

Nanotechnology-based approaches in anticancer research

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Abstract: Cancer is a highly complex disease to understand, because it entails multiple cellular physiological systems. The most common cancer treatments are restricted to chemotherapy, radiation and surgery. Moreover, the early recognition and treatment of cancer remains a technological bottleneck. There is an urgent need to develop new and innovative technologies that could help to delineate tumor margins, identify residual tumor cells and micrometastases, and determine whether a tumor has been completely removed or not. Nanotechnology has witnessed significant progress in the past few decades, and its effect is widespread nowadays in every field. Nanoparticles can be modified in numerous ways to prolong circulation, enhance drug localization, increase drug efficacy, and potentially decrease chances of multidrug resistance by the use of nanotechnology. Recently, research in the field of cancer nanotechnology has made remarkable advances. The present review summarizes the application of various nanotechnology-based approaches towards the diagnostics and therapeutics of cancer.

Keywords: cancer, diagnosis, drug delivery, nanoparticle, nanotechnology, treatment

Introduction

Cancer is a highly heterogeneous complex disease that encompasses a group of disorders characterized by continuous indefinite growth.¹ Through the annals of history, the malaise of cancer has ailed humans. Despite impressive advances in cancer biology, it is the leading cause of death worldwide and remains a challenge. There are over 200 different types of cancer reported all over the globe.² In 2008, approximately 12.7 million cancer cases were reported, causing approximately 7.6 million cancer deaths, out of which 64% of the deaths were reported from economically developing countries.³ The complexity of this disease at genetic and phenotypic levels clarifies its clinical diversity and therapeutic resistance. There is a 5-year relative survival rate of cancer patients,⁴ which provides potential opportunities for early diagnosis and improved treatment, which in turn is highly desirable because of widespread occurrence, high death rate, and recurrence after treatment.⁵

Nanotechnology is an interdisciplinary research field developed with an amalgamation of chemistry, engineering, biology, and medicine, and has various useful applications in cancer biology, such as early detection of tumors, discovery of cancer biomarkers, and development of novel treatments.⁶ It is a rapidly evolving and expanding discipline that has gained public and media interest worldwide. Use of nanotechnology in cancer biology has provided hope within scientific communities of developing novel cancer therapeutic strategies. Nanotechnology involves the creation and/or manipulation of materials at nanometer scale, either by scaling up from single

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groups of atoms or by refining or reducing bulk materials into nanoparticles (NPs).⁷ NPs are typically several hundred nanometers in size and can offer unprecedented interactions with biomolecules present both on the cell surface as well as inside the cell.⁸ NPs can be engineered as nanoplatforms for effective and targeted delivery of drugs, and imaging labels by overcoming many biological, biophysical, and biomedical barriers. For in vitro and ex vivo applications, the advantages of state-of-the-art nanodevices such as nanochips and nanosensors over traditional methods are quite obvious.^{9,10} A variety of NPs are used for diagnosis-cum-therapy of different cancer types, by visualizing tumors and carrying out targeted delivery of drugs with reduced toxic side effects. Cancer related examples of nanodevices include quantum dots (QDs), carbon nanotubes (CNTs), paramagnetic NPs, liposomes, gold NPs (GNPs), magnetic resonance imaging (MRI) contrast agents for intraoperative imaging, and a novel NP-based method for high-specificity detection of DNA and protein.^{6,11–15}

Recent advances have led to development of bioaffinity NP probes for molecular and cellular imaging, targeted NP drugs for cancer therapy, and integrated nanodevices for early screening and detection of cancer. These developments raise exciting opportunities for personalized oncology in which genetic and protein biomarkers are used to diagnose and treat cancer, based on the molecular profiles of individual patients. However, several barriers do exist for in vivo applications of nanodevices in preclinical and clinical use of nanotechnology. Amongst them are biocompatibility, in vivo kinetics, tumor-targeting efficacy, acute and chronic toxicity, ability to escape the reticuloendothelial system, and cost-effectiveness.^{6,16} The development of novel nanotechnology-based approaches towards cancer treatment provides a new ray of hope in the cancer research field. The present review article summarizes the application of various nanotechnology-based approaches towards the diagnostics and therapeutics of cancer.

Use of nanotechnology in cancer treatment

Current therapies and their drawbacks

In the past decade, remarkable progress has been made towards understanding the proposed hallmarks of cancer progression and treatment. With time, the cancer burden is changing for combined types as well as individual types of cancer. However, with ever-increasing incidence, the clinical management of cancer continues to be a grim challenge for the twenty-first century. Over the past couple of decades,

a huge amount of detailed data have been amassed regarding the basic biological processes that become perturbed in cancer, such as disturbances in growth-factor binding, signal transduction, gene transcription control, cell-cycle checkpoints, apoptosis, and angiogenesis.¹⁷ These in turn have prompted the search for rational anticancer drugs and produced a record number of novel compounds, currently being used in cancer treatment trials.

A number of targeted drugs are licensed for routine clinical use, including rituximab, trastuzumab, imatinib, gefitinib, bevacizumab, lapatinib, and cetuximab.¹⁷ Present therapeutic approaches are based on rectifying the damaging mechanism of the genes or by stopping the blood supply to the cancer cells or by destroying the cancer cells itself.¹⁸ Conventional treatment options like surgical excision to remove cancerous parts, radiation therapy, and chemotherapy have their own limitations. Surgery cannot be applied for all types of cancers and might result in loss of an organ, coupled with the risk of cancer recurrence. The radiation approach kills cancerous cells, but it also damages the surrounding healthy cells.¹⁸ The most common therapeutic approach, chemotherapy, is used either alone or in combination with other therapeutic approaches that kill cancer cells by drug toxicity or by preventing cell division, either by stopping the nutrient uptake or by inhibiting the mechanism responsible for cell division.¹⁹ However, this approach is gross and rarely successful for advanced stages of cancer, as pharmacologically active cancer drugs reach the tumor site with poor specificity and dose-limiting toxicity.²⁰ Currently available chemotherapeutic agents are time-tested and confer good disease-free survival only for a limited period of time. Nevertheless, nontarget tissue toxicity and drug resistance curtails the utility of these agents. There is scope to develop newer agents or site-specific delivery systems to transfer these chemotherapeutic agents, which can nullify the major obstacles of toxicity and drug resistance.²¹ The destruction of cancer cells with minimum harm to healthy tissues and delivery of high doses of drug molecules to tumor sites for maximum treatment efficacy are the needs of the hour.

Will nanotechnology-based approaches help in cancer treatment?

Cancer nanotechnology is a rapidly growing field and has made a remarkable contribution to treatment strategies by enabling site-specific release of chemotherapeutic agents, based on their physicochemical characteristics and biological attributes.^{21,22} Several stability and drug-payload studies on NP formulations have shown that they are quite stable with

high carrier capacity, and are suitable for administration of both hydrophilic and hydrophobic substances by various routes.²³ They also have the ability to carry loaded active drugs to cancer cells by selectively utilizing the unique pathophysiology of tumors.²² Due to the advances in synthetic chemistry over the last few decades, different biological nanomaterials have been developed, which can be used for a variety of biological therapies, such as drug delivery, cancer diagnosis, treatment, and imaging.²⁴

The nanotechnology-based drug-delivery system (NDDS) targeted specifically towards cancer cells has several advantages over conventional therapies, such as longer shelf life, improvement in biodistribution of cancer drugs, and administration of both hydrophilic and hydrophobic substances, through oral, nasal, parenteral, and intraocular routes.¹⁹

Current NP systems for cancer therapeutics

There are a variety of NP systems currently being explored for cancer therapeutics.²⁵ The material properties of some NP systems have been developed to enhance their delivery to the tumor site directly.²⁶ Functionalization of NPs by the incorporation of a hydrophilic polymer creates a stealth surface from opsonization.²⁶ Currently used NPs in cancer therapeutics include dendrimers, liposomes, lipid NPs (LNPs), polymeric NPs (PNPs), micelles, protein NPs, ceramic NPs, viral NPs, metallic NPs, and CNTs.^{26–28} Despite extensive research on NP systems for cancer therapeutics, there are only a few NDDSs approved by the US Federal Drug Administration and European Medicines Agency. Specifically, the NDDSs that have been approved include liposomal doxorubicin (Myocet; Elan Pharmaceuticals, Cedar Knolls, NJ), PEGylated liposomal doxorubicin (Doxil; Ortho Biotech, and Caelyx; Schering-Plough), PEGylated liposomal daunorubicin (DaunoXome; Diatos), and the recently approved albumin-bound paclitaxel-loaded NPs (Abraxane; Abraxis Bioscience).²⁶

Various approaches towards cancer treatment with different nanomaterials

Targeted drug delivery via nanocarriers

The development of targeted therapy represents an exciting approach towards cancer treatment.²¹ Different targeting strategies suggest the potential effect of NP systems and will possibly revolutionize current methods in cancer treatment. Different significant events in cancer mechanisms like angiogenesis, uncontrolled cell proliferation, and tumor mass are the targets for NP systems.²⁶ The effectiveness of NP carrier systems reflects their ability to reduce the tumor or related

events without damaging healthy tissues. Moreover, the major improvements offered by NP systems include higher efficacy, lesser side effects, site specificity, efficient delivery, and overcoming multidrug resistance (MDR).²⁹

Targeting tumor cells

The most common targeting strategy is ligand-mediated specific interactions between NPs and cancer cell surface. Longer circulation times and easier endocytosis are the significant factors that need to be considered while choosing target moieties for effective delivery of NPs.³⁰ These ligand-targeted NPs are expected to deliver cytotoxic agents selectively and specifically to tumor cells via receptor-mediated endocytosis, thereby enhancing intracellular drug accumulation. A variety of tumor-targeting ligands, such as antibodies, folate or growth factors, and cytokines have been used to facilitate the uptake of carriers into target cells.³¹ Moreover, monoclonal antibodies and antibody fragments can reduce immunogenicity and improve pharmacokinetics.³² Artificially engineered antibodies have also been reported as a conjugate to thermosensitive liposomes (affisomes) and to poly-(D,L-lactic acid)-polyethylene glycol (PLA-PEG) maleimide copolymer for the delivery of paclitaxel.^{33,34}

Internalization in cancer cells is an important step, because it reduces the dispersal of the drug outside the cancerous cell and can enhance the therapeutic potential of the said drug.^{35–37} Passive targeting approaches have also been reported for the delivery of NPs in angiogenesis as a compensatory mechanism of diffusion. It depends upon the properties of NPs, such as, size, surface nature, and circulation half-life.^{19,38} Passive targeting also involves the use of other innate characteristics of NPs (viz charge) capable of inducing tumor targeting. Cationic liposomes are reported to bind by electrostatic interactions to negatively charged phospholipid head groups, preferentially expressed on tumor endothelial cells.^{19,39} Human cervical carcinoma cells devoid of folate receptor are incapable of cellular uptake of folate-conjugated NPs.⁴⁰ Similarly, enhanced antiproliferative potential of epirubicin going from folate receptor (–) to folate receptor (++) cells was also observed.⁴¹ The uptake of folic acid-conjugated doxorubicin by HeLa cells has also been reported.⁴⁰ The anticancer activity of transferring-conjugated solid-lipid NPs of curcumin on the MCF-7 cell line is also evident.⁴² Several such studies suggested the potential of targeted therapeutic nanoparticles (TNPs) as effective anticancer drug-delivery systems.^{2,22,25,40,43} In an in vivo animal study, targeted TNP-delivered paclitaxel was mainly located

in tumor cells, while nontargeted TNP-delivered paclitaxel was detected at intercellular level.⁴⁴

Targeting the tumor microenvironment

It has been debated that the tumor microenvironment enhanced permeability and retention (EPR) effect might be a key rationale for the development of nanoscale carriers to solid tumors. As an outcome of EPR, nanotherapeutics are expected to improve drug and detection probe delivery, have less adverse effects, and thus result in improved detection and treatment of tumors.⁴⁵ Exploiting the abnormal tumor microenvironment for selective and homogeneous delivery of nanomedicines to tumor sites is another way of cancer treatment.⁴⁶

The caveolae (endothelium transcytosis vesicles)-induced accumulation of an albumin-bound nanomedicine (Abraxane) is initiated by the binding of albumin to a cell surface glycoprotein receptor, suggesting that caveolae targeting might provide a universal portal to pump drugs out of the blood into nearby tissue.⁴⁷ Moreover, arginine–glycine–aspartic acid (RGD) motif has been found to exhibit a strong affinity and selectivity for cell surface in many proteins, thus proclaiming it to be a suitable ligand for tumor targeting by therapeutic NPs.⁴⁸

Targeting recurrent and drug-resistant cancers

Due to the lack of specific ligands, the specific killing of recurrent cancer cells by using targeted TNPs is still unexplored. Although many studies have illustrated the potential use of TNPs to minimize drug resistance, the lack of specific ligands for drug-resistant cancer cells limits the application of targeted TNPs to these aggressive populations. However, targeting of well-progressed cancer environments like recurrence and metastasis is still a challenge. Advanced knowledge of the molecular mechanisms of late-stage cancer has given new leads to target metastatic cancer with NP systems. Moreover, PEGylated liposome modified with a fibronectin-mimetic peptide has been reported to target metastatic colon cancer cells (which overexpress integrins $\alpha 5\beta 1$).⁴⁹

Nanotechnology-based drug-delivery systems

Nanotechnology-based systems can be used to deliver therapeutic entities such as small-molecule drugs, peptides, proteins, and nucleic acids, either alone or in combinations.^{48,50} Their properties could be attributed to their small sizes, lower drug toxicity, maximized bioavailability, cell death, and modification of drug pharmacokinetics.^{51–53} Figure 1 depicts

several nanotechnology based drug delivery systems used in recent years. Several anticancer drugs, such as paclitaxel, doxorubicin, 5-fluorouracil, and dexamethasone, have been successfully formulated using nanomaterials.⁵¹ Polylactic/glycolic acid (PLGA)-based NPs have been formulated to encapsulate dexamethasone.⁵¹ It has also been reported that NPs can escape from the vasculature through the leaky endothelial tissue that surrounds the tumor⁵⁴ and can accumulate in certain solid tumors via the EPR effect.⁵⁵ In contrast, tumor-targeted NPs can enter tumor cells from the extracellular space via receptor-mediated endocytosis.⁵⁶ Significant progress has been made in tumor-targeted nanotherapeutics, and some are already in clinical trials or have been approved by the FDA.³⁷ Kang et al studied doxorubicin-loaded solid-lipid NPs on a doxorubicin-resistant breast cancer cell line (MCF-7/ADR).⁵⁷ Their results showed that doxorubicin-loaded solid-lipid NPs efficiently enhanced apoptotic cell death through higher accumulation of doxorubicin in MCF-7/ADR cells compared with free doxorubicin.

Advantages of NP drug-delivery system

1. To overcome lack of selectivity of anticancer drugs: The major goal of targeted therapies is to target the chemotherapeutics to cancer cells, which ultimately reduces the side effects. To decrease the toxicity and to enhance the selectivity of existing drugs, many NDDSs have been developed in recent years that target specific sites either actively or passively.¹⁹
2. To overcome MDR: Cancers such as non-small-cell lung cancer, and rectal cancer may not respond to standard chemotherapy from the beginning, which is known as primary resistance, while some sensitive tumors respond well to chemotherapy drugs initially but develop acquired resistance later. MDR is mostly due to increased efflux pumps in the cell membrane, such as P-glycoprotein (Pgp). The adenosine triphosphate-binding cassette superfamily includes MDR-associated proteins and breast cancer-resistant proteins. Several studies reported NP-based drug-delivery systems to overcome MDR.^{19,58,59} It was reported that paclitaxel-loaded NP-based drugs could be introduced inside the cells without triggering the Pgp pump in human colorectal tumor.⁶⁰ Also, paclitaxel entrapped in emulsifying wax NPs was shown to overcome drug resistance in a human colon adenocarcinoma cell line (HCT-15).⁵¹ The PLGA NPs were reported to moderately reverse MDR activity in MCF-7/ADR cells.⁵⁹ The activity of doxorubicin and Bcl-2-targeted siRNA on multidrug-resistant A2780/AD human ovarian cancer cells was studied by

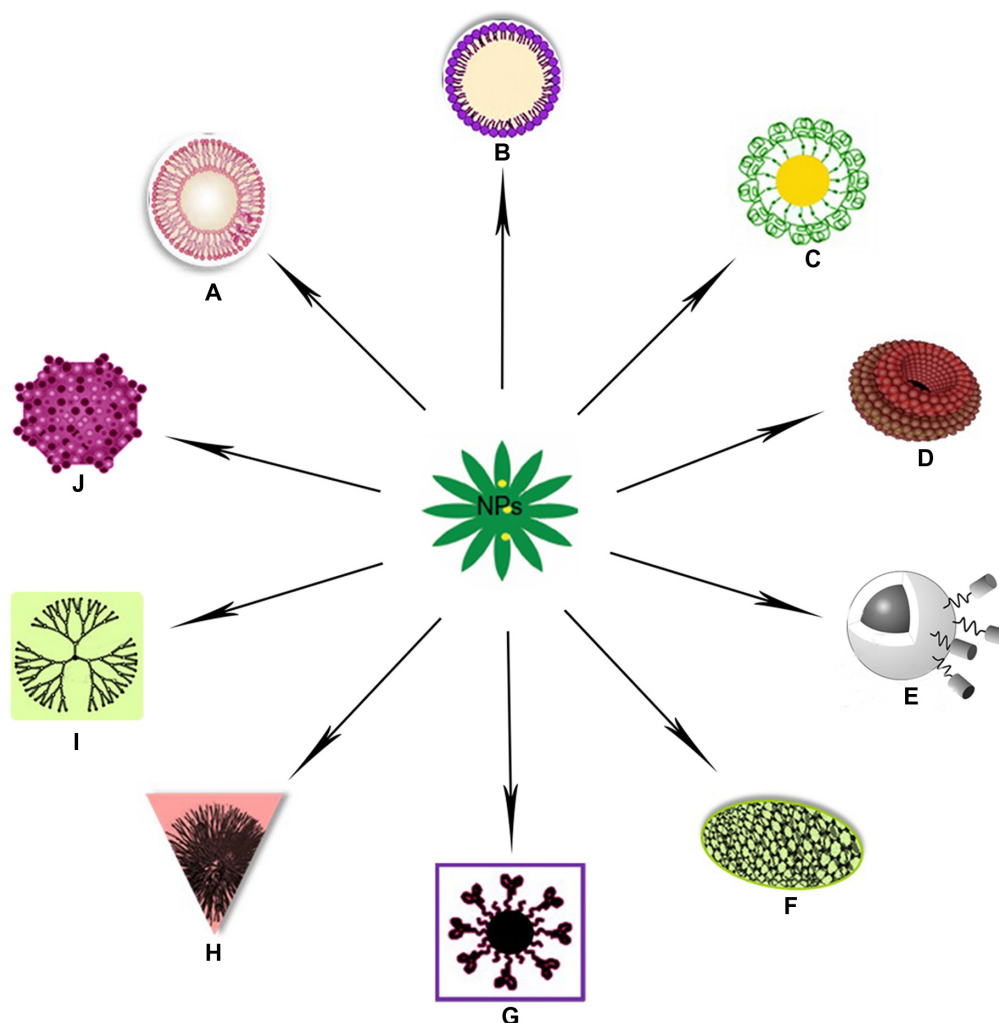


Figure 1 (A–J) Illustration to clarify the described drug-delivery systems of various nanoparticles (NPs). (A) Liposomal nanoparticle; (B) solid lipid nanoparticle; (C) gold nanoparticle; (D) nanodiamond; (E) magnetic nanovector;²²⁰ (F) carbon nanotube; (G) quantum dot nanocarrier; (H) polymeric nanoparticle; (I) dendrimer nanoparticle; (J) virus-mediated nanocarrier.²¹⁹

Chen et al, using mesoporous silica NPs.⁵⁸ They showed that by delivering doxorubicin and Bcl-2 siRNA simultaneously into cancer cells, the Bcl-2 siRNA can effectively silence the Bcl-2 mRNA, significantly suppress non-pump resistance, and substantially enhance the anticancer potential of doxorubicin. Moreover, poloxamer 188 NPs have been reported to overcome MDR in human breast cancer cells.⁶¹

3. To overcome low aqueous solubility of anticancer drugs: Most of the anticancer drugs exhibit poor solubility, which results in reduced bioavailability, increased chances of food effect, frequent incomplete release in dosage form, and higher interpatient variability. Paclitaxel bound to biocompatible proteins is an injectable nanosuspension, that has been approved by the FDA, and is in phase trials for a variety of cancers.^{62,63}

Liposomal nanoparticles as a targeted drug-delivery system

Liposomes are colloidal carriers, formed spontaneously when certain lipids are hydrated in aqueous media. They consist of an aqueous volume entrapped by one or more bilayers of natural and/or synthetic lipids. They are comparatively stable, biocompatible, biodegradable, self-assembled phospholipid membranes with an inner core where hydrophilic drugs could be encapsulated. Liposomes have been suggested as carriers of antineoplastic and antimicrobial drugs, chelating agents, steroids, vaccines, and genetic materials.⁶⁴ They are efficient vehicles for targeted delivery of hydrophobic drugs without eliciting an immune response.²¹ Drugs with widely varying lipophilicities can be encapsulated in liposomes, either in the phospholipid bilayer, in the entrapped aqueous volume, or at the bilayer interface. Drugs encapsulated within this

lipid bilayer are, therefore, protected from extra-liposomal reactions that could alter the effectiveness of the drug.⁶⁵ The lipid bilayer could have a marked effect on the pharmacokinetics and biodistribution of LNPs. In one study, Johnston et al reported significant increase in drug-retention properties in LNPs containing sphingomyelin lipid.⁶⁶ Generally, liposomes have advantages over polymer-based NPs for the formulation of cancer therapeutics.⁶⁷ In most of the cases, the lipid membrane structure mimics the most common structure, which provides a remarkable permeability barrier. Moreover, liposome-encapsulated drugs are protected from extraliposomal reactions, which enhance the effectiveness of the said drug.

LNPs encapsulating therapeutic agents or liposomal nanomedicines represent one of the most advanced classes of drug-delivery systems, with several currently in the market and many more in clinical trials.⁶⁸ There are several examples of LNPs for conventional small-molecule drugs.^{65,69} Curcumin's efficacy as a chemotherapeutic agent against several cancers of the stomach, prostate, breast, and lung has been well documented.^{70,71} Cheng et al reported the development of liposomal curcumin encompassing the anticancer curcumin within liposomal formulations.⁷² In another study, liposome preparations of curcumin exhibited superior antiproliferative and apoptotic effects on six pancreatic cancer cells, and also inhibited pancreatic tumor growth in mouse models.⁷³ Use of the ionophore method of drug loading allows much higher drug:lipid ratios than the conventional citrate method.⁷⁴ This formulation has given excellent results in studies comparing the efficacy of conventional CHOP (chemotherapy treatment composed of cyclophosphamide, doxorubicin, vincristine, and prednisone) and CHOP in which the vincristine has been replaced by LNP-vincristine.^{75,76} Moreover, increased drug retention of vincristine has been reported to be related to increased drug:lipid ratios.⁷⁵ Lipid-based nanocapsules have given a novel approach to scientific communities for encapsulation of poorly soluble drugs like cisplatin.²¹

Drug delivery via carbon nanotubes

CNTs have shown great promise as nanoscale vehicles for targeted drug delivery.² In the past, there have been numerous experimental studies performed *in vitro* and *in vivo* using antibody-functionalized CNTs loaded with chemotherapeutic agents.^{2,77–79} It has been suggested that CNTs could be used as nanocarriers for delivering drugs into the body via intravenous routes.⁸⁰ Drugs can either attach to the outer surface of CNTs via functional groups or be loaded inside the CNTs.² Nanoscale size and ease of cellular uptake makes

single-walled carbon nanotubes (SWCNTs) useful for drug delivery, and their photothermal effects make them potentially useful in a wide range of applications. The poor solubility of SWCNTs could be overcome by functionalization of the surface of the tubes. Multiwalled carbon nanotubes (MWCNTs) have been reported to possess shorter incubation time and relatively higher cytotoxic potential. MWCNT-g-PCA-PTX compared with free drug suggests improved cell penetration of MWCNT-g-PCA-PTX.⁸¹ The grafting of drugs to SWCNTs can be used as a new tool and useful method for potential drug delivery in cancer patients.⁸²

Drug delivery via nanodiamonds

Nanodiamonds (NDs) are attractive agents for use in medicobiological applications, largely due to their greater biocompatibility than other carbon nanomaterials, stable photoluminescence, commercial availability, minimal cytotoxicity, and ease of purification.^{83,84} NDs could be functionalized and conjugated to a variety of molecules for the purpose of cell labeling and drug delivery that can improve their solubility, direct them to specific binding sites on target cells and tissues, and reduce their effects on normal tissues.⁸³ The diversity of ND functionalization broadens the scope of their potential diagnostic and therapeutic applications. They are suitable for controlled drug-delivery applications because of their capability to release drugs slowly and consistently and have abundant capacity for drug loading due to their large surface area:volume ratio.^{85,86} A thin film made of ND clusters loaded with doxorubicin was found to effectively release the drug and kill the target cells.⁸⁵ They could also be used to solubilize and efficiently deliver water-insoluble chemotherapeutic agents to breast and liver tumor cells.^{58,86} NDs can be used for cell labeling and tracing because they do not interrupt cell division or differentiation and have less cytotoxicity as well. They have successfully been used as biomarkers or tracers to label or trace HeLa cells, lung cancer cells, and murine fibroblasts.^{84,87}

Virus-mediated nanocarriers for drug delivery

Different virus-based nanocarriers for drug delivery have also been reported in scientific literature as an emerging nanocarrier platform.^{43,88–90} The versatile hierarchical assembly of viral coat protein subunits provides a natural and easy way of drug packaging. Virus-like particles (VLPs) can easily meet the requirements needed for a drug nanocarrier system, such as biocompatibility, water solubility, and high uptake efficiency. Moreover, VLPs can be modified with polymers such as PEG to improve their half-life in the host

by moderating their immunogenicity.⁴³ Different viruses such as cowpea chlorotic mottle virus (CCMV), a member of the bromovirus group of the Bromoviridae,⁴³ cowpea mosaic virus, and red clover necrotic mosaic virus⁸⁸ have been reported as viral nanocarriers. Recently, nanocarriers based on CCMV have also been suggested.⁸⁹ The incorporation of CdSe/ZnS semiconductor QDs into brome mosaic virus for the design of intracellular microscopic probes and vectors have also been reported.⁹⁰ GNPs could also be encapsulated in VLPs like MS2 bacteriophage in a similar way.⁹¹

Magnetic nanovectors for drug delivery

Use of NP platforms holds the promise of novel and more effective site-specific delivery of therapeutic agents to tumors by magnetic vectoring of magnetically responsive NPs.⁹² Nanovectors based on the three ferromagnetic elements Co, Ni, and Fe can be used either to mediate a hyperthermic effect (predominantly causing tumor necrosis) or to deliver drugs to achieve intratumoral levels. Nickel-based nanomaterials have also been reported for tumor targeting by serving either as inducers of hyperthermia in response to an externally applied magnetic field (as drug-delivery platforms)⁹³ or directly as proapoptotic mediators.⁹⁴ Liu et al synthesized a Co- and Cu-based nonviral carrier for DNA transfer in gene therapy.⁹⁵ Strong binding to DNA involves electrostatic attractions as well as intercalation of the ligands between DNA base pairs, and these complexes efficiently condensed free DNA into globular NPs. Magnetite NPs have reportedly been loaded with daunorubicin to overcome MDR of K562-n/VCR human erythroid leukemia cells.⁹⁶

Enzyme-responsive nanoparticles for drug release

Enzyme-responsive NPs (ENPs), such as polymer-based NPs, liposomes, GNPs, and QDs have been introduced recently and the modulation of their physicochemical properties by the enzymes has been accentuated. Unique physical properties of nanomaterials combined with ENPs are designed with high specificity for triggering the stimulus in drug-delivery schemes.⁹⁷ Hydrolases are the most widely used enzymes for drug delivery, probably due to their facile design.⁹⁸ The attachment of bioactive moieties to carrier through enzyme-cleavable units triggers drug delivery from NP-based carriers.⁹⁹ Similarly, the dispersion of inorganic NPs can be triggered by a hydrolase when the NPs are assembled by biomolecules presenting cleavable units. The rational design and development of a novel enzyme-responsive, doxorubicin peptide-coated magnetic silica NP was conjugated for selectively triggering intracellular

delivery of doxorubicin into the tumor cells with specific protease enzyme expression.¹⁰⁰ The highly efficient doxorubicin is released by the said ENP conjugate upon the specific enzyme interactions in vitro. Some hydrolase-responsive nanomaterials are already under clinical trials.^{98,101} The utilization of oxidoreductase is still in the proof-of-concept stage, and some pioneering examples of their utilization for drug delivery and diagnostics are being debated.⁹⁹ Other enzymes such as kinases and acetyltransferase have also been explored recently in this field.⁹⁹

Dendrimers as nanoparticle drug-carrier system

Dendrimers are a unique class of repeatedly branched polymeric macromolecules with numerous arms extending from the center, resulting in a nearly perfect 3-D geometric pattern.¹⁰² They comprise a series of branches around an inner core, the size and shape of which can be modulated as per the requirement. NPs based on dendrimers and oligonucleotide-linked dendrimers are expected to improve the therapeutic index of cytotoxic drugs by direct delivery to cancerous cells.¹⁰³ There is also hope that delivering drugs by this approach could overcome drug resistance in tumor cells via bypassing Pgp pumps,¹⁰³ which could serve as an attractive modality for drug delivery.^{104–106}

The following properties are associated with dendrimers:^{102,107}

1. Nanoscale sizes that have similar dimensions to important biobuilding blocks.
2. Numbers of terminal surface groups suitable for bioconjugation of drugs, signaling groups, targeting moieties, or biocompatibility groups.
3. Surfaces designed with functional groups to augment or resist transcellular, epithelial, or vascular biopermeability.
4. An interior void space may be used to encapsulate small-molecule drugs, metals, or imaging moieties and facilitates controlled release.
5. Positive biocompatibility patterns that are associated with lower generation of anionic or neutral polar terminal surface groups compared with higher-generation neutral apolar and cationic surface groups.
6. Non- or low immunogenicity associated with most dendrimer surfaces modified with small functional groups or PEG.
7. Surface groups that can be modified to optimize biodistribution, receptor-mediated targeting, therapy dosage, or controlled release of drug from the interior space.
8. Ability to arrange excretion mode from body.

The possibility of a 2,2-bis(hydroxymethyl) propanoic acid-based dendritic scaffold as a delivery carrier for doxorubicin *in vitro* and *in vivo* has been explored.¹⁰⁸ In an attempt to improve the efficacy of doxorubicin, photochemical internalization technology was utilized for site-specific delivery of membrane-impermeable macromolecules from endocytic vesicles into cytosol.¹⁰⁹ Many researchers have also explored the feasibility of cisplatin incorporation in dendrimers.^{102,110,111} Polyamidoamine dendrimer complexes are reported to be used for DNA delivery to cell nucleus due to their high transfection efficiency and very low toxicity.¹⁰² They are favorite candidates to be used as the backbone of multitasking therapeutics because of their well-defined surface functionality, good water solubility, low polydispersity, and lack of immunogenicity.¹⁰² One of the early examples of polyamidoamine dendrimer having a fairly water-soluble nanoformulation has the ability to release cisplatin slowly *in vitro*.¹¹² PEG-dendrimers are generally synthesized by the conjugation of PEG or polyethylene oxide chains to a multifunctional dendritic chain.¹¹³ They constitute a subclass of dendrimers that has attracted numerous researchers because of their prolonged blood circulation time, lower level of toxicity, and relatively lower accumulation in different organs.¹¹⁴

Targeted drug delivery using gold nanoparticles

GNPs are nanometer-sized colloidal suspensions. Multifunctional GNPs have been demonstrated to be highly stable and versatile scaffolds for drug delivery, due to their unique size coupled with their chemical and physical properties.¹¹⁵ The multiple receptor targeting, multimodality imaging, and multiple therapeutic entities holds the promise for a “magic gold bullet” against cancer.⁶ Different subtypes of GNPs such as gold nanospheres, nanorods, nanoshells, nanocages, and SERS NPs have been reported in literature based on their size, shape, and physical properties.^{6,116,117} Colloidal GNPs have great potential to overcome delivery limitations because of their biocompatibility, low toxicity, small size, and tunable surface functionalities.¹¹⁸ In cancer cell cultures, surface properties have been shown to regulate cellular uptake, intracellular release, and distribution in subcellular compartments.¹¹⁸ Modification of the surface properties of GNPs has potential to control accumulation in tumors, where drug payloads are released. GNPs functionalized with a thiolated PEG monolayer capped with a carboxylate group showed an improvement in platinum-based anticancer drugs, such as cisplatin, carboplatin, and oxaliplatin efficacy.¹¹⁹ Recently, GNPs conjugated with bovine serum albumin

have been reported as one of the best vehicles for drug release.¹²⁰ A modification of GNPs was reported, in which 11-mercapto-undecanoic acid-modified GNPs conjugated with chloroquine act as an efficient cancer therapy.¹²¹

Quantum-dot nanocarrier systems for drug delivery

QDs are spherical light-emitting NPs composed of a semiconductor material. They are a promising class of fluorescent probes for cancer screening from biological fluids, classification of tumor from biopsies, and for high-resolution biomolecular and cellular imaging.^{122,123} Initial results displayed by QDs synthesized by using reduced glutathione and TGA co-capped CdTe with good biological compatibility and high fluorescence intensity are quite encouraging for the treatment of colorectal cancer.¹²⁴ Novel folate-conjugated carboxymethyl chitosan-ferroferrocene oxide-doped cadmium telluride QD NPs were developed with high drug-loading efficiency, low cytotoxicity, and favorable cell compatibility, and are promising candidates for carboxymethyl chitosan-based targeted drug delivery and cellular imaging.¹²⁵ Codelivery of siRNA and chemotherapeutic agents has been developed recently to combat MDR in cancer therapy. Several QDs functionalized by β -cyclodextrin coupled to amino acids have been reported recently, some of which can be used to facilitate the delivery of siRNA. The intrinsic fluorescence of the QDs provides an opportunity to track the system by laser confocal microscopy. These multifunctional QDs are promising vehicles for the codelivery of nucleic acids and chemotherapeutics, and for real-time tracking during cancer treatment.¹²⁶ Recent advances have led to multifunctional NP probes that are highly bright and stable under complex *in vivo* conditions. Polymer-encapsulated QDs are essentially nontoxic to cells and small animals, but their long term *in vivo* toxicity and degradation needs to be carefully studied. Nonetheless, bioconjugated QDs have raised new possibilities for ultrasensitive and multiplexed imaging of molecular targets in living cells and animal models.¹²³ Moreover, magnetic NPs and QDs have been used effectively to achieve early diagnosis and recurrence prevention of lung cancer micrometastases in peripheral blood.¹²⁷

Solid-lipid nanoparticles as targeted drug-delivery system

SLNs are physiological lipid-based NP systems that offer physical stability, protection of labile drugs from degradation, easy preparation, and low toxicity.⁵⁷ SLNs have been developed as an alternative system to the existing traditional carriers, such as liposomes and polymeric NPs. They are new-generation submicron-sized lipid emulsions where

the liquid lipid has been substituted by a solid lipid. SLNs offer unique properties, such as small size, large surface area, high drug loading, and the interaction of phases at the interfaces, and are attractive for their potential to improve performance of pharmaceuticals, nutraceuticals, and other materials.¹²⁸ Recently, several studies reported the use of SLNs as tumor-targeted drug-delivery systems.^{42,57,129,130} SLNs loaded with doxorubicin were studied in Pgp-over-expressing MCF-7/ADR cell lines.⁵⁷ Efficiently enhanced apoptotic cell death via higher accumulation of doxorubicin by SLNs in cancer cells was also reported.⁵⁷ Targeted delivery of a potent anticancer compound, namely curcumin, incorporated with transferrin-mediated SLNs has also been reported in MCF-7 breast cancer cells.⁴² More recently, high-pressure homogenized emodin-loaded SLNs gave positive antitumor activity *in vitro*.¹²⁹ Moreover, cationic lipid-bound oligonucleotide-loaded SLNs are also reported as a potential new approach for carrying anti-miRNA inhibitors for cancer therapy.¹³⁰

Polymeric nanoparticles as targeted drug-delivery system

PNPs offer a promising means of targeted drug delivery of chemotherapeutic drugs with enhanced efficacy, reduced toxicity, controlled and long-term release rates, prolonged bioactivity, increased patient compliance due to less administration frequency, and the ability to codeliver multiple drugs with synergistic effects at the same site.¹³¹ A protective covering provided by the PNPs to chemotherapeutic drugs limits their interaction with healthy cells.¹³¹ PNPs are available in different types, such as nanospheres, in which the particles' entire mass is solid and molecules may be adsorbed at the sphere surface or encapsulated within the particle matrix, and nanocapsules, in which the entrapped substances are confined to a cavity consisting of a liquid core (either oil or water) surrounded by a solid material shell.¹³² The properties of PNPs need to be optimized depending on the particular application. In order to achieve the properties of interest, the mode of preparation plays a vital role.¹³²

There is scope for many variations in polymer chemistry where PNPs can be easily manipulated without loss of their desired physical, chemical, and biological properties.¹³³ PNPs are often produced by pairing PEG with the polymer of NPs. PEG inhibits binding of plasma proteins to the surface of PNPs, which provides prolonged systemic circulation time, prevents recognition by the reticuloendothelial system,¹³⁴ and an EPR effect in different types of tumors.¹³¹ Hu et al reported prolonged circulation and enhanced anticancer potential of

one anticancer agent, endostar, a novel recombinant human endostatin, which was approved by the Chinese State Food and Drug Administration in 2005 with PNPs (PEG-PLGA nanoparticles).¹³⁵

PNPs coupled with ligands and aptamers have been reported as a way to actively target cancerous cells that further induce receptor-mediated endocytosis for intracellular delivery. A novel injectable formulation of an anticancer drug, sirolimus, using PNP has been reported as an attractive new therapeutic approach for cancer therapy that offers improved pharmacokinetic features and gets readily dispersed in physiological media without any surfactants or cosolvents. PNP-sirolimus effectively inhibited tumor cell proliferation and tumor mass growth and enhanced radiation-induced cell death by inhibition of the mammalian target of rapamycin (mTOR), mTOR pathway and activation of autophagy.¹³⁶

Anticancer drugs loaded in PLGA-lecithin-PEG NPs and functionalized with AS1411 antinucleolin aptamers have been reported as having high encapsulation efficiency and superior sustained drug release for differential targeted drug delivery in cancer treatment.¹³⁷ Aptamer-functionalized PEG-PLGA nanoparticles have been reported with enhanced delivery of paclitaxel as a therapeutic application in the treatment of gliomas.¹³⁸ Targeted delivery of an anticancer agent, cisplatin, against prostate cancer cells by aptamer-functionalized Pt(IV) prodrug-PLGA-PEG NPs have been also reported with higher efficiency compared with free cisplatin.¹³⁹ Enhanced cellular uptake of folic acid-conjugated PLGA-PEG nanoparticles loaded with vincristine sulfate have been reported to contain significantly higher cellular uptake in the folic acid receptor over expressed MCF-7 cells, compared to PLGA-mPEG NPs without folic acid modification.¹⁴⁰ A novel redox-sensitive biodegradable polymer with "trimethyl-locked" benzoquinone has been reported for the preparation of paclitaxel-incorporated NPs to release paclitaxel in response to chemically triggered reduction.¹⁴¹ With further development in these technologies and other methods of triggering and targeting, polymer nanomaterials will provide improved cancer treatment methods.

Use of NPs in cancer treatment via thermal ablation

There is a great demand for minimally invasive options to conventional surgery like thermal ablative technologies for the treatment of cancer.¹⁴² Radio-frequency, microwave, and laser-based hyperthermia allow less invasive treatments but still require insertion of a probe into the lesion to be treated.¹⁴³ Recent advances in nanoscale materials have

provided a potentially noninvasive means of heating cells to cytotoxic levels. Temperatures above 42°C induce cell death in some tissues. Cells of any nature (cancerous or normal) heated to temperatures in the range of 41°C–47°C begin to show signs of apoptosis,¹⁴⁴ while temperatures above 50°C are associated with less apoptosis and more necrosis.¹⁴⁵ Hyperthermic adjuvant chemotherapy plays an increasing role in multimodality cancer treatment.¹⁴⁶ It has been suggested that minimal tissue hyperthermia increases the blood flow and yields higher concentrations of chemotherapeutic agents, which results in enhanced therapeutic effect. Similar to chemotherapy, radiotherapy is also more effective using hyperthermia as an adjunct treatment.¹⁴⁵ Targeted hyperthermia is achieved by using nanoscale metallic particles that convert electromagnetic energy into heat. Metal NPs exemplify the potential for application in targeted hyperthermic therapy, specifically iron oxide NPs, gold-silica nanoshells, solid-gold NPs and CNTs.^{145,147} Iron oxide NPs have been used as both diagnostic and therapeutic nanoscale materials to treat deep tissue tumors. With certain drawbacks, iron NPs continue to be actively investigated because of their minimal toxicity and potential for rapid heating.¹⁴⁸ GNPs could be heated with shortwave radio-frequency fields. By labeling GNPs with antibodies against particular cancer cells, higher concentrations of GNPs could be achieved. Once the particles are internalized, radio-frequency fields applied to cells result in localized heat and killing of cancer cells.¹⁴⁵

CNTs are capable of being targeted and delivered to specific cells either through direct covalent functionalization or through noncovalent wrapping of targeting moieties. The absorption characteristics of CNTs have been utilized as hyperthermic enhancers using near-infrared absorptions.¹⁴⁹ CNTs significantly absorb intense heat release under capacitively coupled radio-frequency fields, similar to GNPs.¹⁵⁰ QDs also have great potential in photodynamic therapy, where they act as either photosensitizers themselves or as a carrier.^{151,152} Although nanotechnology is a relatively new field of investigation, using targeted NPs in the hyperthermic treatment of cancer cells is expected to be a viable option for cancer treatment.¹⁴⁵ Moreover, these days application of nanotechnology for laser thermal-based killing of cancerous cells, targeted with absorbing NPs, is becoming an extensive area of research.¹⁵³

Use of nanotechnology in cancer treatment via gene therapy

Gene therapy can be defined as the transfer of genetic material to the cells of an individual, in order to ensure a

targeted molecular intervention and achieve a higher level of specific action than conventional cytotoxic chemotherapy. Recently, the gene-silencing technique has gained much popularity. There are three major nucleic acid-based gene-silencing molecules viz antisense oligodeoxyribonucleic acids, siRNA, and micro RNA. Despite so much promise of gene therapy, a number of difficulties remain to be conquered; the most important is the need for more efficient gene-delivery systems. The success of a gene therapy largely depends upon the activity induced by the target genes and the efficiency of gene delivery resulting from the combined effects of the delivery vector and the applied delivery route.¹⁵⁴ Various NPs have been investigated for gene delivery, including GNPs,¹⁵⁵ silver NPs,¹⁵⁶ CNTs,¹⁵⁷ liposomes,¹⁵⁸ polymersomes,¹⁵⁹ and polyplexes. Use of NPs enables targeting of tumor tissue through the EPR effect in gene therapy. Kim et al reported a mannosylated chitosan NP-mediated cytokine gene-delivery system for cancer immunotherapy.¹⁶⁰ Recently, thiolated chitosan NPs as a delivery system for antisense therapy has also been reported in T47D breast cancer cells.¹⁶¹ A nanosized immune liposome-based delivery complex was also reported, which preferentially targets and delivers molecules useful in gene medicine, including plasmid DNA and antisense oligonucleotides to tumor cells.¹⁶² This tumor-targeting NP delivery vehicle can also deliver siRNA to both primary and metastatic cells.

A gene-delivery vehicle, nonviral vector DOTAP: cholesterol has been suggested as an effective systemic gene-delivery vector that efficiently delivers tumor-suppressor genes to disseminated lung tumors.¹⁶³ A promising role of novel nonviral gene carrier heparin-polyethyleneimine (HPEI) NP-mediated ms-T34 A in C-26 colon carcinoma therapy was also reported.¹⁶⁴ PEI-derived NPs such as poly(ϵ -caprolactone)-pluronic-poly(ϵ -caprolactone)-grafted PEI (PCFC-g-PEI), folic acid-PCFC-isophorone diidocyanate-PEI and HPEI are reported as gene-therapy delivery systems for renal cell carcinoma.¹⁶⁵ It has been reported in the literature that QDs may also function as gene-delivery vehicles if modified with lipofectamine,^{166,167} encapsulated in poly(maleic-anhydride-alt-1-decene), surface-modified with dimethylamino propylamine and poly(maleic-anhydride-alt-1-decene) coated. QDs have been reported as siRNA delivery vehicles and exhibit up to a 20-fold increase in silencing effect and a sixfold decrease in toxicity compared with other common delivery agents.¹⁶⁷

Gene delivery could also be achieved by the use of CNTs.² They are a good nonviral vector for gene therapy,

and can cross the cell membrane by an endocytosis process.¹⁶⁸ CNTs are promising vectors for gene and protein, for which they have shown much greater fluorescent activity of protein and DNA when conjugated to single-walled NTs.¹⁶⁹ The delivery of siRNA for treatment of tumor cells using functionalized MWCNTs and liposomes showed that the siRNA delivered via MWCNTs achieved significant inhibition of tumor growth.¹⁷⁰ Dendrimers have also been reported to enter tumors and carry either chemotherapeutic agents or genetic therapeutics. They could be conjugated to fluorochromes and possess the ability to enter into cells. They can then be detected within the cell in a manner compatible with sensing apparatus for evaluation of physiologic changes.¹⁷¹ DNA-assembled polyamidoamine dendrimer clusters for cancer cell-specific targeting were also reported. Dendrimer-5-fluorouracil (5FU) conjugates were prepared by acetylation, and upon hydrolysis release free 5FU, thus minimizing the toxicity of 5FU.¹⁷² In earlier studies, researchers have shown that cisplatin encapsulated in dendrimer polymers has increased efficacy and is comparatively less toxic.¹⁷³ Novel gene-silencing strategy in gene therapy involving the use of siRNA can be conjugated to phospholipid-functionalized SWNTs using a cleavable disulfide linker, resulting in efficient gene silencing and subsequent death of the targeted cell.¹⁴⁹ Amino-functionalized MWNT-siRNA complexes have shown successful suppression of tumor and prolonged survival in lung tumor in animal model.¹⁷⁰ CNTs cationically functionalized with PEI have been shown to be capable of complexing with siRNA and significantly increasing silencing activity and cytotoxicity.¹⁷⁴ A highly efficient, low-toxic, and specifically targeting gene-delivery vector (H1) was also reported.¹⁷⁵ Peritumoral injection of H1/pIL-2 NPs displayed effective antitumor potential. Researchers have derived PEI-coated virus-like particles from an adeno-associated virus type 2 delivery system for potential therapy of breast cancer.¹⁷⁶

Nanotechnology-based approaches in cancer diagnostics

One of the major reasons behind low survival rates among cancer patients is the failure to detect the disease at an early stage. Modern imaging technologies have made enormous advances. However, the effective early detection of precancerous and neoplastic lesions remains an elusive goal. The challenge is to find a test that can detect clinically apparent cancer cases at an early stage, long before the symptoms become visible. A variety of highly integrated nanotechnology

platforms to diagnose cancer hold considerable promise. NP probes, nanocantilever, nanowire, and nanotube arrays are expected to solve the problem of early detection of different types of cancer.¹³ Nanomaterials possess long-term stability and enable the design of powerful bioassays for simultaneous measurements of multiple markers of a disease.¹⁷⁷ Nanomaterial-based probes involve the use of nanotechnology to meet the demands of in vitro diagnostics for increased sensitivity and rapid detection from complex environmental systems.¹⁷⁸ NPs labeled in vivo signaling moiety offer better molecularly targeted specificity than conventional imaging techniques. Various combinations of multifunctional NPs have been established with imaging agents like radioisotopes, lanthanides, or fluorophores.¹⁷⁹ Hopefully, nanotechnology could lead to a paradigm shift in cancer detection, diagnosis, and therapy.

Some of the existing technologies for cancer imaging include noninvasive imaging techniques, such as computed tomography (CT), magnetic resonance imaging (MRI), positron emission tomography (PET), ultrasound scans, single-photon emission CT, optical imaging, and macroscopically visualizing tumors.¹⁸⁰ Unfortunately, most of these imaging techniques are somewhat limited to detecting abnormalities at the microscopic level. It is possible, however, to combine existing optical imaging technologies with sophisticated NP-based optical contrast agents for high-resolution in vivo cancer imaging. Recently, this strategy has been successfully demonstrated for detecting tumors.¹⁸¹ However, developing reliable early detection approaches from serum, other biological fluids, or any sample obtained through minimal or noninvasive procedures remains of supreme importance.

Various nanotechnology platforms for the detection of cancer

Approaches for the in vivo detection and monitoring of cancer biomarkers

To identify malignancies based on their molecular expression profiles, all imaging technologies require contrast agents, comprising a signal-amplifying material conjugated to a molecular recognition and targeting agent such as an antibody.¹³

In vitro detection

Various novel assemblies of NP systems have been reported as nanoprobe for the early detection of cancer. Nanomaterials such as NPs, nanowires, nanotubes and nanodevices have been explored as ultrasensitive probes to

detect cancer biomarkers.¹⁷⁸ NP probes with molecularly targeted recognition agents might provide information on the presence, relative abundance, and distribution of cancer signatures and markers associated with the tumor microenvironment.¹³ Because of their unique properties, QDs are one of the well-established NPs that cover applications such as cellular imaging, immunoassays, in vivo imaging, and sensing.^{182,183} QD-based microfluidic protein chip detection and optically addressed fluorescence resonance energy transfer (FRET) probes are being used in signal transduction to design the detection systems for nucleic acids, proteins, peptides, and small molecules.^{184,185} The increased number of acceptors linked to QDs can significantly amplify the signal of the target through enhanced energy-transfer efficiency. This ultrasensitive nanoprobe based on the FRET system had the capability to detect low concentrations of DNA, which have great potential to diagnose DNA mutations of tumor cells in clinical samples.¹⁷⁸ Methylation-specific QD-FRET offers great promise for its translational use in early cancer diagnosis, prognostic assessment of tumor behavior, and monitoring response to therapeutic agents in clinical samples.¹⁸⁶ QD-based bioluminescence resonance energy transfer (BRET) conjugated to bioluminescent proteins uses biochemical energy to excite fluorophores and offers additional advantages over FRET system for various bioanalyses.¹⁸⁷ A nanoplatform of protease probing based on the BRET system was designed to detect protease activity in complex biological samples.¹⁸⁸ The sensitivity of this technology has great clinical value to cancer diagnosis and monitoring.

Nanophotonic methods also possess important applications in bioanalyses and spectroscopy using plasmonic NPs and can monitor the molecular binding events and changes in molecular conformation with exquisite sensitivity.¹⁷⁸ The said nanoprobe could be used to follow cell-signaling pathways at the single-molecule level in the living cells, which was not possible by conventional single-molecule-imaging techniques.^{189,190} The multiplex protein detection of potential cancer markers was achieved using highly sensitive giant magnetoresistive sensors capable of detecting even ten magnetic nanotags.¹⁹¹ Recently, a magnetic NP-based method has been reported, which is an alternate to PCR methods and enzyme-linked immunosorbent assays for in vitro detection of viruses and circulating tumor cells in biological samples.¹⁷⁸ Although this technique is fast, sensitive, and suitable in complex biological media, it could be broadly applied to detect different biomarkers and biological species with increased sensitivity and specificity, and might become a truly portable, easy-to-use and low-cost device

for point-of-care in vitro diagnostics of different types of cancer in future.¹⁷⁸

Ex vivo detection

NPs have also shown promise for the ex vivo detection of cancer biomarkers.¹³ The ex vivo detection technique offers the potential advantages of readily identifying the conjugate markers, yielding specific information on their tissue distribution, introducing new protocols that include cell surface, endocellular, and microenvironmental antigens in the same test. Gold-nanoshell-based immunoassays have been developed for the detection of various analytes of subnanogram-per-milliliter quantities.¹⁹² Fluorescent NPs have been used for an ultrasensitive DNA-detection system,¹⁹³ and QD bioconjugates with targeting antibodies have been used to recognize molecular signatures, including *ERBB2*.¹⁹⁴ In one study, fluorophore-laden silica beads were used for the optical identification of leukemia cells in blood samples.¹⁹⁵

Nanotubes, nanowires, and nanodevices

Nanocantilevers, nanowires, and nanotubes also have many applications in biology and medicine because of their unique structures and properties. Use of these devices facilitates the transition from single-biomarker to multiple-biomarker cancer diagnostics, prognostics, and treatment. Nanocantilevers are in the class of nanotechnology-based methods that deflect and change resonant frequencies as a result of affinity-binding of biomarker proteins or nucleic acid hybridization events occurring on their free surfaces.¹⁹⁶ The breakthrough potential afforded by nanocantilevers resides in their extraordinary multiplexing capability.¹⁹⁷ In one study, Majumdar used microcantilevers to detect SNPs in a 10-mer DNA target oligonucleotide without using extrinsic fluorescent or radioactive labeling.¹⁹⁸

Devices based on nanowires are emerging as a powerful and general platform for ultrasensitive, direct electrical detection of biological and chemical species. Nanowire and nanotube arrays might contain several thousand sensors on a single chip, therefore offering even greater multiplexing advantages.¹³ A silicon (Si) nanowire field-effect device was developed, in which distinct nanowires and surface receptors are incorporated into arrays.¹⁹⁹ The capability of Si-nanowire probes for multiplexed real-time monitoring of protein markers with high sensitivity and selectivity in clinically relevant samples opens up substantial possibilities for the diagnosis and treatment of cancer.²⁰⁰ SWCNTs exhibit distinct electrical and spectroscopic properties such as photo luminance in the near-infrared range and strong

resonant Raman scattering.²⁰¹ They promise great potential applications in biological detection and imaging because of their unique Raman properties.²⁰²

Multifunctional nanomaterials

Combined uses of multiple-platform diagnostic nanotechnologies are emerging day by day. Multifunctional nanomaterials are highly sensitive, stable, detectable, biocompatible, and targetable.¹⁷⁸ The development of multiple integrated nanoprobe systems includes metals, oxides, polymers, enzymes, or other components to give the system, required functionality, and specificity.^{115,178} The combination of QDs and iron oxide NPs can create a single NP probe that may lead to clinically useful measurements and images of cancer at molecular levels both *in vitro* and *in vivo*.¹⁰⁰ The combination of metallic NPs and magnetic NPs is another type of multifunctional NP, which is likely to lead to new applications in biomedicine because metallic NPs possess intrinsic properties and functions as optical-contrast agents and probes.¹⁷⁸ Park et al reported the simultaneous dual-mode imaging (MRI and fluorescence imaging) *in vitro* and *in vivo*, using hybrid nanostructures containing magnetic NPs and QDs.²⁰³ A polymeric multifunctional optical nanoprobe platform was developed and successfully demonstrated the applications of multifunctional nanomaterials, such as real-time oxygen sensors, *in situ* optical probes, magnetic pH sensors, and stimuli-responsive magnetic optical probes.²⁰⁴

Recent developments in nanotechnology-based cancer detection

A cyclic RGD peptide-conjugated type II CdTe/CdS QD formulation was reported for targeting and imaging of pancreatic tumor vasculature in live animals.²⁰⁵ Lanthanoid trivalent ion-based luminescence agents have been reported as optical probes that have more potential than QDs.^{206,207} The efficacy of europium-catalyzed SWCNT was reported as a novel excellent visible candidate probe for targeted cellular imaging.²⁰⁷ Lewis et al reported noble metal NPs coated with cerium and luminescent europium complexes for bioimaging and potential biodelivery applications.²⁰⁸

Theranostic MRI is now receiving growing interest in imaging-guided drug delivery, monitoring treatment, and personalized administration.²⁰⁹ Recently, an SWCNT-Au-PEG nanocomposite was reported as an interesting optical theranostic probe for cancer imaging and therapy.²¹⁰ Luminescence-quenched shell cross-linked NPs as photonic nanoprobe are reported to detect a protease, which is a viable marker for cancer imaging *in vivo*.²¹¹ Polysorbate 80-coated

temozolomide-loaded PLGA-based superparamagnetic NPs have been characterized as drug carriers and good MRI contrast agents for diagnosing malignant brain glioma.²¹² A multifunctional nanoprobe is prepared by functionalizing SnO₂ NPs with both folic acid as targeting moiety and a gene probe to inhibit or recognize intracellular miRNA levels for target cell-specific imaging and *in situ* detection of intracellular miRNA.²¹³ A sensitive and selective sensor for detecting colon cancer based on NP covalent modified antihuman epithelial cell adhesion-molecule antibody was also developed.²¹⁴

Recently, a highly efficient multifunctional nanoplateform, multicolor luminescent NaYF₄:Yb³⁺,Er³⁺ upconversion NPs were reported by Liu et al,²¹⁵ which could be utilized in image-guided cancer photodynamic therapy. For the first time, Wang et al reported a synthesis of biocompatible triplex Ag@SiO₂@mTiO₂ core-shell NPs that possessed a combined capacity for fast and multiplexed fluorescence-surface-enhanced Raman scattering F-SERS labeling for imaging as well as drug loading for cancer therapy.²¹⁰ In one study, optimized magnetic-particle imaging tracers using iron oxide NPs traced significant signal intensity and better spatial resolution compared with commercial NPs developed for MRI.²¹⁶ High-contrast imaging can be achieved by gold nanorods with targeted tumor-growth-factor receptors as compared to nontargeted gold nanorods. This can facilitate imaging and demarcating tumor margins during surgical resection.²¹⁷ In one study, hyaluronan-coated superparamagnetic iron oxide NPs with high magnetic relaxivity were reported as a promising system with enhanced uptake and enabled MRI of cancer cells for targeted drug delivery.²¹⁸

Future perspectives

Nanotechnology is a rapidly developing field that has given new hope in the treatment of various diseases. Early detection and treatment of cancer remains a challenge to the scientific community. Moreover, different strategies have been explored in recent years for cancer detection and therapy. Application of nanotechnology in cancer treatment seems to solve these limitations, which has given new hope to humanity. Specific targeting of cancer cells was also the major challenge faced by conventional therapeutic approaches of cancer treatment. Recently, various NP-based drug-delivery systems such as liposomes, dendrimers, diamondoids, QDs, viral NPs, and CNTs have shown encouraging results in cancer therapy. Properties like prolonged existence in systemic circulation, enhanced drug localization, and their efficacy make the NP-based model an excellent one. One of the major challenges in cancer treatment, ie, MDR, can also be overcome by these NP formulations. In

the light of our review, we expect that in future, different NP formulations would serve as “Trojan horses” in the field of cancer diagnostics and its treatment. Hereby, we find it pertinent to highlight that toxicity and immune-system induction should be given due consideration before finalizing the use of any NP formulation for diagnostic and treatment purposes.

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