Toxicology of Nanoparticles in Drug Delivery

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Accepted: 19 October 2021 / Published online: 24 November 2021
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Abstract
Nanoparticles have revolutionized biomedicine especially in the field of drug delivery due to their intriguing properties such as systemic stability, level of solubility, and target site specificity. It can, however, be both beneficial and damaging depending on the properties in different environments, thus highlighting the importance of nanotoxicology studies before use in humans. Different types of nanoparticles have been used in drug delivery, and this review summarizes the recent toxicity studies of these nanoparticles. The toxicological evaluation of three widely used nanoparticles in drug delivery that are metal, lipid, and protein nanoparticles has been discussed in detail. Studies have recorded several toxic effects of various nanoparticles such as metal-based nanoparticles have been linked to increased oxidative stress and have the potential to infiltrate the cell nucleus and protein-based nanoparticles have been observed to have hepatotoxicity and nephrotoxicity as their adverse effects. Considering the increasing application of nanoparticles in drug delivery and the growing concerns of regulatory authorities regarding the toxicity of nanocarriers in living organisms, it requires urgent attention to identify the gap in toxicity studies. The review highlights the gap in toxicity studies and potential focus areas to overcome the existing challenges.

Keywords Nanoparticles · Nanotoxicology · Toxicity assessment · Protein nanoparticles · Metal nanoparticles · Lipid nanoparticles

Abbreviations
DXR Doxorubicin
HSA Human serum albumin
RA Rheumatoid arthritis
CLT Celastrol
CPP Cell-penetrating peptide
GNPs Gelatin nanoparticles
RFRTs Ferritin derivative apoferritin
TIMP-GLIA Tolerogenic immune-modifying nanoparticles containing gliadin
FNPs Fibroin nanoparticles
SFN-DXR Silk fibroin nanoparticle bearing doxorubicin
TSFN Tween 80–coated silk fibroin nanoparticles
SF Silk fibroin
HPLC High-performance liquid chromatography
IC50 Half maximal inhibitory concentration
LD50 Median lethal dose

Introduction
The recent years have marked an unprecedented growth in nanotechnology due to its wide range of applications [1]. The rising popularity can be attributed to their astonishing properties which sets them apart from any other material. This has allowed their application to be expanded in the field of biomedicine, for the purpose of drug delivery, nutraceuticals, diagnostics/imaging, production of biocompatible materials, and much more [2, 3]. The technological advancements using nanoparticles have gained importance in such a short period due to impressive physical and chemical properties. A particle having size less than 100 nm (any one dimension) is characterized as a nanoparticle as per the new definition. These nanostructures possess great surface...
to area ratio, show quantum effects, and are highly mobile when present in free state [4]. The conventional drugs face problems like low specificity, less bioavailability, and high toxicity. These problems can be substantially overcome by use of nanoparticle-based drug delivery systems [5]. Metal, non-metal, polymers, and biopolymers have been used for preparation of nanoparticles/nanoformulations [6, 7]. For therapeutics, a nanoparticle can be used in two ways by acting itself (1) as a drug or (2) as a carrier for another substance [7, 8]. Since these nanoparticles are entering the biological system, they interact with various biological pathways and biomolecules found within living beings. Where on one side the unique properties of nanoparticles makes them ideal for drug delivery, the same properties also raise challenges, if not optimized efficiently [9]. As mentioned before, nanoparticles have a great surface to area ratio, which in one case provides benefit of efficient drug loading, but on the contrary, it can also increase chances of interaction with other biomolecules when present in a living organism. Some of the other properties that attract the concerns of scientists when using nanoparticles as drug delivery systems are their particle size (which allows easy access to various organs), charged surface (increasing chances of non-specific interaction), solubility, surface coating, and more. Various researches over the years support the claim that nanoparticles can be a bane or boon. We mostly know about the tremendous benefits that nanoparticles offer, but there is very limited information about toxicity, non-specific protein interaction, translocation to secondary target organs, etc. [10]. Some of the in vivo and in vitro studies have already highlighted the toxic effects of silica, gold, silver, and titanium nanoparticles. Since the size of the nanoparticles used in these studies is similar to the size used in biomedicine, it becomes imperative to assess the toxicological parameters. This review tries to highlight toxicity studies for some of the commonly used nanoparticles in drug delivery (Figs. 1 and 2).

**Carbon-Based Nanoparticles**

Nanoparticle-based drug formulations have become prominent in recent years due to their properties of targeted and sustainable drug release, enhanced solubility of the drug, and minimum or no adverse drug reactions. Among numerous nanocarriers, carbon nanotubes have been one of the major focuses because of their wide application in cancer therapies [11]. However, the application of carbon nanotubes has been also expanded to non-anticancer drugs, either as a carrier or as an adjuvant [12]. Carbon nanotubes are made from graphene sheets. These sheets are rolled into different shapes (such as ellipsoid, hollow, sphere, and tubes) and sizes. The carbon nanotubes are majorly classified into two types that are SNWCT (single-walled carbon nanotubes) and MWCNT (multi-walled carbon nanotubes) [11, 13, 14]. The advanced physical, mechanical, and chemical properties of CNT raise some concern among researchers when used in drug delivery systems. This is because CNTs can move across the cell membranes while commuting the carrier biomolecule. Therefore, understanding the cytotoxicity of CNTs becomes very crucial [15].

Studies have shown that CNTs can cause neurotoxicity, pulmonary toxicity, immune toxicity, embryotoxicity, genotoxicity, hepatotoxicity, and cardiovascular toxicity [16–22]. Gholamine et al. studied the toxicological impact of SWNTs and MWCNT in mice. The mice were randomized into five groups and injected with MWCNT and SWNTs suspended in PBS (phosphate-buffered saline). After 2 weeks of observation, mice of each group were evaluated for a brain factor known as the brain-derived neurotrophic factor (BNDF). The factors support overall neuron health by promoting sprouting, improving survival, as well as controlling other aspects such as memory, learning, locomotion, and more. SWNTs and MWCNT showed anxiogenic and depression effects, while also impacting the gene expression of BNDF [16]. Fujita et al. conducted studies to understand the in vitro and in vivo pulmonary toxicity caused by SWNTS. It was found that SWNTS induced inflammation and slower recovery and...
enhanced the expression of genes governing cell proliferation, inflammation, and reactive oxygen species generation. However, the level of toxic impact is dependent on the size of SWNTs used [23]. Reproductive and developmental toxicity of CNTs was observed in mice after intravenous administration. Fetal malformations, miscarriages, and teratogenic and high levels of ROS were some of the observations [24]. A study by Shang et al. highlights damage caused to the kidney and brain by SWCNTs by promoting ROS in mice [25]. Zeinabad et al. studied the CNTs interaction with tau protein and PC12 cells. Results showed that both MWCNT and SWCNT induced different modes of cell death pathway. Moreover, CNTs led to neurotoxicity and protein conformational changes [26••]. Cao et al. mention about the toxicological impact of CNT to the vascular system, wherein they induce formation of atherosclerotic plaque, poor heart rate, and vasomotor dysfunction among animal models [27]. CNTs have been used effectively to deliver anti-tumor drug paclitexal in animal models. The study showed significant tumor suppression and effective release of the drug with minimal side effects. However, the study lacks any toxicity assessment for short-term or long-term period [28].

Metal-Based Nanoparticles

Superparamagnetic iron oxide (SPIIONS) is one of the FDA-approved nanoparticles, used widely in diagnostics as well as therapeutic application (hyperthermia, magnetic drug delivery, treatment of anemia) [29]. However, the recent years have observed expansion of SPIONS as potential drug delivery systems to treat cancer and other diseases [30]. But, the concerns related to toxicity of SPIONS have somehow limited their use until their property is enhanced. This is the reason that some of the previously FDA-approved SPIONS for use in medical imaging have been pulled off concerning their toxicity issues and anaphylactic reactions. Existing studies have demonstrated challenges like ROS, LDH leakage, inflammation, DNA damage, and alteration in mitochondrial functioning [31–33]. Currently, Feraheme™ or ferumoxytol of AMAG pharmaceuticals is the only approved drug used in anemic patients suffering from chronic kidney disease (CKD) [34–36]. Umirem, Gastromark, and Feridex are some of the other FDA-approved SPIONS used as imaging agents [34, 37].

SPIONS are also known to impact the complement system leading to innate immunity-related side effects. One such SPION-based drug known to alter the complement system is Feridex™, which is now discontinued by FDA [38–40]. Another concern with use of SPIONS is they show well-known Haber–Weiss and Fenton reactions as the free ions (Fe2+) released after lysosomal ingestion react with oxygen or hydrogen peroxide in the powerhouse of cell (mitochondria), forming free radicals which are responsible for genotoxicity, DNA damage, and cellular inflammation [41, 42]. Some of the recent studies have highlighted that the toxicity of SPIONS can be reduced by use of coating agents. Polymers like polyethylene glycol (PEG), poly vinyl (PV), dextran, and chitosan can enhance the half-life and stability of SPIONS and prevent aggregation within cells [42, 43]. Recently, Ferreti et al. evaluated the biocompatibility of magnetic particles. The study finds that when magnetic iron nanoparticles are coated with zwitter-ionic ligands, they showed good stability and biodistribution and required levels of renal clearance [44]. Toxicity of SPIONS is a critical issue which needs to be considered before its extensive use as a drug delivery system. Though lots of research have been carried out to understand the toxicity relation, it is crucial to
dig deeper to enhance their property further. This will lead to more safe use of SPION nanoparticles and FDA approval for drug-based purposes. Similar to SPIONs, other metal nanoparticles like titanium, gold, silver, and zinc have been used for drug delivery but showed significant toxicity.

**Lipid-Based Nanoparticles**

Alec D Bangham discovered liposome back in the 1960s at the University of Cambridge (Babraham Institute). Liposomes are small and spherical structures forming an enclosed compartment, composed of phospholipid layers. The cavity formed at the center due the interaction between hydrophilic head and hydrophobic tail allows them to act as amphipathic nanocarriers for a wide range of drugs in therapies [45, 46]. About more than 40 liposome-based drugs are either approved for use in the market or are undergoing clinical trials [47, 48]. This increases interest in understanding the interaction of liposomes within the living system and so does their toxic effects. Upon entry into the living system, liposomes interact with a wide range of biomolecules such as LDL, HDL, and opsonins. The encounter with opsonins leads to activation of RES recognition, which further tries to eliminate it from the system. Other lipoproteins in the blood are known to reduce the stability of liposomes within the biological system by leading to rearrangement of surface lipids. This is one of the major concerns when using liposomes as a drug delivery system [49]. To overcome this challenge, liposomes are PEGylated as they prevent opsonization and support evasion from phagocytic cells [47, 50].

Doxil® is an FDA-approved PEGylated liposome-based drug used for treatment of tumor [51]. However, Gabizon et al. highlighted that encapsulated doxorubicin alters particle clearance and phagocytic functioning of cells in the liver. Recently, FDA has approved emergency use of two lipid nanoparticle-based mRNA COVID vaccines of Pfizer-BioNTech, proven to be 95% effective [52, 53]. However, a major challenge of vaccine instability has been observed. Adjustments of lipid nanoparticle structure would be necessary to enhance and improvise the vaccine stability.

In 2018, FDA EC approved another lipid RNAi-based drug named ONPATTRO by Alnylam Pharmaceuticals for diseases caused due to altered TTR (Transthyretin) protein. Risk and safety studies associated with the drug use indicate infusion-related reactions, which can be reduced with vitamin A supplementation and premedication with certain drugs like antihistamines and corticosteroids. Some of the common adverse reactions observed with use of drugs include respiratory symptoms, headache, and change in blood pressure. However, there are limited studies to understand the use of this drug among pregnant women, patients with renal issues, etc. [48, 54, 55].

Some of the PEG-liposomal formulations are negatively charged; this often leads to complement activation, hypersensitivity reactions (HSRs), cardiopulmonary distress, and anaphylactoid reactions. Hoven et al. studied the impact of negatively charged PEGylated liposome nanocarriers. Liposomes with variations in PEG in terms of liposomal size, chain length, surface concentration, and anchor molecule were done. While other variations caused no to mild effects, PEG anchored with cholesterol showed maximum complement activation [56]. Another similar study is conducted by Szebeni et al., where they evaluated the doxil and hynic PEG liposomes for complement activation and hypersensitivity [57].

**Protein-Based Nanoparticles**

Protein-based nanoparticles have received much attention in recent times due to their biocompatibility, amphiphilicity, easy biodegradability, and reduced toxicity. Albumin, gelatin, ferritin, fibroin, and casein are some of the widely used protein-based nanoparticles for drug delivery [58].

Human serum albumin is a flexible protein that is used to make albumin-based nanocarriers (ANCs) for the administration of cancer treatments [59, 60]. Because of the various drug binding sites found in the albumin molecule, a large amount of drug can be integrated into the particle matrix with serum albumin nanoparticles [61]. HSA may be a valuable analytical indicator for people with autoimmune diseases. Thanks to breakthroughs in biotechnology and protein science, many albumin-associated pharmaceutical preparations, such as nab-paclitaxel (Abraxane®), albumin nanoparticles manufactured utilizing nab technology, are already on the market [62]. The FDA has also approved a paclitaxel formulation (Abraxane). Abraxane differs from previous paclitaxel formulations in that it is attached to albumin particles, which confer water solubility; previous formulations had to be administered in the solvent Cremophor®, which raised allergic reactions and limited dosing [63].

While looking at the preferred outcomes, we must not ignore the drawbacks. Albumin NPs have been claimed to be unstable and extremely damaging to healthy cells. To overcome this and to lengthen their half-life in an aqueous environment and/or limit the development of protein macro-aggregates, the freshly synthesized albumin NPs must be stabilized or cross-linked. Thermal treatment, high hydrodynamic pressure, and enzymatic cross-linking using genipin or transglutaminase are some of the ways that can be used to stabilize protein nanoparticles. Glutaraldehyde cross-linking is one of the most commonly utilized ways for stabilizing albumin nanoparticles in general [64]. Steinhauser et al. used heat denaturation of HSA to produce particles loaded with ASOs and found out that HSA nanoparticles cross-linked...
with glutaraldehyde exhibited no toxicity in cell culture at concentrations of up to 5 mg/ml [65].

Langer [66], used pH and buffer titration studies were used to assess the stability and electrical behavior of the HSA nanoparticles, with a typical titration profile of prepared nanoparticles throughout a pH range of 3 to 10. The HSA nanoparticles' isoelectric point (pI) was calculated to be 5.05. The nanoparticles became unstable at pH values around pI, as evidenced by PCS: particle diameter increased from 250 nm to roughly 2.7 μm (Langer, 2003). Even at higher pH values when the nanoparticles had a strong surface charge, the particle aggregation remained largely irreversible. They concluded that pH values that lead to neutral particle surface charges should be avoided when handling protein-based nanoparticles. In an HSA nanoparticle suspension, the pI value, as well as the buffer content, was found as critical criteria for particle stability [66].

Albumin nanocarriers are used in the treatment of a variety of disorders in addition to cancer. HSA-coupled TRAIL delivery, according to Byeon et al., could be a promising therapy for rheumatoid arthritis (RA), a chronic autoimmune disease marked by extreme synovial hyperplasia and joint destruction. They also found that rats given CLT-encapsulated HSA nanoparticles had milder hepatotoxicity, nephrotoxicity, and cardiotoxicity than rats given CLT alone, according to serum analysis and histopathological examination [67].

Mesken et al. recently released a study on plasmid transmission in HEK 293 T cells loaded onto HSA nanoparticles coated with cell-penetrating peptide (CPP). They were prepared using the desolvation method. There was little cytotoxicity and no significant change in performance when nanoparticle-mediated transfection was examined at low plasmid concentration levels [68]. Even though numerous studies and tests have been conducted to assess the efficiency of HSA nanocarriers and their potentially hazardous effects at various doses, sufficient evidence suggesting the detrimental toxicity of this nano-carrier is still lacking.

Gelatin nanoparticles have recently been offered as a feasible option for parenteral formulations due to their low cost, biocompatibility, biodegradability, minimal antigenicity, and application in a variety of formulations [69]. The US Food and Drug Administration considers gelatin to be a natural, biodegradable, and biocompatible substance that can be utilized to deliver resveratrol, cycloheximide, doxorubicin, and other medications for tumor therapy [70].

In terms of nanoparticle formation, gelatin has low mechanical strength and a fast breakdown rate. To boost mechanical strength and minimize the rate of breakdown and solubility in aqueous solutions, gelatin nanoparticles must be physically, physiologically, or chemically cross-linked with various cross-linking agents, such as GA [71]. In comparison to uncross-linked particles, cross-linking of GNPs is necessary to provide stability, spherical shape, and increased in vivo circulation time. GNPs that have not been cross-linked have been discovered to be unstable and to agglomerate during storage. Furthermore, substantial applications of GNPs as a drug/vaccine delivery vehicle have been explored in a variety of sectors. However, there is still a serious issue with the usage of animal-derived gelatin, which poses a danger of infection with transmissible spongiform encephalopathy [72].

Experiments on the toxicity of GNPs after administration have been carried out. In a study, paclitaxel-loaded gelatin nanoparticles were used to treat dogs with intravesical bladder cancer, and they achieved a 2.6 higher drug concentration in tumors than control dogs given Cremophor EL with commercial paclitaxel injection. Intraperitoneal injections of doxorubicin-loaded glutaraldehyde cross-linked gelatin nanoparticles into rats were used in the experiment. The electrocardiograms and body weights of the animals were monitored for side effects, and the results revealed that control nanoparticles (no medication) had no toxicity. Although the anticancer drug’s potency was increased, repeated administration of the formulation (doxorubicin-loaded gelatin nanoparticles) revealed substantial cardiotoxicity [73].

In another study, healthy rats were exposed to doxorubicin-loaded gelatin nanoparticles that had been cross-linked by glutaraldehyde. The researchers discovered that linking DXR to gelatin nanoparticles boosted the drug’s cardiotoxicity in the lab. The significant toxicity of DXR-loaded nanoparticles might be due to the drug’s covalent binding to the carrier since DXR was attached to the protein matrix of nanoparticles through glutaraldehyde [74].

Gelatin nanofibers have also been proposed for wound healing processes in tissue engineering. Although electrospun pure gelatin nanofibers have been successfully developed, their poor mechanical characteristics and quick disintegration profile have limited their use in wound healing. Many people have tried to combine gelatin with other natural and synthetic polymers to take advantage of gelatin’s excellent biocompatibility while increasing the mechanical and physical qualities of nanofibers [75].

Ferritin has unique qualities in various disciplines, such as restricted synthesis, nanodevices, disease detection and therapy, drug delivery, vaccine development, and bioassay. These qualities make ferritin a suitable and powerful nanoplatform [76]. Given its presence not just within every cell of the human body but also in the extracellular space and circulating plasma, the iron storage protein ferritin is likely the greatest prospect for therapeutic use among NPs. The production of ferritin nanoparticles in microorganisms necessitates time-consuming purifying procedures. As a result, producing ferritin nanoparticles in large quantities is challenging. Furthermore, ferritin nanoparticles’ high drug loading capacity is limited by their tiny size [77].
The most widely studied technique for anticancer drug delivery is doxorubicin (DXR) encapsulation in ferritin nanocages. DXR is a commonly used cytotoxic medication that has good anticancer activity and is used to treat a variety of solid cancers. However, because it is linked with numerous significant toxicities when supplied at large dosages, its usage is dose-limited. This tailored nanoformulation enhances the drug tumor uptake and accumulation, tumor growth suppression, and circulation half-life and decreased DXR cardiotoxicity [78].

In the experiment conducted by Todd et al., an exceptional tumor inhibition rate of 83.64 percent was re-counted when a genetically improved ferritin derivative apoferritin (RFRTs) was photo-irradiated by a 671-nm laser. Toxicity of the skin is one of the most prevalent side effects of photodynamic treatment; with this RFRT formulation, they found low toxicity to the skin and other tissues [79].

Although ferritin-based drug delivery has been employed for cancer therapy, ferritin was nearly always supplemented with recognition ligands to accomplish tumor-specific targeting in practically all published studies. It causes problems because these additional surface changes undermine the inherent tumor-specific binding of natural ferritin and disrupt its in vivo performance and biocompatibility due to the changed surface physicochemical characteristics of ferritin nanoparticles. Furthermore, many existing approaches for drug loading into ferritin include dismantling ferritin nanocages under a harsh acidic pH, which irreversibly destroys ferritin protein cages and causes harm to the spherical protein surface. Ferritin’s irreparable destruction will significantly impact their in vivo stability and medication delivery efficiency [80].

Despite the fact that numerous researches, in their clinical trials, have revealed that this NP can decrease the harmful effects of cytotoxic medications, there is little information on its harmful effects. Before the probable application of this substance in humans, the safety profile, including pharmacology and toxicology, should be thoroughly examined. The development of gliadin NPs has benefitted medication delivery and controlled release applications [81]. Although the desolvation approach can be used to make gliadin particles, it has significant drawbacks, such as limited drug loading efficiency and the difficulty to separate particles from the aqueous phase [82]. Antisolvent precipitated gliadin NPs are vulnerable to pH, heating, and salt effects, resulting in aggregation and instability. They were particularly susceptible to aggregation at pH 6.0–7.0, with a rapid rise in particle size and even the formation of a precipitate [83].

Furthermore, when it comes to achieving the desired outcome with the usage of gliadin nanoparticles while avoiding serious side effects, it is critical to pay attention to the dosage. TIMP-GLIA tolerance was investigated in mice after immunization by Freitag et al. TIMP-GLIA was given several hours after priming (day 0) and again after immunization on day 7. While 0.025 mg/mouse (1.25 mg/kg) therapy failed to show effect, dosages of 0.25–2.5 mg/mouse (12.5–125 mg/kg) were linked to significant reductions in ear edema when compared to controls. Experiments using gliadin recall also revealed a dose-dependent impact. T cell proliferation was unaffected in spleen cells from animals given 0.025 mg/mouse TIMP-GLIA, but two infusions of 0.25–2.5 mg/mouse/dose resulted in a considerable reduction of T cell growth. TIMP-GLIA had no observed adverse impact level of 75 mg/kg, according to the study [84].

Multiple studies have been conducted to determine the stability of GNPs [85]. In line with the research, the gliadin nanoparticle suspensions showed fair stability to aggregation and sedimentation over a very slim pH range that is from pH 4.5 to 6.0, when exposed to a selection of food processing conditions. Short-term thermal handleings at temperatures above 40 °C disrupted particle suspensions, increasing the size of the particle and sedimentation. The increased hydrophobic attraction between the protein particles and/or the increased particle collision frequency may have caused the instability seen at higher temperatures. The protein nanoparticles’ poor aggregation stability in pH, salt, and temperature conditions often encountered in food products would be a serious issue for their use in many real-world food systems. As a result, more tactics are needed to improve aggregation stability. Coating gliadin nanoparticles with pectin is a viable method of creating stable functional ingredients, particularly for application in the food sector [86].

The creation of fibroin nanoparticles (FNPs) for a variety of biomedical applications has recently received a lot of attention. FNPs can encapsulate a variety of medicinal chemicals, including small and large molecules, proteins, enzymes, vaccines, and genetic elements, due to their versatility and chemical modifiability [87]. Pandey et al. used the MTT assay to assess the cytotoxicity of the medication containing silk fibroin nanoparticles on the rat and human glioblastoma cell lines C-6 and LN-229, respectively. They also calculated the capacity of free and drug-bearing nanoparticles to impede cell growth. SFN-DXR had an IC50 value of 3.68 M and 1.44 M for 24 and 48 h against C-6 rat glioma cell lines, respectively, which was lower than DXR. Similarly, when TSFN (Tween 80–coated silk fibroin nanoparticles) with DXR is compared to free DXR, the TSFN with DXR has a considerably larger cytotoxicity impact [88].

Similarly, Mishra et al. used solvent precipitation procedures to make curcumin-loaded SF nanoparticles and tested their efficiency in triggering apoptosis in a metastatic breast cancer cell line in vitro. Curcumin release from nanoparticles indicated an initial burst release of more than 50% in the first 24 h, with curcumin release continuing for up to 7 days. Although lower concentrations of curcumin-loaded SF nanoparticles seemed to be less hazardous after 96 h of
treatment resulted in 10 times greater than the polyphenol delivery as an oral solution, thus proving the hypothesis of improved efficacy of casein nanoparticles as oral carriers [94].

Toxicity and therapeutic efficacy such as the IC50, LD50, and EC50 of the casein can be evaluated by standard pharmaceutical techniques. A 3-month dose-repeated toxicity test was conducted to examine the safety of casein nanoparticles when administered orally to animals. It was reported that the males and females treated with the maximum dose of 500 mg/Kg bw developed comprehensive hyperchloremia after almost a month. A few cases of hypernatremia in the females were also reported. It was concluded that the formation of drug aggregates with casein nanoparticles results in highly localized concentrations at the sites of deposition that are associated with local toxicity [95]. This study indicates the side effects of protein-based nanoparticles, casein, although further studies are required for the detailed analysis of casein nanoparticle’s toxicity and adverse effects.

Polymeric Nanoparticles

Polymeric nanoparticles are made from natural or synthetic polymers and are especially evident in smart drug delivery. They are being studied extensively in the pharmaceutical industry as carriers for controlled/sustained release in medication delivery systems. Safety difficulties, toxicity threats, inadequate biocompatibility, and physiological challenges all contribute to the activity of these nanopharmaceuticals being limited [96]. The downside of these nanoparticles is toxic monomer aggregation, residual material linked with them, and toxic degradation process [97].

Polymeric NPs’ toxicity is affected by quantum size effects, which are linked to oxidative stress, cytotoxicity, and genotoxicity [98]. Upon investigation of the toxicity of a variety of poly(lactide-co-glycolic acid) (PLGA) nanoparticles on human-like THP-1 macrophages, Nadège et al. found that the slightly cytotoxic chitosan polymer conferred significant cytotoxicity to PLGA nanoparticles when used as a nanoparticle stabilizer. Poly vinyl alcohol and poloxamer 188 polymers also conferred significant cytotoxicity to PLGA nano. These findings revealed that stabilizers employed in the formulation of PLGA nanoparticles have a significant toxicological role when used at high concentrations, which could have consequences for local toxicity of PLGA-based nanomedicine [99].

The differential lung toxicity of positively and negatively charged PEG-polylactic acid (PLA) nanoparticles was explored following daily endotracheal instillation of BALB/c mice while examining the impacts of nanoparticle surface charge for polymeric particles. PEG-PLA nanoparticles based on cationic stearylamine induced higher local and systemic harmful effects [100]. Synthetic nanoparticles
and organic solvents that have been demonstrated to be harmful are employed to manufacture these nanoparticles. Natural polymers and synthesis methods using less harmful solvents have been investigated as potential alternatives to synthetic polymers [101]. Because of their poor circulation stability and targeting inefficiency, using polymeric NPs as chemotherapeutic drug delivery vehicles is generally difficult [102]. Multiple experiments have been going on in order to address these issues such as Palanikumar et al. developed biocompatible and biodegradable pH-responsive hybrid NPs. A cross-linked bovine serum albumin shell was added to these nanosystems based on a drug-loaded PLGA core to decrease interactions with serum proteins and macrophages. As a result, the drug-loaded NPs demonstrated strong anti-cancer action both in vitro and in vivo while causing no harm to healthy tissue [103].

Silica Nanoparticles

Properties like ease of surface modification, synthesis, tunable pore size, efficient thermal stability, and biocompatibility make silica nanoparticles a great choice for drug delivery [104, 105]. Recently, silica nanoparticles have been used to deliver cargos like anti-cancer drugs (camptothecin and doxorubicin in breast cancer treatment), nucleic acid, and proteins (e.g., RGD peptide and aptamers) via controlled released method [106–108]. The growing application often demands assessment of potential risks of toxicity associated with the nanoparticles. Some studies have highlighted the toxicity of silica nanoparticles which leads to lung diseases and neurotoxicity and impacts the immune system. The physicochemical properties of silica nanoparticles are known to induce toxicity at cellular level. In a study by Bancos et al., the impact of silica nanoparticles on macrophages was evaluated using the RAW 264.7 cell line. Exposure to 10-nm silica nanoparticles led to reduced phagocytosis and impacted cytokine production [109]. Gonzalez et al. identified that silica nanoparticles induce cytotoxic effects on lung carcinoma cells [110]. Similar studies by Yang et al. highlighted cytotoxicity in human vascular endothelial cells due to silica nanoparticles [111]. Micronucleus assay and in vitro comet-based analysis have been carried out in some studies to understand the damage caused by silica nanoparticles at DNA level. It was observed that silica nanoparticles show dose-dependent genotoxic effects in these studies on lung cells [112]. In another study, Lucarelli et al. used U-937 cells and ELISA-based assay to evaluate the change in response to silica nanoparticles; they observed M1 polarization and altered IL-1 beta and TNF-alpha production [113]. Xifei et al.’s study highlights that silica nanoparticles lead to increased levels of reactive oxygen species and depletion of glutathione at intracellular levels [114]. Thus, it is very important to evaluate and regulate the toxicity assessment of nanoparticles before use in humans.

Challenges and Future Prospects

The field of nanotechnology has seen remarkable improvement post-1980 as many nano-based drug delivery products have been approved in the market. However, a range of challenges are faced before these nano-based agents enter into the market which include efficient development of nano-carriers, toxicity assessment, and ethical and regulatory requirements. The safety and risk ratio both to the patient and to the manufacturer is not yet clearly defined. The risk/benefit ratio itself faces challenges in laying the details as a strong framework for evaluation has not been demarcated. For nano-based products to be used, it is very crucial that the pre-clinical and clinical testing guidelines be set in detail. Nanoscience has recently stepped in drug delivery and understanding of toxicity, pharmacology, and immune responses of these nano-products still need a lot of deep research.

One common challenge in defining the safety and risk of nanoparticles is that they have different properties depending upon their size, surface area, and surface charge, making them difficult to categorize the risks. Moreover, nanoparticles can move within the biological system easily due to their effective biocompatibility. They can cross cell membranes, placenta, blood–brain barrier, and more, thus increasing the chances of non-specific interaction, which can eventually lead to unwanted accumulation and toxicity at cellular level. Clinical trials to assess these risks can be of great advantage, but the current scenario faces challenges of efficient risk assessment, risk management, and risk communication. Sometimes, extrapolation of animal-based studies into humans can be detrimental. For example, a study involving monoclonal antibodies showed severe illness among the human subjects, even though the results were promising in animal subjects. Learning from these kinds of cases needs to be considered when working with nanomedicines. Post-clinical trials, the side effects need to be informed to the respective safety agency to avoid potential risks.

Despite the above challenges, it can be anticipated that the efficiency of nanomedicines will make their demand grow further in coming years. The application of nano-carriers in cancer treatment has attracted the eye of many researchers, which becomes more evident from the fact that most of the FDA-approved nano-based drugs were cancer related. All in all, the coming years will see a rise in the application of nano-based drug delivery agents for a variety of diseases, and there will be significant growth in the market size of nanomedicines.
Conclusion

Over decades, nanoparticles have been considered to be a wonder of modern science due to their wide applications. The use of nanoparticles in biomedicine has very well marked that standard, as superior therapeutic systems have emerged, helping us conquer various diseases and enhance the quality of human life. However, there still exist many challenges to effectively utilize the full potential of nanoparticles. Toxicity of nanoparticles is one the major barriers that need urgent attention. The safety assessment is vital for use in living beings to recognize potential risks and create preventive measures. As we have seen, there is very limited data available for certain types of nanoparticles used in drug delivery. Therefore, before pushing nanoparticles into the therapeutic market, it is crucial to assess the risk to benefit ratio. Another reason is that the property of nanoformulations within a biological system is yet to be studied in great detail. This has made the regulatory authorities take necessary steps which emphasizes understanding the safety and toxicities associated with nanoparticles used in drug delivery systems. Thus, toxicological studies are needed to provide a more detailed analysis of nanoparticles used in drug delivery. It will help researchers to enhance the properties of nanoparticles further to reduce the toxicity and create nanoformulations which are effective but also non-toxic for therapeutic use.

Declarations

Conflict of Interest Swati Sharma, Roza Parveen, and Dr. Biswa Prasun Chatterji declare that they have no conflict of interest.

Human and Animal Rights and Informed Consent This article does not contain any studies with human or animal subjects performed by any of the authors.

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