweat, urine, orange juice, onto dermal wipes and into air. In case of fabrics silver leaching was noticed via dissolution. In case of consumer products used by children silver leach is very minimum, also bioavailable silver can be iconic than normal form of silver⁶⁸. There are specific approaches for antimicrobial packaging as below:

Table 3: Antimicrobials into polymers and food packaging applications

Antimicrobial agents	Polymeric agents	Target organisms	Used References
Organic acids and anhydrides: acetic, benzoic, malic, sorbic	Edible layers, ethylene vinyl acetate, linear low-density polyethylene	Molds	[69]
Metal: Ag	Different polyolefins	Bacterial, yeast and fungal	[70]
Fungicide: Imazalil, Benomyl	low density polyethylene	Mold strains	[71]
Bacteriocins :Lacticin Nisin, pediocins	Edible layers, LDPE, cellulose	Bacterial strains of type Gram positive	[72]
Enzymes: Glucose, Lysozyme	PS, Cellulose acetate	Bacterial strains of type Gram negative	[73]
Oxidase	Edible films	Bacteria	[74]
Chelating agent: EDTA	Edible layers	Bacterial strains of type Gram negative	[75]
p-coumaic acids, Cinnamic, caffeic	Nylon, PS (polystyrene)	Molds, yeast, bacteria	[76]
plant extracts, seed extract,	LDPE, cellulose	Molds, yeast, bacteria	[77]

Migration of nanoparticles from food composites to food is also examined by insignificance level of migration inductively and attached ICP-MS (Inductively Coupled Plasma Mass Spectrometry) and based on improved specifications, Ag/PVC nanocomposites films are much promising of long-term food packaging applications⁷⁸.

Nanotechnology approaches in “water purification technology”

Availability of Safe and clean water now a days is a big concern worldwide. There are some specific steps by which this challenge can also be resolved, for example, development of green materials, polymers, adsorbents, and nanomaterials

used for purification of water⁶². There is defined procedure and protocol to develop new materials for water treatment⁶³. Refer below flow diagram in Fig. 5 for one of the water purification approaches:

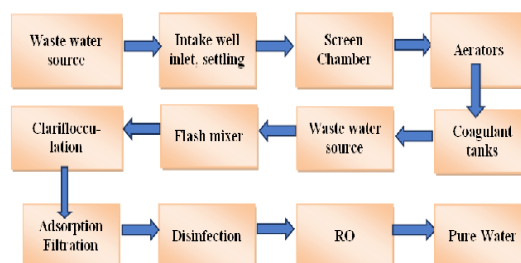


Fig. 5. Waste water treatment by use of nanomaterial.

For water treatment, if method used in combinations, it would show more effective results for water treatment. Specific absorbent may help with fast cleaning of water. There are other specific protocols which can be used for removing pollutants in water.

Associated risks or limitations of Nanotechnology:

In present scenario, world is facing several

types of medical challenges which cause the degradation of healthy life. Nanotechnology is playing a vital role in eliminating the infectious challenges through innovative developments. Based on novel innovations of nanotechnology there is much relief in healthcare sector but still there are many factors that function as major limitations in growth of nanotechnology. Few of them are summarized in the table below:

Table 4: Limitations of Nanotechnology

Sr. No.	Risk Factors	Description of associated risk factors	Used References
1	Cost	Initial set up or stages required high input amount	[2]
2	Safety	Usage beyond the shelf and not referring IFU	[23]
3	Consumption	Contact with toxic nanoparticles	[79]
2	Development time	From research & development to final approval it takes long time in case of drug development	[80]
3	Nanotoxicity	For sensitive domains, it is not easy to control toxicity	[81, 82]
4	Control release	Few specific nanoparticles are difficult to control release	
5	Manufacturing and scale up challenges	Pharmaceutical science always faces challenges in manufacturing and scales up	[83, 84]
6	Target areas and bioavailability	In medical field, it is hard to target area such as lungs, Poor bioavailability can alter the properties of nanomaterials and have adverse impacts	[85]

For above risks or limitations, regulatory support is required to develop and sustain advancement in nanotechnology. ISO (International Organization for Standardization), ASTM (American Society for Testing and Materials) and FDA (Food and Drug Administration), can play a vital role in continual improvement in product and processes.

More studies are required to understand the toxic aspects of nanomaterials. These studies can be planned with risk-based approaches.

Applications of Nanotechnology:

Nanotechnology plays a vital role in diagnosis and disease control due to its antibacterial properties. To sustain a healthy life, free from any type of bacterial infection is possible through innovative approach of nanotechnology.

Nanotechnology is useful in different sector due to its advanced applications. Different fields like medicine, food, household, water, hospitals, labs, and pharmaceuticals are taking advantage of these innovative approaches to nanotechnology. Few of them are explained in the table below:

Table 5: Nanotechnology's applications

Application factor	Description of application	Used References
Medical bioengineering	Diagnostic tests and tissue bioengineering	[86]
Medical devices	Develop biocompatibility, resist the biofilm creation	[87]
Food science	Detect pesticides, pathogens, and toxins	[88]
Food packaging	Develop intelligent food packaging system	[20]
Wound management	Wound healing by metal oxide, bio-kill approach in ICU	[89, 90]
Paints and coatings	Used in automotive, aerospace, construction, and hygiene industry	[91]
Antibacterial agent	Wide particle size range, high stability, variable morphological characteristics, and stable physical and chemical properties	[92, 93]
Water treatment	Medicinal items, consumer products, and agriculture	[49, 63]
Laboratories & industries	Semiconductor nanocrystalline oxides have the capability to absorb substrates	[53]
Photocatalysis	Chemically stable, capability of transmission in infra and visible regions	[94]
Coatings	Device coating which helps in infection control	[95]
Nanocomposites	Key role in sensors, catalysis, and electronic devices due to their thermal stability	[94]
Hospital textiles	Antimicrobial efficient Fabrics	[17, 53]
Food applications	Development of supplements, cooking oil catalyzation, crop pesticides, novel food packaging, and increase of food texture	[96]
Wound healing Management	Nanostructure of plant extracts helps to release phytochemicals at the wound site	[95]
Pharmaceuticals nano systems	Carbon nanotubes , quantum dots, nano shells, nanobubbles, paramagnetic nanoparticles, liposomes, niosomes, dendrimers, nano emulsions, and solid lipid nanoparticles	[46, 97, 98]
Biodegradable polymers	Food packaging edibility, renewability, and biocompatibility	[99]
SHOMAL	Developed for disinfection and disease prevention for poultry, livestock, and aquatics	[27]
Silver nanoparticles	Weapon against antibiotic resistant strains, animal husbandry, water/air filters, biomedical, and food packaging and antiviral applications, powerful weapon against COVID-19 infection	[100, 101, 102]
Iodine	Useful for orthopedics antibacterial therapy	[56]
Quantum dots	Effective against Staphylococcus aureus, epidemic methicillin-resistant, Staphylococcus aureus, Pseudomonas aeruginosa and Escherichia coli	[103]
Disease diagnosis with treatment	Cervical cancer detection through tumor markers, photodynamic therapy, drug delivery system, in-vivo imaging, therapy techniques, and tissue engineering	[104, 105]
lipid nanoparticles	Used in drug delivery and biomedical applications	[29, 30]
Coating on SS	Used in treatment of burn with antibacterial coating	[106]

CONCLUSION

Future prospectives

Nanotechnology is the interlinked branch of various sciences which have nanoparticles within range of 1 nm to 100 nm. Nanoparticles exist in different forms like silver nanoparticles, gold nanoparticles, surface coating metals, nanocomposites, Quantum dots, nano polymers. Due to antimicrobial properties of silver nanoparticles, it has shown effective disinfection. As a disinfectant, silver nanoparticles are used to develop air filters, animal farming, fabrics, biomedical and food wrapping items. Silver nanoparticles development is possible by three major methods i.e., biological methods (by algae, bacteria, plant extracts, fungus), physical methods (laser, arc-discharge, ball-milling, vapor condensation) and chemical methods (reduction, electrochemical, irradiation, pyrolysis). Out of these methods, biological method is effective and preferable for synthesis of silver nanoparticles.

To overcome various infectious challenges in biomedical field, silver nanoparticles are much promising agent. Risk associated with reusable medical devices can be minimized through surface coating of silver nanoparticles. Such coatings have high potential to improve the life of reusable medical devices. Based on specific features of silver nanoparticles, it is used in biomedical applications (devices, wound healing, hospital acquired infections), industries applications (textile, food packaging, animal husbandry), environmental applications (air disinfectants, water disinfectants, medical premises), drug delivery, catalytic actions, biomagnetic separations, biodetection and labelling. Silver nanoparticles are effective against different type of bacterial infections, and they are effective agents for the cases of antibiotic resistance.

As per these properties and applications, usage of silver nanoparticles can save time, efforts, and cost. This new field of silver nanoparticles can function as a strong backbone for industrial challenges and therapeutic issues. Reusable medical devices and medical consumables are always prone to get infectious if not fulfill the criteria of appropriate cleaning, disinfection, and sterilization. Non-fulfillment of monitoring, effectiveness checks and required evaluations are the possible root cause of current infectious stage. COVID-19 was one of the

examples for such type of infectious environment. Nanotechnology in aspects of different nanomaterials can be the solution to these challenges. Silver nanoparticles have immense scope to develop innovative medical devices and related consumables. For development of antimicrobial material/ devices, more research work is required to understand the concept of biocompatibility (sensitivity/ toxicity).

DISCUSSION

Nanotechnology provides a strong platform to evaluate antimicrobial properties of different nanomaterials. As per review, silver nanomaterials can be used as an effective weapon against antibiotic resistance. Life of reusable medical devices can be increased through coating of silver nanomaterials. Nanotechnology can help in different areas like food, medicine, water, and air but there are certain limitations which require more scientific advancement and research. Development of nanomaterial required high input amount and time for initial set up with appropriate safety and consumption. After from that if devices are developed as implants, then biocompatibility evaluation is the area which required more experiments are research.

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Conflict of interest

Authors declare that they have no competing interests

Data Availability Statement

This statement does not apply to this article.

Ethical approval

This study did not involve any human participants, animal subjects, or any material that requires ethical approval.

REFERENCES

1. Jones, K.E., Patel, N.G., Levy, M.A., Storeygard, A., Balk, D., Gittleman, J.L. and Daszak, P., *Nature*, **2008**, *451*, 990–993.
2. Deshmukh, S.P., Mullani, S.B., Koli, V.B., Patil, S.M., Kasabe, P.J., Dandge, P.B., Pawar, S.A. and Delekar, S.D., *Photochem. Photobiol.*, **2018**, *94*, 1249–1262.
3. Humphries, R.M. and McDonnell, G., *J. Clin. Microbiol.*, **2015**, *53*, 3118–3125.
4. Ko, Y.S., Joe, Y.H., Seo, M., Lim, K., Hwang, J. and Woo, K., *J. Mater. Chem. B*, **2014**, *2*, 6714–6722.
5. Nimodiya, A.R. and Ajankar, S.S., *Int. J. Creative Res. Thoughts*, **2022**, *1*, 1–4.
6. Deshmukh, S.P., Patil, S.M., Mullani, S.B. and Delekar, S.D., *Mater. Sci. Eng. C*, **2018**, *1*, 1–12.
7. Roberts, C.G., *Am. J. Infect. Control*, **2013**, *41*, S77–S80.
8. Reprocessing Medical Devices in Health Care Settings: Validation Methods and Labeling - Guidance for Industry and Food and Drug Administration Staff, **2015**.
9. Kumar, D., Sonawane, U., Gohil, M.K., Pol, R., Patil, A.S., Mittal, R. and Agarwal, A.K., *Trans. Indian Natl. Acad. Eng.*, **2020**, *5*, 299–303.
10. Mahmoud, E.R.I., Algahtani, A. and Tirth, V., *Met. Mater. Int.*, **2020**, *26*, 12540–12550.
11. Ben-Sasson, M., Lu, X., Bar-Zeev, E., Zodrow, K.R., Nejati, S., Qi, G., Giannelis, E.P. and Elimelech, M., *Water Res.*, **2014**, *62*, 260–270.
12. Dankovich, T.A. and Gray, D.G., *Environ. Sci. Technol.*, **2011**, *45*, 1992–1998.
13. Mehta, Y., Gupta, A., Todi, S., Myatra, S.N., Samaddar, D.P., Patil, V., Bhattacharya, P.K. and Ramasubban, S., *Indian J. Crit. Care Med.*, **2014**, *18*, 149–163.
14. Nadaf, N.Y. and Kanase, S.S., *Dig. J. Nanomater. Bios.*, **2015**, *10*, 1189–1199.
15. Sharifi-Rad, J., Hoseini Alfatemi, S.M., Sharifi Rad, M. and Iriti, M., *Ann. Med. Health Sci. Res.*, **2014**, *4*, 863–868.
16. Peter, A., Mihaly-Cozmuta, L., Mihaly-Cozmuta, A., Nicula, C., Indrea, E. and Barbu-Tudoran, L., *J. Food Process Eng.*, **2014**, *37*, 596–608.
17. von Goetz, N., Lorenz, C., Windler, L., Nowack, B., Heuberger, M. and Hungerbühler, K., *Environ. Sci. Technol.*, **2013**, *47*, 9979–9987.
18. Echegoyen, Y. and Nerin, C., *Food Chem. Toxicol.*, **2013**, *62*, 16–22.
19. Mitrano, D.M., Barber, A., Bednar, A., Westerhoff, P., Higgins, C.P. and Ranville, J.F., *J. Anal. At. Spectrom.*, **2012**, *27*, 1131–1137.
20. Chmielowiec-Korzeniowska, A., Tymczyna, L., Dobrowolska, M., Banach, M., Nowakowicz-Dąbek, B., Bryl, M., Drabik, A., Tymczyna-Sobotka, M. and Kolejko, M., *Open Chem.*, **2015**, *13*, 1269–1274.
21. Owusu, E.G.A., Ph.D. Thesis, University College London, **2020**, 1–282.
22. Seavey, R., *Am. J. Infect. Control*, **2013**, *41*, S111–S117.
23. Khan, I., Saeed, K. and Khan, I., *Arabian J. Chem.*, **2019**, *12*, 908–931.
24. Boyce, J.M., *Antimicrob. Resist. Infect. Control*, **2016**, *5*, 10.
25. Weber, D.J. and Rutala, W.A., *Am. J. Infect. Control*, **2013**, *41*, S67–S71.
26. Glover, R.D., Miller, J.M. and Hutchison, J.E., *Nano Lett.*, **2011**, *11*, 8950–8957.
27. Nia, J.R., Using of Nano Silver in Poultry, Livestock, and Aquatic Industry, US Patent, **2009**, 1–3.
28. Chaloupka, K., Malam, Y. and Seifalian, A.M., *Trends Biotechnol.*, **2010**, *28*, 580–588.
29. Burdus, A.-C., Gherasim, O., Grumezescu, A.M., Mogoanta, L., Ficai, A. and Andronescu, E., *Nanomaterials*, **2018**, *8*, 681.
30. Sim, S. and Wong, N.K., *Biomed. Rep.*, **2021**, *14*, 1–9.
31. Valavanidis, A., Nanotechnological Breakthroughs in Medical Devices, ResearchGate, **2019**, 1–28.
32. Ruggeri, M., Bianchi, E., Rossi, S., Vigani, B., Bonferoni, M.C., Caramella, C., Sandri, G. and Ferrari, F., *Pharmaceutics*, **2020**, *12*, 815.
33. Pantic, I., *Rev. Adv. Mater. Sci.*, **2014**, *37*, 15–19.
34. Li, Y.-J., Harroun, S.G., Su, Y.-C., Huang, C.-F., Unnikrishnan, B., Lin, H.-J., Lin, C.-H. and Huang, C.-C., *Adv. Healthcare Mater.*, **2016**, *1*, 1–10.
35. Wigginton, N., De Titta, A., Piccapietra, F., Dobias, I., Nesatyy, V.J. and Latmani, R.B., *Environ. Sci. Technol.*, **2010**, *44*, 2163–2168.
36. Weckman, T. and Laasonen, K., *Phys. Chem.*

- Chem. Phys.*, **2015**, 17, 1–13.
37. Marassi, V., Di Cristo, L., Smith, S.G.J., Ortelli, S., Blosi, M., Costa, A.L., Reschiglian, P., Volkov, Y. and Prina-Mello, A., *R. Soc. Open Sci.*, **2017**, 4, 170726.
 38. Misumi, S., Okada, A., Okamoto, Y. and Inoue, M., *Procedia CIRP*, **2013**, 5, 486–491.
 39. Weckman, T. and Laasonen, K., *Phys. Chem. Chem. Phys.*, **2015**, 17, 1–13.
 40. Chen, X., Myung, Y., Thind, A., Gao, Z., Yin, B., Shen, M., Cho, S.B., Cheng, P., Sadtler, B., Mishra, R. and Banerjee, P., *J. Mater. Chem. A*, **2017**, 5, 1–39.
 41. Badanayak, P. and Vastrad, J.V., *Pharma Innovation J.*, **2021**, 10, 1023–1027.
 42. Xian, G., Xiong, J., Zhao, H., Xian, L., Fan, H., Li, Z. and Du, H., *Wear*, **2020**, 460–461, 1–10.
 43. Kamble, S., Bhosale, K., Mohite, M. and Navale, S., *Int. J. Adv. Res. Sci. Commun. Technol.*, **2023**, 1, 1–7.
 44. Vu, D., *Eng. Technol. Appl. Sci. Res.*, **2023**, 10367–10371.
 45. Trivinho-Strixino, F., Delgado-Silva, A.O., Santos, J.S., Rodrigues, A., Mambrini, G.P. and Sikora, M.S., *Plasma*, **2023**, 235–249.
 46. Marchewka, J., Kołodziejczyk, E., Bezkosty, P. and Sitarz, M., *Sci. Rep.*, **2023**, 1–15.
 47. Boyce, J.M., *Antimicrob. Resist. Infect. Control*, **2016**, 5, 10.
 48. Al-Maamori, M.H., Al-Tamimi, D.H. and Braihi, A., Mattingley Publ. Co., Inc., **2020**, 23142–23164.
 49. Liu, Y., Rosenfield, E., Hua, M. and Mi, B., *Water Res.*, **2013**, 47, 2949–2958.
 50. Patil, S.M., Deshmukh, S.P., Dhodamania, A.G., Morea, K.V. and Delekar, S.D., *Curr. Org. Chem.*, **2017**, 21, 821–833.
 51. Xu, Q., Aa, S., Gao, Y., Guo, L., Creagh-Flynn, J., Zhou, D., Greiser, U., Dong, Y., Wang, F., Tai, H., Liu, W. and Wang, W., *Adv. Mater.*, **2018**, 1, 1–38.
 52. Pal, S., Nisi, R., Stoppa, M. and Licciulli, A., *ACS Omega*, **2017**, 2, 3632–3639.
 53. Ilic, V., Saponjic, Z., Vodnik, V., Lazovic, S., Dimitrijevic, S., Jovanc, P., Nedeljkovic, J.M. and Radetic, M., *Ind. Eng. Chem. Res.*, **2010**, 49, 7287–7293.
 54. Duncan, T.V., *J. Colloid Interface Sci.*, **2011**, 363, 1–24.
 55. Mohan, A.N. and Manoj, B., *Mater. Chem. Phys.*, **2019**, 232, 137–144.
 56. Teng, W., Zhang, Z., Wang, Y., Ye, Y., Yinwang, E., Liu, A., Zhou, X., Xu, J., Zhou, C., Sun, H., Wang, F., Zhang, L., Cheng, C., Lin, P., Wu, Y., Gou, Z., Yu, X. and Ye, Z., *Small*, **2021**, 1, 1–13.
 57. Li, Y.-J., Harroun, S.G., Su, Y.-C., Huang, C.-F., Unnikrishnan, B., Lin, H.-J., Lin, C.-H. and Huang, C.-C., *Adv. Healthcare Mater.*, **2016**, 1, 1–10.
 58. Shi, H., Wang, W., Zhang, L., Tang, Z. and Fan, J., *J. Environ. Chem. Eng.*, **2021**, 9, 1–11.
 59. Kovacova, M., Markovic, Z.M., Humpolicek, P., Micusik, M., Švajdlenkova, H., Kleinova, A., Danko, M., Kubat, P., Vajdak, J., Capakova, Z., Lehocky, M., Munster, L., Todorovi, B.M. and Spitalsky, Z., *ACS Biomater. Sci. Eng.*, **2018**, 1, 1–50.
 60. Joe, Y.H., Park, D.H. and Hwang, J., *J. Hazard. Mater.*, **2016**, 301, 547–553.
 61. Teng, W., Zhang, Z., Wang, Y., Ye, Y., Yinwang, E., Liu, A., Zhou, X., Xu, J., Zhou, C., Sun, H., Wang, F., Zhang, L., Cheng, C., Lin, P., Wu, Y., Gou, Z., Yu, X. and Ye, Z., *Small*, **2021**, 1, 1–13.
 62. Saleh, T.A., *Environ. Technol. Innov.*, **2021**, 1, 1–18.
 63. Yang, W., Hu, W., Zhang, J., Wang, W., Cai, R., Pan, M., Huang, C., Chen, X., Yan, B. and Zeng, H., *Chem. Eng. J.*, **2020**, 1, 1–36.
 64. Levard, C., Hotze, E.M., Lowry, G.V. and Brown, G.E., *Environ. Sci. Technol.*, **2012**, 46, 6900–6914.
 65. Reed, R.B., Zaikova, T., Barber, A., Simonich, M., Lankone, R., Marco, M., Hristovski, K., Herckes, P., Passantino, L., Fairbrother, D.H., Tanguay, R., Ranville, J.F., Hutchison, J.E. and Westerhoff, P.K., *Environ. Sci. Technol.*, **2016**, 50, 4018–4026.
 66. Zille, A., Fernandes, M.M., Francesko, A., Tzanov, T., Fernandes, M., Oliveira, F.R., Almeida, L., Amorim, T., Carneiro, N., Esteves, M.F. and Souto, A.P., *ACS Appl. Nano Mater.*, **2015**, 4, 7020–7028.
 67. Manna, J., Goswami, S., Shilpa, N., Sahu, N. and Rana, R.K., *ACS Appl. Mater. Interfaces*, **2015**, 1, 1–25.
 68. Quadros, M., Pierson, R.IV, Tulve, N., Willis, R., Rogers, K., Thomas, T. and Marr, L.C., *Environ. Sci. Technol.*, **2013**, 1, 1–31.
 69. Cuq, B., Gontard, N. and Guilbert, S., *Active Food Packaging*, **1995**, 111–142.

70. Ishitani, T., Active Packaging for Food Quality Preservation in Japan, Royal Society of Chemistry, **1995**, 177–188.
71. Weng, Y., Chen, M. and Chen, W., *LWT - Food Sci. Technol.*, **1999**, 32, 191–195.
72. Prudencio, C.V., dos Santos, M.T. and Vanetti, M.C.D., *J. Food Sci. Technol.*, **2015**, 52, 5408–5417.
73. Beacham, I.R., *Int. J. Biochem.*, **1979**, 10, 877–883.
74. Yousefi, M., Nematollahi, A., Shadnoush, M., Mortazavian, A.M. and Khorshidian, N., *Front. Nutr.*, **2022**, 1, 1–14.
75. Abarca, R.L., Medina, J., Alvarado, N., Ortiz, P.A. and Carrillo, B.L., *PLoS ONE*, **2022**, 1, 1–19.
76. Konzock, O., Tous-Mohedano, M., Cibin, I., Chen, Y. and Norbeck, J., *AMB Express*, **2023**, 1, 1–10.
77. Ordon, M., Zdanowicz, M., Nawrotek, P., Stachurska, X. and Mizielska, M., *Int. J. Mol. Sci.*, **2021**, 1, 1–17.
78. Azlin-Hasim, S., Cruz-Romero, M.C., Morris, M.A., Padmanabhan, S.C., Cummins, E. and Kerry, J.P., *Food Bioprocess Technol.*, **2016**, 1, 1–13.
79. Liu, Y., Rosenfield, E., Hua, M. and Mi, B., *Water Res.*, **2013**, 47, 2949–2958.
80. Herzog, F., Clift, M.J.D., Piccapietra, F., Behra, R., Schmid, O., Petri-Fink, A. and Rothen-Rutishauser, B., *Part. Fibre Toxicol.*, **2013**, 10, 11.
81. Gliga, A.R., Skoglund, S., Wallinder, I.O., Fadeel, B. and Karlsson, H.L., *Part. Fibre Toxicol.*, **2014**, 1, 1–17.
82. Pulit-Prociak, J. and Banach, M., *Open Chem.*, **2016**, 14, 76–91.
83. Levard, C., Hotze, E.M., Lowry, G.V. and Brown, G.E., *Environ. Sci. Technol.*, **2012**, 46, 6900–6914.
84. Kumari, S. and Sarkar, L., *J. Sci. Res.*, **2021**, 65, 42–46.
85. Calderon, L., Han, T., McGilvery, C.M., Yang, L., Subramaniam, P., Lee, K.-B., Schwander, S., Tetley, T.D., Georgopoulos, P.G., Ryan, M., Smith, R., Chung, K.F., Liou, P.J., Zhang, J. and Mainelis, G., *Atmos. Environ.*, **2017**, 160, 42–52.
86. Agboklu, M., *J. Nanotechnol. Res.*, **2024**, 1, 1–5.
87. Ramasamy, M. and Lee, J., *Biomed Res. Int.*, **2016**, 1, 1–17.
88. Banach, M., Tymczyna, L., Chmielowiec-Korzeniowska, A. and Pulit-Prociak, J., *Bioinorg. Chem. Appl.*, **2015**, 1, 1–15.
89. Duncan, T.V., *J. Colloid Interface Sci.*, **2011**, 363, 1–24.
90. Behbudi, G., *Adv. Appl. NanoBio-Technol.*, **2021**, 2, 63–67.
91. Edwards-Jones, V., *Lett. Appl. Microbiol.*, **2009**, 49, 147–152.
92. Haleem, A., Javaid, M., Singh, R.P., Rab, S. and Suman, R., *Glob. Health J.*, **2023**, 7, 70–77.
93. Agboklu, M., *J. Nanotechnol. Res.*, **2024**, 1, 1–5.
94. Dunnill, C.W., Page, K., Aiken, Z.A., Noimark, S., Hyett, G., Kafizas, A., Pratten, J. and Wilson, M., *J. Phys. Chem. Solids*, **2018**, 115, 127–136.
95. Hajjalyani, M., Tewari, D., Sobarzo-Sánchez, E., Nabavi, S.M., Farzaei, M.H. and Abdollahi, M., *Int. J. Nanomed.*, **2018**, 13, 5023–5043.
96. Cushen, M., Kerry, J., Morris, M., Cruz-Romero, M. and Cummins, E., *Trends Food Sci. Technol.*, **2012**, 24, 30–46.
97. Ansari, N., Ali, J. and Zulfequar, M., Austin Publishing, **2021**, 71–90.
98. Wagener, S., Dommershausen, N., Jungnickel, H., Laux, P., Mitrano, D.M., Nowack, B., Schneider, G. and Luch, A., *Environ. Sci. Technol.*, **2016**, 1, 1–17.
99. Metak, A.M. and Ajaal, T.T., *Int. Schol. Sci. Res. Innov.*, **2013**, 7, 1103–1109.
100. Li, Y.-J., Harroun, S.G., Su, Y.-C., Huang, C.-F., Unnikrishnan, B., Lin, H.-J., Lin, C.-H. and Huang, C.-C., *Adv. Healthcare Mater.*, **2016**, 1, 1–10.
101. Chen, X., Xu, J., Ji, B., Fang, X. and Qian, J., *Front. Bioeng. Biotechnol.*, **2023**, 1, 1–13.
102. Behbudi, G., *Adv. Appl. NanoBio-Technol.*, **2021**, 2, 63–67.
103. Owusu, E.G.A., Ph.D. Thesis, University College London, **2020**, 1–282.
104. Agboklu, M., *J. Nanotechnol. Res.*, **2024**, 1, 1–5.
105. Huang, Y., Guo, X., Wu, Y., Chen, X., Feng, L., Xie, N. and Shen, G., *Signal Transduct. Target. Ther.*, **2024**, 1, 1–50.
106. Shirani, S., Emadi, R., Eslami, A. and Saboori, A., *Mater. Today Commun.*, **2025**, 1, 1–15.