

Editorial

A Short Overview on the Nanoparticle-Based Smart Drug Delivery Systems

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Recently, nanotechnology has been extensively used in different areas such as catalysts, biosensors, food industry, pharmaceuticals as well as gene delivery, etc. Nanoparticles are particles with size ranged between 1 and 100 nanometers and may exhibit size related properties that differ significantly from those observed in their own bulk materials^[1-3].

Pharmaceutical nanoparticles, can beneficially enhance therapeutic efficacy by improving the drug bioavailability^[4,5]. Controlled release formulations based on nanoparticles provide better penetration and allow slow and controlled release of active pharmaceutical ingredients at the target site^[6-9].

There are many types of drug delivery systems, which have been developed for targeted and controlled release purposes. Smart or intelligent drug delivery systems are the main group of these systems which are capable to adjust drug release rates in response to a trigger^[10,11]. These systems are divided to two groups: open-loop control systems and closed-loop control systems^[12]. In open-loop control systems external triggers e.g. magnetic, thermal, ultrasonic, and electric are essential for drug release while in a closed-loop type, the drug release rate is controlled without any external intervention^[13]. The drug release from the latter group is controlled by an internal trigger such as: pH or temperature - responsive drug delivery systems, urea-responsive drug delivery systems, glucose-responsive insulin delivery systems, inflammation-induced pulsatile release systems, morphine triggered naltrexone delivery systems, the systems utilizing molecular targeting agents such as antibody or aptamer and the systems utilizing chelation^[13,14].

Variety of molecular targeting agents have been documented such as enzymes, humanized antibodies and single-chain Fv generated from murine hybridoma or phage display, aptamers, mini bodies and peptides^[15-17]. The drug delivery systems that utilize these agents are developed to various applications such as isolation, detection and release of therapeutic agent^[18]. From the pharmaceutical point of view, functionalization of the nanoparticles with molecular targeting agents has shown promising application in the biomedical areas^[19,20].

Biodegradable and biocompatible smart polymers are able to release the entrapped drugs at the appropriate time and site of action in response to specific physiological triggers. The responses may alter from swelling/contraction to disintegration^[15,21,22]. Aliphatic polyesters, such as polylactic acid (PLA) and polyglycolic acid (PGA), as well as their copolymer poly(lactic-co-glycolic acid) (PLGA) are the mostly used polymers in developing controlled release nanoparticles. Chitosan and its derivatives are very beneficial materials for drug delivery purposes. Chitosan nanoparticles targeted using magnetic nanoparticles, antibodies or aptamers have been assessed for crossing from the blood-brain barrier^[23].

Mesoporous silica nanoparticles can apply as stimuli-responsive materials to create smart delivery systems. Targeting agents such as nucleic acids can be employed to cap the pores of these nanoparticles as biomolecular cap and therefore these nanoparticles can act as gate keepers. Either intracellular or external triggers, such as changes in pH, light, enzymatic activity, electromagnetic field or ultrasound are applied to remove the cap from nanoparticle, which result in the release of drug. Furthermore, optical and magnetic contrast agents can also be introduced into nanoparticles to form multipurpose drug delivery systems^[24,25].

Magnetic nanoparticles can be applied for physically directing of the drugs into their desired and accessible target cells or tissues by use of an external magnetic field^[23]. Magnetic nanoparticles have attracted a special attention in nanomedicine due to advantages such as ideal surface modification, competency to move in cellular level by an external magnetic field and ability to hyperthermia effect^[20,26]. Functionalization of these nanoparticles with molecular targeting agents improves their biomedical application especially in cancer therapy intentions.

The hollow structures of carbon nanotubes can be loaded with the drug molecules for extended release of the drugs^[27,28]. Functionalization of single walled nanotubes by targeting agents has been used to selectively target of drugs by pH-dependent release profile^[29].

Despite of many challenges, there is a great attention for the development of smart drug delivery systems. Smart drug

delivery systems based on polymeric nanoparticles may have extensive applications in oral drug delivery of biological drugs, which are sensitive to both gastric acid and enteric enzymes. Smart drug delivery systems can also be applied in the field of smart diagnostics^[30].

References

- Adibkia, K., Bozorgmehr, Z., Dastmalchi, S., et al. Evaluating retardation and physicochemical properties of co-ground mixture of Na-diclofenac with Magnesium stearate. (2012) *Res Pharm Sci* 7(5): S657.
- Kumar, D., Jain, N., Gulati, N., et al. Nanoparticles laden *in situ* gelling system for ocular drug targeting. (2013) *J Adv Pharm Technol Res* 4(1): 9-17.
- Dizaj, S.M., Jafari, S., Khosroushahi, A.Y. A sight on the current nanoparticle-based gene delivery vectors. (2014) *Nanoscale Res Lett* 9(1): 252.
- Adibkia, K., Omid, Y., Siah, M.R., et al. Inhibition of endotoxin-induced uveitis by methylprednisolone acetate nanosuspension in rabbits. (2007) *J Ocul Pharmacol Ther* 23(5): 421-32.
- Mahanty, S., Sruti, J., Patra, C.N., et al. Particle design of drugs by spherical crystallization techniques. (2010) *Int J Pharm Sci Nanotech* 3(2): 912-918.
- Aouada, F.A., De Moura, M.R., Orts, W.J., et al. Polyacrylamide and methylcellulose hydrogel as delivery vehicle for the controlled release of paraquat pesticide. (2010) *J Mater Sci* 45(18): 4977-4985.
- Adibkia, K., Ghanbarzadeh, S., Shokri, M.H., et al. Micro-porous surfaces in controlled drug delivery systems: design and evaluation of diltiazem hydrochloride controlled porosity osmotic pump using non-ionic surfactants as pore-former. (2013) *Pharm Develop Technol* 19(4): 507-12.
- Adibkia, K., Alaei-Beirami, M., Barzegar-Jalali, M., et al. Evaluation and optimization of factors affecting novel diclofenac sodium-eudragit RS100 nanoparticles. (2012) *Afr J Pharm Pharmacol* 6(12): 941-947.
- Adibkia, K., Hamedeyazdan, S., Javadzadeh, Y. Drug release kinetics and physicochemical characteristics of floating drug delivery systems. (2011) *Expert Opin Drug Deliv* 8(7): 891-903.
- Kasagana, V.N., Karumuri, S.S.M. T. Recent Advances in Smart Drug Delivery Systems. (2011) *Int J Nov Drug Deliv Tech* 1(3): 201-207.
- Crommelin, D.J., Florence, A.T. Towards more effective advanced drug delivery systems. (2013) *Int J Pharm* 454(1): 496-511.
- Doyle, F.J. 3rd, Huyett, L.M., Lee, J.B., et al. Closed-loop artificial pancreas systems: engineering the algorithms. (2014) *Diabetes Care* 37(5): 1191-1197.
- Wise, D.L. Handbook of pharmaceutical controlled release technology. (2000) CRC Press. 902.
- Alvarez-Lorenzo, C., Concheiro, A. Intelligent drug delivery systems: polymeric micelles and hydrogels. (2008) *Mini Rev Med Chem* 8(11): 1065-1074.
- Fahmy, T.M., Fong, P.M., Goyal, A., et al. Targeted for drug delivery. (2005) *Materials Today* 8(8): 18-26.
- Sun, H., Zhu, X., Lu, P.Y., et al. Oligonucleotide Aptamers: New Tools for Targeted Cancer Therapy. (2014) *Mol Ther Nucleic Acids* 3: e182.
- Trujillo, C.A., Nery, A.A., Alves, J.M., et al. Development of the anti-VEGF aptamer to a therapeutic agent for clinical ophthalmology. (2007) *Clin Ophthalmol* 1(4): 393-402.
- Yoshida, R., Sakai, K., Okano, T., et al. Pulsatile drug delivery systems using hydrogels. (1993) *Adv Drug Deliv Revs* 11(1): 85-108.
- Okazawa, A., Maeda, H., Fukusaki, E., et al. In vitro selection of hematoporphyrin binding DNA aptamers. (2000) *Bioorg Med Chem Lett* 10(23): 2653-2656.
- Zhou, W., Huang, P.-J., Ding, J., et al. Aptamer-based biosensors for biomedical diagnostics. (2014) *Analyst* 139(11): 2627-2640.
- Huang, G., Gao, J., Hu, Z., et al. Controlled drug release from hydrogel nanoparticle networks. (2004) *J Control Release* 94(2-3): 303-311.
- Zakeri-Milani, P., Loveymi, B.D., Jelvehgari, M., et al. The characteristics and improved intestinal permeability of vancomycin PLGA-nanoparticles as colloidal drug delivery system. (2013) *Colloids Surf B: Biointer* 103: 174-1781.
- Bruno, J.G. A review of therapeutic aptamer conjugates with emphasis on new approaches. (2013) *Pharmaceuticals* 6(3): 340-357.
- Park, Y.-H., Bae, H., Jang, Y., et al. Effect of the size and surface charge of silica nanoparticles on cutaneous toxicity. (2013) *Mol Cell Toxicol* 9(1): 67-74.
- Santra, S. Fluorescent silica nanoparticles for cancer imaging. (2010) *Methods Mol Biol* 624: 151-162.
- Gijs, M.A. Magnetic bead handling on-chip: new opportunities for analytical applications. (2004) *Microfluidics and Nanofluidics* 1(1): 22-40.
- So, H.-M., Park, D.-W., Chang, H., et al. Carbon Nanotube Biosensors with Aptamers as Molecular Recognition Elements. (2010) *Methods Mol Biol* 625: 239-49.
- Nam, K., Eom, K., Yang, J., et al. Aptamer-functionalized nanopattern based on carbon nanotube for sensitive, selective protein detection. (2012) *J Mater Chem* 22(44): 23348-23356.
- Shao, W., Arghya, P., Yiyong, M., et al. Carbon Nanotubes for Use in Medicine: Potentials and Limitations. (2013).
- James, H.P., John, R., Alex, A., et al. Smart polymers for the controlled delivery of drugs—a concise overview. (2014) *Acta Pharm Sin B* 4(2): 120-127.