1

Introduction

Sreekanth Thota 1,2,3 and Debbie C. Crans 3

¹ Fundação Oswaldo Cruz – Ministério da Saúde, National Institute for Science and Technology on Innovation on Neglected Diseases (INCT/IDN), Center for Technological Development in Health (CDTS), Av. Brazil
² Universidade Federal do Rio de Janeiro, Programa de Desenvolvimento de Fármacos, Instituto de Ciências Biomédicas, Av. Carlos Chagas Filho, Rio de Janeiro, RJ 21941-902, Brazil
³ Colorado State University, Department of Chemistry, Fort Collins, Colorado 80523, USA

1.1 History of Metal Complexes

1.1.1 Introduction

Pharmaceutical science, which studies the design, action, delivery, and disposition of drugs, is an important field in drug research. Humans have made several sincere attempts for the search of new drugs in order to cure and control different diseases. Although possible remedial measures are available at present to tackle any disease, scientists are increasingly trying to find superior and more effective drugs [1]. Over the last 50 years some "wonder drugs" have played a crucial role in diminishing the global burden of infectious diseases. New drugs are constantly being screened for their potential biological properties. Among the category of new drugs that are receiving much attention are metal-based drugs [2]. Precious metals have been used for medicinal purposes for at least 3500 years. Among them, gold has played a crucial role in a variety of medicines in China and Arabia [3].

1.1.2 Metal Complexes

Medicinal inorganic chemistry is in the early days of its development, although there are now a significant number of clinical trials involving metal compounds or other agents that interfere with metabolic pathways for metals, both for therapy and for diagnosis [4]. In chemistry, metal complexes are nothing but reactions between metals and ligands [5]. Biomedical applications of several metal coordination compounds in recent years have provided a substantial contribution to the augmentation of more impressive diagnostic and therapeutic agents [6]. Metal coordination compounds and metal ions are known to effect cellular processes in a dramatic way [7]. Metal coordination complexes offer biological and chemical diversity that is distinct from that of organic drugs.

1.1.3 **Metal Complexes in Medicine**

In the ancient history of medicine, extraordinarily, many metal-based drugs played a crucial role as anti-infective agents. The increasing medicinal application of metals and metal complexes day by day is gaining clinical and commercial significance [8]. The development of metals containing anticancer drugs has been in the 1960s with the synthesis of Platinum compounds. Cisplatin is one of the most extensively used antineoplastic drugs, specifically for the treatment of ovarian and testicular cancers [9, 10]. The success of cisplatin and its analogs has accelerated a resurgence of inorganic medicinal chemistry and the search for complexes of other precious metals [Ru, Va, Zn, Cu, Ag, Gold, Pd] with interesting biological properties [11–17]. Among them, particularly ruthenium compounds have attracted significant attention with two compounds, namely, NAMI-A and KP1019, advancing through clinical trials [18]. Many precious metals and metal compounds have succeeded in the clinic over the last few decades. Platinum compounds are the most extensively used chemotherapeutic agents, silver compounds have been useful as antimicrobial agents, and gold compounds are used widely in the treatment of rheumatoid arthritis. Scientists have been investigating over the past 25 years several metal-based compounds and such return of interest in metal-based drugs can be witnessed in several recent articles [19-24].

1.2 Nanotechnology

1.2.1 Introduction

In today's world, nanotechnology is a relatively new field, but its structural nanometer dimensions and functional devices are not new, and in fact, these materials have much significance. In recent years, we found a plethora of literature explaining the recent advances in nanotechnology [25-33]. Nanotechnology has the potential to provide novel, paradigm-shifting solutions to medical problems. Nanotechnology, which has been defined as the engineering and manufacturing of materials at the atomic and molecular scale, offers exclusive tools for developing safer and more efficient medicines (nanomedicines), and provides several potential advantages in drug formulation and delivery. Nanotechnology refers to an emerging field of science that includes preparation and development of various nanomaterials. Nowadays, nanomaterials are widely used in many fields including biomedicine, consumer goods, and energy production [34-37]. The purpose of nanomaterials in biotechnology combines the fields of material science and biology.

Development of Nanotechnology

In recent years, disparate products of nanotechnology have played a key role in adding a novel armamentarium of the rapeutics to the pipelines of pharmaceutical industries. The nanotechnology fever we are experiencing now began when the United States launched the National Nanotechnology Initiative [38], the world's first program of its kind, in 2000. Nanotechnology usage may possibly achieve many advantages: (i) improved delivery of poorly water-soluble drugs; (ii) targeted delivery of drugs in a cell- or tissue-specific manner; (iii) drugs transcytosis beyond the tight endothelial and epithelial barriers; (iv) improved delivery of large macromolecule drugs to intracellular sites of action; (v) co-delivery of multiple drugs or therapeutic modality for combination therapy; (vi) improvement in drug delivery through visualization of sites by combining therapeutic agents with imaging modalities [39]; and (vii) real-time read on the in vivo efficacy of an agent [40]. Nanotherapeutics has the potential to actively target tumors, increasing the therapeutic effectiveness of a treatment while limiting side effects. This improved therapeutic index is one of the great promises of nanotechnology [41].

1.2.3 Nanotechnology in Medicine

In pharmaceutical trade, a new molecular entity (NME) that exhibits significant biological activity but meager water solubility, or a very terse circulating half-life, will likely face significant challenges in progress or will be assumed undevelopable [42]. Nanotechnology may revolutionize the rules and possibilities of drug discovery and change the landscape of pharmaceutical industries. In medicine, nanotechnology application may be referred to as nanomedicine that explains various intriguing possibilities in the healthcare sector. The major current and promising applications of nanomedicine include, but are not limited to, drug delivery, in vivo imaging, in vitro diagnostics, biomaterials, therapy techniques, and tissue engineering [28]. In oncology, nanomaterials can enable targeted delivery of imaging agents and therapeutics to cancerous tissues; nanoscale devices enable multiplexed sensing for early disease detection and therapeutic monitoring. The

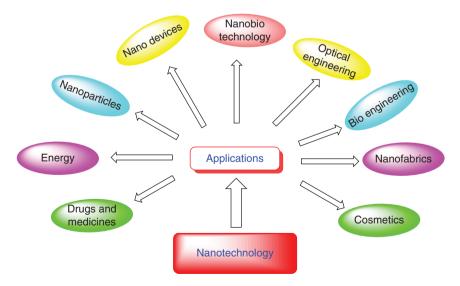


Figure 1.1 Applications of Nanotechnology.

drug delivery field application of nanotechnology is widely expected to change the landscape of pharmaceutical and biotechnology industries in the foreseeable future [40, 43–45]. Nanotechnology attracts scientists because of a wide variety of applications, which includes drugs and medicines, energy, nanoparticles, nanodevices, nanobiotechnology, optical engineering, bioengineering, nanofabrics, and cosmetics (Figure 1.1).

1.3 Nanoparticles

1.3.1 Introduction

Any intentionally produced particle that has a characteristic dimension from 1 to 100 nm and has properties that are not shared by non-nanoscale particles with the same chemical composition has been called a nanoparticle [46, 47]. Nanoparticles demonstrate a particularly useful platform, describing exclusive properties with potentially wide-ranging therapeutic applications [48]. The enormous diversity of nanoparticles was described (Figure 1.2). Nanoparticles made of polymers (NPs) are of particular interest as drug delivery systems because of their synthetic versatility as well as their tunable properties (e.g., thermosensitivity and pH response). Nanoparticles offer exciting prospects for improving delivery, cell

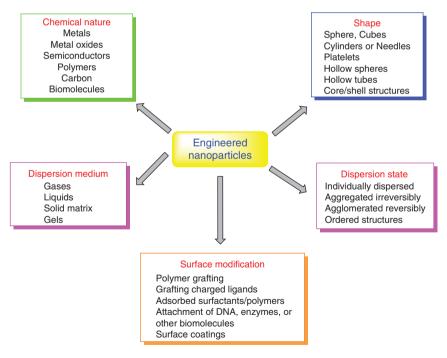


Figure 1.2 Various features of engineered nanoparticles.

uptake, and targeting of metallodrugs, especially anticancer drugs, to make them more effective and safer. Transition metal nanoparticles synthesis has been extensively investigated in recent years because of its many unique physical (electronic, magnetic, mechanical, and optical) and chemical properties. Nanoparticles are often in the range 10–100 nm and this is the same size as that of human proteins.

Development of Nanoparticles

The primary intention in designing nanoparticles as a delivery system is to manage particle size, surface properties, and release of pharmacologically active agents in order to obtain the site-specific action of the drug at the therapeutically optimal rate and dose regimen [49]. Nanoscale particles developed using organic molecules as building blocks have been widely examined for drug and gene delivery. For example, polymer, polymersome, and liposome constructs for controlled release of proteins and polymeric micelles, macromolecules, and long-circulating polymeric nanoparticles are in different stages of clinical and preclinical development [29]. In the 1960s, Bangham and Horne produced the first nanoparticle-based platform for medical application based on liposomes. In the following decades, nanoparticles gathered more scientific and general interest and developed rapidly [50].

1.3.2.1 Liposome-Based Nanoparticles

Liposomes are small sphere-shaped particles, formed by one or more phospholipid bilayers that can be made from cholesterol and natural phospholipids. Depending on the design, they can range from 10 nm up to micrometers [51].

1.3.2.2 Polymeric Nanoparticles

Polymeric nanoparticles might be the most widely used nanoparticle carriers and have been extensively investigated in this regard. They could be formed by biodegradable, biocompatible, and hydrophilic polymers such as poly (D,L-lactide), poly(lactic acid) (PLA), poly(D,L-glycolide) (PLG), poly(lactide-coglycolide) (PLGA), and poly-(cyanoacrylate) (PCA) [52-54].

1.3.2.3 Metal Nanoparticles

Metal nanoparticles are attractive materials in many fields ranging from physics (hard or soft magnetic materials, optics, microelectronics) to catalysis [55]. Noble metal nanoparticles with spherical shape and sharp size distribution such as gold were formed progressively by the chemical reduction method supported by ultrasonic device [56]. The capability to integrate metal nanoparticles into biological systems has had a huge impact in biology and medicine. Some noble metal nanoparticles have been attracting huge interest from the scientific community owing to their awesome properties and diversity of applications, which include gold and silver [57]. The three important properties of gold nanoparticles that have attracted intensive interest are that they are easily prepared, have low toxicity, and readily attach to molecules of biological interest [58].

1.3.3 Nanoparticles in Science and Medicine

Over the past few years nanoparticles have emerged as a key player in modern medicine. Nanoparticles have significance ranging from being contrast agents in medical imaging to being carriers for gene delivery into individual cells [59]. Nanoparticles represent an extraordinarily charming platform for a distinct array of biological significance. In cancer therapy there has been an enormous amount of interest in the preparation and significance of nanoparticles [60]. NPs can easily be conjugated with biomolecules, and thus, they can act as labels for signal amplification in biosensing and biorecognition assays. These strategies can significantly enhance detection sensitivity; even a single molecule can be detected in an ideal case [34]. The exclusive properties and adequacy of nanoparticles emerge from a peculiarity of attributes, including the similar size of nanoparticles and biomolecules such as polynucleic acids and proteins. Additionally, nanoparticles can be formed with a huge range of metal and semiconductor core materials that convey favorable properties such as fluorescence and magnetic behavior [40]. Nanoparticles can afford significant improvements in traditional biological imaging of cells and tissues using fluorescence microscopy as well as in modern magnetic resonance imaging (MRI) of various regions of the body. MRI technique is extensively used in modern medicine, specifically in the diagnosis and treatment of most diseases of the brain, spine, and the musculoskeletal system. Superparamagnetic iron oxide (SPIO) nanoparticles can also be used to visualize features that would not otherwise be detectable by conventional MRI [61]. Several such SPIO nanoparticles have been used in modern MRI [62, 63]. Nanoparticles have already been used for a wide range of applications both in vitro and in vivo. Nowadays various nanoparticles are used in biomedicine. A list of some of the applications of nanomaterials in biology or medicine is given below:

- 1) Drug and gene delivery [64–75]
- 2) Fluorescent biological labels [76–80]
- 3) Detection of proteins [81]
- 4) Biodetection of pathogens [82]
- 5) Medical imaging [83]
- 6) Probing of DNA structure [84]
- 7) Tissue engineering [85]
- 8) Phagokinetic studies [86]
- 9) Tumor destruction via heating (hyperthermia) [87]
- 10) Separation and purification of biological molecules and cells [88]
- 11) Cancer cell imaging [89]
- 12) Treatment of cancer [26, 69, 90, 91].

Some of the potential applications of nanoparticles are in antibacterial creams and powders (Ag), biolabeling and detection (Au, Ag, quantum dots), bone growth promoters (hypoxyapatite ceramics), cancer diagnostics and targeted drug delivery (magnetic nanoparticles, metal nanoparticles), biocompatible coatings for implants, cell, receptor, antigen, and enzyme imaging (quantum dots), MRI contrast agents (Fe₂O₃, Fe₃O₄), gene delivery (CNT), and dental composites [92].

The most widely used nanoparticles in everyday life and in research laboratories are silver nanoparticles (AgNPs). This huge degree of AgNP commercialization has been due to their significant antimicrobial and antifungal properties. Many manufacturers claim that potential AgNP toxicity is minimal or nonexistent. In medical practice silver nanoparticles are commonly used as an integral part of both surgical and nonsurgical equipment such as wound dressings, bandages, and catheters [93].

1.4 Nanotechnology-Supported Metal Nanoparticles

Drug loading into NPs can be achieved by three techniques: (i) covalent attachment to the polymer backbone, (ii) adsorption to the polymer surface, or (iii) entrapment in the polymer matrix during preparation of the NPs. In most cases metallodrug polymer systems have been formulated by covalent attachment of the metal-based drug to the polymer backbone. Drug delivery system efficiency can be optimized; for that, carriers must be sufficiently small for the impressive diffusion of the drug-carrier composite into the targeted cellular environment. Hence, metal nanoparticles, owing to their small size, can be excellent candidates as drug carriers [49, 50]. There has been a great deal of development in the field of gold-nanoparticle-mediated cancer therapy in vitro and in vivo in the last 10 years. In recent years, several metal nanoparticles have been widely used [94-107].

Acknowledgment

This work was supported by National Council for Scientific and Technological Development (CNPq), Coordenação de Aperfeiçoamento de Pessoal de Nível Superior (CAPES), Oswaldo Cruz Foundation (Fiocruz).

References

- 1 Kamatchi, T.S., Chitrapriya, N., Kim, S.K., Fronczek, F.R., and Natarajan, K. (2013) Influence of carboxylic acid functionalities in ruthenium (II) polypyridyl complexes on DNA binding, cytotoxicity and antioxidant activity: Synthesis, structure and in vitro anticancer activity. Eur. J. Med. Chem., **59**, 253-264.
- 2 Kostova, I. (2006) Ruthenium complexes as anticancer agents. Curr. Med. Chem., 13 (9), 1085-1107.
- 3 Allardyce, C.S. and Dyson, P.J. (2001) Ruthenium in medicine: current clinical uses and future prospects. Platinum Metals Rev., 45 (2), 62-69.
- 4 Barry, N.P.E. and Sadler, P.J. (2013) Challenges for metals in medicine: How nanotechnology May help to shape the future. ACS Nano, 7 (7), 5654–5659.
- 5 Turel, I. (2015) Practical applications of metal complexes. *Molecules*, **20** (5), 7951-7956.

- 6 Thota, S., Vallala, S., Yerra, R., and Barreiro, E.J. (2015) Design, synthesis, characterization, cytotoxic and structure activity relationships of novel Ru(II) complexes. Chin. Chem. Lett., 26, 721-726.
- 7 Reedijk, J. and Bouwman, E. (1999) Metalloenzymes with a quinone co factor. Bioinorg. Catalysis., 12, 563-585.
- 8 Guo, Z. and Sadler, P.J. (1999) Medicinal inorganic chemisty. Adv. Inorg. Chem., 49, 183-306.
- 9 Reedijk, J. (1996) Improved understanding in platinum antitumor chemistry. Chem. Commun., 12, 801-806.
- 10 Wong, E. and Giandomenico, C.M. (1999) Current status of platinum based antitumor drugs. Chem. Rev., 99, 2451-2466.
- 11 Clarke, M.J. (2003) Ruthenium metallopharmaceuticals. Coord. Chem. Rev., **236**, 209–233.
- 12 Crans, D.C. (2015) Antidiabetic, chemical and physical properties of organic vanadates as presumed transition state inhibitors for phosphatases. J. Org. Chem., 80 (24), 11899-11915.
- 13 Gao, C.Y., Qiao, X., Ma, Z.Y., Wang, Z.G., Lu, J., Tian, J.L., Xu, J.Y., and Yan, S.P. (2012) Synthesis, characterization, DNA binding and cleavage, BSA interaction and anticancer activity of dinuclear zinc complexes. Dalton Trans., 41, 12220-12232.
- 14 Santini, C., Pellei, M., Gandin, V., Porchia, M., Tisato, F., and Marzano, C. (2014) Advances in copper complexes as anticancer agents. Chem. Rev., 114 (1), 815-862.
- 15 Eckhardt, S., Brunetto, P.S., Gagnon, J., Priebe, M., Giese, B., and Fromm, K.M. (2013) Nanobio silver: It's interactions with peptides and bacteria, and its uses in medicine. Chem. Rev., 113 (7), 4708-4754.
- 16 Shaw III, C.F. (1999) Gold-based therapeutic agents. Chem. Rev., 99 (9), 2589-2600.
- 17 Abu-Surrah, A.S., Al-Sadoni, H.H., and Abdalla, M.Y. (2008) Palladium-based chemotherapeutic agents: routes toward complexes with good antitumor activity. Cancer Therapy., 6, 1-10.
- 18 Rathgeb, A., Bohm, A., Novak, M.S., Gavriluta, A., Domotor, O., Tommasino, J.B., Enyedy, E.A., Shova, S., Meier, S., Jakupec, M.A., Luneau, D., and Arion, V.B. (2014) Ruthenium-nitrosyl complexes with glycine, L-alanine, L-valine, L-proline, D-proline, L-serine, L-threonine, and L-tyrosine: synthesis, X-ray diffraction structures, spectroscopic and electrochemical properties and antiproliferative activity. *Inorg. Chem.*, 53, 2718 - 2729.
- 19 Wang, X., Wang, X., and Guo, Z. (2015) Functionalization of platinum complexes for biomedical applications. Acc. Chem. Res., 48 (9), 2622-2631.
- 20 Simon, P. (2012) Ruthenium compounds as anticancer agents. Educ. Chem., 26-29.
- 21 Willsky, G.R., Halvorsen, K., Godzala, M.E., Chi, L.H., Most, M.J., Kaszynski, P., Crans, D.C., Goldfine, A.B., and Kostyniak, P.J. (2013) Coordination chemistry may explain pharmacokinetics and clinical response of vanadyl sulfate in type 2 diabetic patients. Metallomics, 5 (11), 1491-1502.

- 22 Crans, D.C., Schoeberl, S., Gaidamauskas, E., Baruah, B., and Roess, D.A. (2011) Antidiabetic vanadium compound and membrane interfaces: interface facilitated metal complex hydrolysis. J. Biol. Inorg. Chem., 16, 961–972.
- 23 Allardyce, C.S. and Dyson, P.J. (2016) Metal-based drugs that break the rules. Dalton Trans., 45, 3201-3209.
- 24 Naggar, M.E., Shehadi, I., Abdou, H.E., and Mohamed, A.A. (2015) Gilded hope for medicine. Inorganics., 3, 139-154.
- 25 Farokhzad, O.C. and Langer, R. (2009) Impact of nanotechnology on drug delivery. ACS Nano, 3 (1), 16-20.
- 26 Abadeer, N.S. and Murphy, C.J. (2016) Recent progress in cancer thermal therapy using gold nanoparticles. J. Phys. Chem. C, 120 (9), 4691-4716.
- 27 Dreaden, E.C., Alkilany, A.M., Huang, X., Murphy, C.J., and El-Sayed, M.A. (2012) The golden Age: gold nanoparticles for biomedicine. Chem. Soc. Rev., **2012** (41), 2740–2779.
- 28 Shi, J., Votruba, A.R., Farokhzad, O.C., and Langer, R. (2010) Nanotechnology in drug delivery and tissue engineering: from discovery to applications. Nano Lett., 10, 3223-3230.
- 29 Huang, H.C., Barua, S., Sharma, G., Dey, S.K., and Rege, K. (2011) Inorganic nanoparticles for cancer imaging and therapy. J. Controlled Release, 155, 344-357.
- 30 Park, K. (2013) Facing the truth about nanotechnology in drug delivery. ACS Nano, 7 (9), 7442-7447.
- 31 Kratz, F. and Warnecke, A. (2012) Finding the optimal balance: challenges of improving conventional cancer chemotherapy using suitable combinations with nano-sized drug delivery systems. J. Controlled Release, 164, 221 - 235.
- 32 Weiss, P.S. (2010) Nanoscience and nanotechnology: present and future. ACS Nano, 4 (4), 1771–1772.
- 33 Goh, D., Tan, A., Farhatnia, Y., Rajadas, J., Alavijesh II, M.S., and Seifalian, A.M. (2013) Nanotechnology-based gene-eluting stents. Mol. Pharmaceutics, **10**, 1279-1298.
- 34 Benešová, I., Dlabková, K., Zelenák, F., Vaculovič, T., Kanický, V., and Preisler, J. (2016) Direct analysis of gold nanoparticles from dried droplets using substrate-assisted laser desorption single particle-ICPMS. Anal. Chem., 88, 2576–2582.
- 35 Dudkiewicz, A., Tiede, K., Loeschner, K., Jensen, L.H.S., Jensen, E., Wierzbicki, R., Boxall, A.B.A., and Molhave, K. (2011) characterization of nanomaterials in food by electron microscopy. Trends Anal. Chem., 30, 28 - 43.
- 36 Martin, C.R. and Mitchell, D.T. (1998) Peer reviewed: Nanomaterials in analytical chemistry. Anal. Chem., 70, 322A-327A.
- 37 Gowda, S.R., Reddy, A.L.M., Zhan, X., and Ajayan, P.M. (2011) Building energy storage device on a single nanowire. Nano Lett., 11, 3329-3333.
- 38 The White House Office of the Press Secretary. National Nanotechnology Initiative: Leading to the Next Industrial Revolution. (2000) http://clinton4 .nara.gov/WH/New/html/20000121_4.html.

- 39 Liong, M., Lu, J., Kovochich, M., Xia, T., Ruehm, S.G., Nel, A.E., Tamanoi, F., and Zink, J.I. (2008) Multifunctional inorganic nanoparticles for imaging, targeting, and drug delivery. ACS Nano., 2, 889-896.
- 40 Ferrari, M. (2005) Cancer nanotechnology: opportunities and challenges. Nat. Rev. Cancer, 5, 161-171.
- 41 Farrell, D., Alper, J., Ptak, K., Panaro, N.J., Grodzinski, P., and Barker II, A.D. (2008) Recent advances from the national cancer institute alliance for nanotechnology in cancer. ACS Nano., 4 (2), 589-594.
- 42 Wagner, V., Dullaart, A., Bock, A.K., and Zweck, A. (2006) The emerging nanomedicine landscape. *Nat. Biotechnol.*, **24**, 1211–1217.
- 43 Whitesides, G.M. (2003) The 'Right' size in nanobiotechnology. Nat. Biotechnol., **21**, 1161–1165.
- 44 Zhang, L., Gu, F.X., Chan, J.M., Wang, A.Z., Langer, R.S., and Farokhzad, O.C. (2008) Nanoparticles in medicine: therapeutic applications and developments. Clin. Pharmacol. Ther., 83, 761-769.
- 45 Langer, R. (1990) New methods of drug delivery. Science., 249, 1527-1533.
- 46 Hansen, S.F., Larsen, B.H., Olsen, S.I., and Baun, A. (2007) Categorization framework to aid hazard identification of nanomaterials. Nanotoxicology., 1, 243 - 250.
- 47 Auffan, M., Rose, J., Bottero, J.Y., Lowry, G.V., Jolivet, J.P., and Wiesner, M.R. (2009) Towards a definition of inorganic nanoparticles from an environmental, health and safety perspective. Nat. Nanotechnol., 4, 634-641.
- 48 De, M., Ghosh, P.S., and Rotello, V.M. (2008) Applications of nanoparticles in biology. Adv. Mater., **20**, 4225–4241.
- 49 Mohanraj, V.J. and Chen, Y. (2006) Nanoparticles a review. Tropical Journal of Pharmaceutical Research, 5 (1), 561-573.
- 50 Bangham, A.D. and Horne, R.W. (1964) Negative staining of phospho lipids and their structural modification by surface-active agents as observed in the electron microscope. J. Mol. Biol., 8 (5), 660-668.
- 51 Torchilin, V.P. (2005) Recent advances with liposomes as pharmaceutical carriers. Nat. Rev. Drug Discovery, 4 (2), 145–160.
- 52 Mundargi, R.C. et al. (2008) Nano/Micro technologies for delivering macromolecular therapeutics using poly (D, L-lactide and co-glycolide) and its derivatives. J. Controlled Release, 125 (3), 193-209.
- 53 Zhang, K. et al. (2014) PEG-PLGA copolymers: Their structure and structure-influenced drug delivery applications. J. Controlled Release, 183, 77-86.
- 54 Brannon-Peppas, L. (1995) Recent advances in the use of biodegradable microparticles and nanoparticles in controlled drug delivery. Int. J. Pharm., **116** (1), 1–9.
- 55 Costa, N.J.S., Guerrero, M., Colliere, V., Teixeira-Neto, E., Landers, R., Philippot, K., and Rossi, L.M. (2014) Organometallic preparation of Ni, Pd, and NiPd nanoparticles for the design of supported nanocatalysts. ACS Catal., 4, 1735–1742.
- 56 Kikuo, Okuyama and Wuled, Lenggoro (2004). Nanoparticle Preparation and Its Application - A Nanotechnology Particle Project in Japan. Proceedings of the 2004 International Conference on MEMS, NANO and Smart Systems (ICMENS'04).

- 57 Sreeprasad, T.S. and Pradeep, T. (2013) Noble Metal Nanoparticles. 303-388.
- 58 Giasuddin, A.S.M., Jhuma, K.A., and Mujibul Haq, A.M. (2012) Use of gold nanoparticles in diagnostics, surgery and medicine: a review. Bangladesh J Med Biochem, 5 (2), 56-60.
- 59 Murthy, S.K. (2007) Nanoparticles in modern medicine: state of the art and future challenges. Int. I. Nanomed., 2 (2), 129–141.
- 60 Zou, L., Wang, H. et al. (2016) Current Approaches of Photothermal Therapy in Treating Cancer Metastasis with Nanotherapeutics. Theranostics., **6** (6), 762–772.
- 61 Mornet, S., Vasseur, S., Grasset, F. et al. (2004) Magnetic nanoparticle design for medical diagnosis and therapy. J. Mater. Chem., 14, 2161–75.
- 62 Huh, Y.M., Jun, Y.W., and Song, H.T. et al. (2005) In vivo magnetic resonance detection of cancer by using multifunctional magnetic nanocrystals. J. Am. Chem. Soc., 127, 12387-12391.
- 63 Harisinghani, M.G., Barentsz, J., and Hahn, P.F. et al. (2003) Noninvasive detection of clinically occult lymph-node metastases in prostate cancer. N. Engl. J. Med., 348, 2491-2499.
- **64** Duncan, R. (2003) The dawning era of polymer therapeutics. *Nat Rev Drug* Discov., 2, 347-360.
- 65 Panatarotto, D., Prtidos, C.D., Hoebeke, J., Brown, F., Kramer, E., Briand, J.P., Muller, S., Prato, M., and Bianco, A. (2003) Immunization with peptide-functionalized carbon nanotubes enhances virus-specific neutralizing antibody responses. Chemistry & Biology, 10, 961-966.
- 66 Muller, R.H., Mader, K., and Gohla, S. (2000) Solid lipid nanoparticles (SLN) for controlled drug delivery – a review of the state of the art. Eur. J. Pharm. Biopharm., **50**, 161–177.
- 67 Mah, C., Zolotukhin, I., Fraites, T.J., Dobson, J., Batich, C., Byrne, B. et al. (2000) Microsphere-mediated delivery of recombinant AAV vectors in vitro and in vivo. J. Mol Therapy., 1, S239.
- 68 Torchilin, V.P. and Weissig, V. (eds) (2003) Liposomes (Practical approach), Oxford Univ Press, Oxford.
- 69 Allen, T.M. and Cullis, P.R. (2004) Drug delivery systems; entering the main stream. Science, 303, 1818-1822.
- 70 Lai, C.Y., Trewyn, B.G., Jeftinija, D.M., Jeftinija, K., Xu, S., Jeftinija, S., and Lin, V.S.Y. (2003) A mesoporous silica nanosphere based carrier system with chemically removable CdS nanoparticle Caps for Stimuli responsive controlled release of neurotransmitters and drug molecules. J. Am. Chem. Soc., **125**, 4451–4459.
- 71 Hong, R., Han, G., Fernandez, J.M., Kim, B.J., Forbes, N.S., and Rotello, V.M. (2006) Glutathione mediated delivery and release using monolayer protected nanoparticle carriers. J. Am. Chem. Soc., 128, 1078-1079.
- 72 Giri, S., Trewyn, B.G., Stellmaker, M.P., and Lin, V.S.Y. (2005) Stimuli-responsive controlled-release delivery system based on mesoporous silica nanorods capped with magnetic nanoparticles. Angew. Chem. Int. Ed., **44**, 5038–5044.
- 73 Polizzi, M.A., Stasko, N.A., and Schoenfisch, M.H. (2007) Water soluble nitric oxide releasing gold nanoparticles. Langmuir., 23, 4938-4943.

- 74 Yang, Q., Wang, S.H., Fan, P.W., Wang, L.F., Di, Y., Lin, K.F., and Xiao, F.S. (2005) Ph-Resposive carrier system based on carboxylicacid modified mesoporous silica and polyelectrolyte for Drug delivery. Chem. Mater., 17, 5999-6003.
- 75 Neuman, D., Ostrowski, A.D., Absalonson, R.O., Strouse, G.F., and Ford, P.C. (2007) Photosensitized NO release from water soluble nanoparticle assemblies. J. Am. Chem. Soc., 129, 4146-4147.
- 76 Wang, S., Mamedova, N., Kotov, N.A., Chen, W., and Studer, J. (2002) Antigen/antibody immunocomplex from CdTe nanoparticlebioconjugates. Nano Lett., 2, 817-822.
- 77 Sapsford, K.E., Berti, L., and Medintz, I.L. (2006) Materials for fluorescence resonance enrgy transfer analysis: beyond traditionaldonor-acceptor combinations. Angew. Chem. Int. Ed., 45, 4562-4589.
- 78 Bruchez, M., Moronne, M., Gin, P., Weiss, S., and Alivisatos, A.P. (1998) Semiconductor nanocrystals as fluorescent biological labels. Science, 281, 2013-2016.
- 79 Dubertret, B., Calame, M., and Libchaber, A.J. (2001) Single-mismatch detection using gold quenched fluorescent oligonucleotides. Nat. Biotechnol., **19**, 365–370.
- 80 Keren, K., Berman, R.S., Buchstab, E., Sivan, U., and Braun, E. (2003) DNAtemplated carbon nanotube field-effect transistor. Science, 302, 1380-1382.
- 81 Nam, J.M., Thaxton, C.S., and Mirkin, C.A. (2003) Nanoparticles-based bio-bar codes for the ultrasensitive detection of proteins. Science., 301, 1884-1886.
- 82 Edelstein, R.L., Tamanaha, C.R., Sheehan, P.E., Miller, M.M., Baselt, D.R., Whitman, L.J., and Colton, R.J. (2000) The BARC biosensor applied to the detection of biological warfare agents. Biosensors Bioelectron, 14, 805-813.
- 83 Medarova, Z., Pham, W., Farrar, C., Petkova, V., and Moore, A. (2007) In vivo imaging of siRNA delivery and silencing tumors. Nat. Med., 13 (3), 372 - 377.
- 84 Mahtab, R., Rogers, J.P., and Murphy, C.J. (1995) Protein-sized quantum dot luminescence can distinguish between "straight", "bent", and "kinked" oligonucleotides. J. Am. Chem. Soc., 117, 9099-9100.
- 85 de la Isla, A., Brostow, W., Bujard, B., Estevez, M., Rodriguez, J.R., Vargas, S., and Castano, V.M. (2003) Nanohybrid scratch resistant coating for teeth and bone viscoelasticity manifested in tribology. Mat Resr Innovat, 7, 110–114.
- 86 Parak, W.J., Boudreau, R., Gros, M.L., Gerion, D., Zanchet, D., Micheel, C.M., Williams, S.C., Alivisatos, A.P., and Larabell, C.A. (2002) Cell motility and metastatic potential studies based on quantum dot imaging of phagokinetic tracks. Adv. Mater., 14, 882-885.
- 87 Shinkai, M., Yanase, M., Suzuki, M., Honda, H., Wakabayashi, T., Yoshida, J., and Kobayashi, T. (1999) Intracellular hyperthermia for cancer using magnetite cationic liposomes. J. Magn. Magn. Mater., 194, 176-184.
- 88 Salata, O.V. (2004) Applications of nanoparticles in biology and medicine. *Journal of Nanobiotechnology.*, **2**, 1–6.
- 89 Jain, P.K., Lee, K.S., El-Sayed, I.H., and El-Sayed, M.A. (2006) Calculated absorption and scattering properties of gold nanoparticles of different size,

- shape, and composition: Applications in biological imaging and biomedicine. J. Phys. Chem. B, 110 (14), 7238-7248.
- 90 Johnstone, T.C., Kulak, N., Pridgen, E.M., Farokhzad, O.C., Langer, R., and Lippard, S.J. (2013) Nanoparticle Encapsulation of Mitaplatin and the Effect Thereof on In Vivo Properties. ACS Nano, 7 (7), 5675-5683.
- 91 Ibrahim, N.K., Desai, N., Legha, S. et al. (2002) Phase I and pharmacokinetic study of ABI-007, a cremophor-free, protein-stabilized, nanoparticle formulation of paclitaxel. Clin Cancer Res. 8, 1038-44.
- 92 Nagarajan, R. (2008) Nanoparticles: Building Blocks for Nanotechnology, ACS Symposium Series, American Chemical Society, Washington, DC.
- 93 Pantic, I. (2014) Application of silver nanoparticles in experimental physiology and clinical medicine: current status and future prospects. Rev. Adv. Mater. Sci., 37, 15-19.
- 94 Jain, S., Hirst, D.G., and O'Sullivan, J.M. (2012) Gold nanoparticles as novel agents for cancer therapy. Br J Radiol., 85 (1010), 101–113.
- 95 Huang, X. and El-Sayed, M.A. (2010) Gold nanoparticles: optical properties and implementations in cancer diagnosis and photothermal therapy. J. Adv. Res., 1, 13-28.
- 96 Spivak, M.Y., Bubnov, R.V., Yemets, I.M., Lazrenko, L.M., Tymoshok, N.O., and Ulberg, Z.R. (2013) Gold nanoparticles - the theranostic challenge for PPPM:nanocardiology application. *The EPMA Journal*, 4, 1,1–17.
- 97 Abou El-Nour, K.M.M., Eftaiha, A., Al-Warthan, A., and Ammar, R.A.A. (2010) Synthesis and applications of silver nanoparticles. Arabian J. Chem., 3 (3), 2010.
- 98 Tran, Q.H., Nguyen, V.Q., and Le, A.T. (2013) Silver nanoparticles: synthesis, properties, toxicology, applications and perspectives. Adv. Nat. Sci.: Nanosci. Nanotechnol., 4 (3), 2013.
- 99 Mulfinger, L. (2007) Synthesis and study of silver nanoparticles. J. Chem. Educ., 84 (2), 322.
- 100 Gawande, M.B., Goswami, A., Felpin, F.X., Asefa, T., Huang, X., Silva, R., Zou, X., Zboril, R., and Varma, R.S. (2016) Cu and Cu-based nanoparticles: synthesis and applications in catalysis. Chem. Rev., 116 (6), 3722-3811.
- 101 Yang, Y., Sun, C., Ren, Y., Hao, S., and Jiang, D. (2014) New route toward building active ruthenium nanoparticles on ordered mesoporous carbons with extremely high stability. Scientific reports, 4.
- 102 Maggioni, D., Fenili, F., Alfonso, L.D., Donghi, D., Panigati, M., Zanoni, I., Marzi, R., Manfredi, A., Ferruti, P., Alfonso, G.D., and Ranucci, E. (2012) Luminescent rhenium and ruthenium complexes of an amphoteric poly(amidoamine) functionalized with 1,10-phenanthroline. Inorg. Chem., **2012** (51), 12776–12788.
- 103 Cookson, B.J. (2012) The preparation of palladium nanoparticles. *Platinum* Metals Rev., **56** (2), 83–98.
- 104 Chen, H., Wei, G., Ispas, A., Hickey, S.G., and Eychumuller, A. (2010) Synthesis of palladium nanoparticles and their applications for surface enhanced raman scattering and electrocatalysis. J. Physical Chemistry C., 114 (50), 21976-21981.

- 105 Raj, A.T., Ramanujan, K., Thangavel, S., Gopalakrishan, S., Raghavan, N., and Venugopal, G. (2015) Facile synthesis of vanadium-pentoxide nanoparticles and study on their electrochemical, photocatalytic properties. J. Nanosci. Nanotechnol., 15 (5), 3802-3808.
- 106 Pinto, S., D'Ornelas, L., and Betancourt, P. (2008) Synthesis and characterization of vanadium nanoparticles on activated carbon and their catalytic activity in thiophene hydrodesulphurization. Appl. Surf. Sci., 254 (17), 5390-5393.
- 107 Natalio, F., Andre, R., Hartog, A.F., Stoll, B., Jochum, K.P., Wever, R., and Tremel, W. (2013) Vanadium pentoxide nanoparticles mimic vanadium haloperoxidases and thwart biofilm formation. Nat. Nanotechnol., 7, 530-535.