

Available online on 15.03.2019 at <http://jddtonline.info>

Journal of Drug Delivery and Therapeutics

Open Access to Pharmaceutical and Medical Research

© 2011-18, publisher and licensee JDDT, This is an Open Access article which permits unrestricted non-commercial use, provided the original work is properly cited

Open  Access

Mini Review

Nanoparticle therapeutics: a new treatment of cancer

Di Kang

China Pharmaceutical University Nanjing, Jiangsu, 211198, China

Article Info: Received 31 Jan 2019; Review Completed 07 March 2019; Accepted 09 March 2019; Available online 15 March 2019**Cite this article as:**Kang D, Nanoparticle therapeutics: a new treatment of cancer, Journal of Drug Delivery and Therapeutics. 2019; 9(2):394-396 <http://dx.doi.org/10.22270/jddt.v9i2.2415>***Address for Correspondence:**

Di Kang, China Pharmaceutical University Nanjing, Jiangsu, 211198, China

INTRODUCTION

To get speedy recovery from any deadly diseases, the most important thing is getting proper medication¹⁻⁹. In the passing years, it has been seen that cells and specific genes in the human body are acquiring drug resistance capability against certain drugs which needs to work correctly for getting rid of any particular diseases. Therefore, the introduction of nanoparticle and exosomes are gaining importance in drug manufacturing units. It is not only helping the drug get administered into the exact position where it needs to get delivered but also hastening speedy recovery of the patient from a lethal disease¹⁰⁻¹⁹.

Numerous methods are used to get rid of the problem like drug resistance ability, impassability of drugs through intercellular pathways and gene silencing activities.

It has been seen that few specific cancers like glioblastoma are more robust to get diagnosed in the initial stages. Hence, during the final stages, it becomes impossible for chemotherapeutic drugs to get administered into the exact location of the disease.

Nanoparticles are used in this regard for early detection and treatment of such specific brain tumor cases²⁰⁻³¹. Apart from that, exosomes, as extracellular organelles can be used to get

proteomic and genetic information from long distance intercellular communication. Along with it, exosomes are used for proper drug delivery on the exact location where the disease is mainly predominant.

Similarly, psychosis is another fatal disease that is impacting thousands of people worldwide. Nanoparticles are used for the treatment of this disease for delivering the drug. Nanoparticles of 1-500 nanometers are used primarily to administer antipsychotic drugs into the human body so that the administered drugs can get easily absorbed through the blood-brain barrier³²⁻⁴⁴.

To control the impact of multidrug resistance among the human sometimes excessive amount of drugs get administered, but it only causes intolerance of toxicity and death of the patient. Along with it, histone modification and DNA methylation have been introduced to minimize the issues in this regards. Synthesis of oligonucleotides after some chemical modification and synthesis of some nano-sized camptothecin for combined drug delivery are also few steps that are getting followed. Doxorubicin (DOX) is the best medication used for curing the malignant tumor. Apart from that, lung and cervical carcinomas T7 peptide conjugated nanoparticles are mainly used.

Table 1: Nanoparticles with Dual Drug Delivery to Control MDR

Materials	Agent 1	Agent 2	Cancer cell lines	Ref
Poly(ϵ -caprolactone) and poly(ethyl ethylene phosphate)	Doxorubicin	NA	MCF-7/ADR	(38)
Poly(lactic-co-glycolic acid) and polyethylene glycol	Vincristine	NA	MCF-7/ADR	(42)
Pluronic-P105	Doxorubicin	Paclitaxel	MCF-7/ADR	(46)
Poly(ethylene oxide)-poly(propylene oxide)-poly(ϵ -caprolactone)	Docetaxel	Chloroquine	MCF-7 and MCF-7/ADR	(47)
VE and tocopherol poly(ethylene glycol)succinate	Paclitaxel	5-Fluorouracil	KB-8-5 and KB-3-1	(48)
Poly (lactic-co-glycolic acid)	Docetaxel	TPGS	HeLa	(50)
EPC, DOTAP, cholesterol and PEG2KPE	Paclitaxel	Tariquidar	SKOV-3 and SKOV-3TR	(51)
CEA and AHM	Doxorubicin	Verapamil	NCI/ADR-RES	(53)
Chitosan	Doxorubicin	Pyroglutathione	HepG-2	(54)
TPGS2000 and PEG2000-DSPE	Doxorubicin	Curcumin	MCF7/ADR	(55)
Poly(D,L-lactide-co-glycolide)	Doxorubicin	Curcumin	K562	(56)
1-Palmitoyl-2-azelaoyl-sn-glycero-3-phosphocholine	Doxorubicin	Ceramide	P388/ADR	(57)
Precirol ATO 5,Squalene, SPC, Tween-80 and DOTAP	Doxorubicin/Paclitaxel	siRNA targeting MRP1 and BCL2	A549	(62)
PAMAM and PEG-2K-DOPE	Doxorubicin	siRNA targeting GFP	A549 cells and C166-GFP	(63)

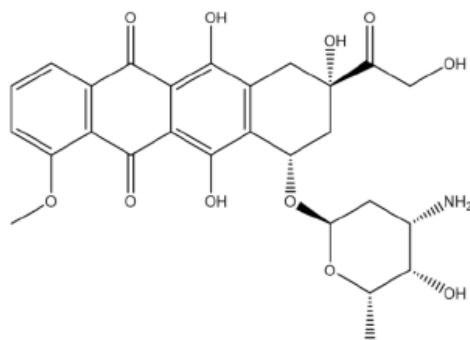


Figure 2: Chemical Structure of DOX

Different strategies and process have been invented for careful drug delivery into the affected areas. To control the multi-drug resistance (MDR), the process to introduce multiple chemotherapeutic drugs is used nowadays. Apart from this, nanoparticles used for antipsychotic disease treatment is comparatively low while it is mainly used for cancer treatment⁴⁵⁻⁵¹. This is only for the lack of knowledge about psychosis that restricts the drugs from being designed. Similarly, safe designing of the EMs cell for gene therapy is used for the proper treatment of cancer cells of the human body. On the other hand, DNA methylation represses the growth and spread of cancer cell for the long term. Non-small-cell lung cancer (NSCLC) is a lethal form of lung cancer that accounts for 80% of total lung cancer cases around the world. NSCLC is less responsive for chemotherapy, radiotherapy and any treatment.

References

1. Kang, C., Qin, J., Osei, W. & Hu, K. Age-dependent Mitochondrial Targeting Of Protein Kinase C Epsilon In Cardioprotection. *The FASEB Journal* (2017).
2. Han, R., Sun, Y., Kang, C., Sun, H. & Wei, W. Amphiphilic dendritic nanomicelle-mediated co-delivery of 5-fluorouracil

and doxorubicin for enhanced therapeutic efficacy. *Journal of Drug Targeting* 2017; **25**:140-148.

3. Yan, G., *et al.* Application of Real-Time Cell Electronic Analysis System in Modern Pharmaceutical Evaluation and Analysis. *Molecules* **23**, 3280 (2018).
4. Duan, Y., *et al.* Bioactivity evaluation-based ultra high-performance liquid chromatography coupled with electrospray ionization tandem quadrupole-time-of-flight mass spectrometry and novel distinction of multi-subchemome compatibility recognition strategy with Astragali Radix-Fructus Corni herb-pair as a case study. *J Pharm Biomed Anal* 2016; **129**:514-534.
5. Sun, Y., *et al.* Co-delivery of dual-drugs with nanoparticle to overcome multidrug resistance. *European Journal of BioMedical Research* 2016; **2**:12-18.
6. Ai, R., *et al.* Comprehensive epigenetic landscape of rheumatoid arthritis fibroblast-like synoviocytes. *Nature communications* 2018; **9**:1921.
7. Ai, R., *et al.* Comprehensive epigenetic landscape of rheumatoid arthritis fibroblast-like synoviocytes. *Nat Commun* 2018; **9**:1921.
8. Fan, S., *et al.* Computationally expanding infinium HumanMethylation450 BeadChip array data to reveal distinct DNA methylation patterns of rheumatoid arthritis. *Bioinformatics* 2016; **32**:1773-1778.
9. Zhu, Y., *et al.* Constructing 3D interaction maps from 1D epigenomes. *Nature communications* 2016; **7**:10812.
10. Liu, F., Sun, Y. & Kang, C. Controlling Amphiphilic Functional Block Copolymers' Self-Assembly: From Structure to Size. (2016).
11. Song, L., *et al.* Crocetin inhibits lipopolysaccharide-induced inflammatory response in human umbilical vein endothelial cells. *Cellular Physiology and Biochemistry* 2016; **40**:443-452.
12. Sun, Y., Kang, C., Liu, F. & Song, L. Delivery of antipsychotics with nanoparticles. *Drug Development Research* 2016; **77**:393-399.
13. Kang, C., *et al.* Delivery of nanoparticles for treatment of brain tumor. *Current Drug Metabolism* 2016; **17**:745-754.
14. Xue, X., *et al.* Discovery of novel inhibitors disrupting HIF-1 α /von Hippel-Lindau interaction through shape-based screening and cascade docking. *PeerJ* 2016; **4**:e2757.
15. Hersch, S.J., *et al.* Divergent protein motifs direct elongation factor P-mediated translational regulation in *Salmonella enterica* and *Escherichia coli*. *MBio* 2013; **4**:e00180-00113.

16. Hersch, S.J., *et al.* Divergent protein motifs direct elongation factor P-mediated translational regulation in *Salmonella enterica* and *Escherichia coli*. *MBio* 2013; **4**:e00180-00113.
17. Shuhong, X., *et al.* Dynamic expression of AQP4 in early stage of ischemia/reperfusion rats and cerebral edema. *Chinese Pharmacological Bulletin* 2016; **32**:1433-1441.
18. Peng, J., *et al.* Enhanced Liver Regeneration After Partial Hepatectomy in Sterol Regulatory Element-Binding Protein (SREBP)-1c-Null Mice is Associated with Increased Hepatocellular Cholesterol Availability. *Cellular Physiology and Biochemistry* 2018; **47**:784-799.
19. Ngo, V., *et al.* Epigenomic analysis reveals DNA motifs regulating histone modifications in human and mouse. *Proceedings of the National Academy of Sciences*, 201813565 (2019).
20. Ngo, V., Wang, M. & Wang, W. Finding de novo methylated DNA motifs. *bioRxiv*, 043810 (2018).
21. Yang, Z., *et al.* Functional exosome-mimic for delivery of siRNA to cancer: in vitro and in vivo evaluation. *Journal of Controlled Release* 2016; **243**:160-171.
22. Kang, C., Hernandez, V.A. & Hu, K. Functional interaction of the two-pore domain potassium channel TASK-1 and caveolin-3. *Biochimica et Biophysica Acta (BBA)-Molecular Cell Research* 2017; **1864**:1537-1544.
23. Waller, A.P., *et al.* GLUT12 functions as a basal and insulin-independent glucose transporter in the heart. *Biochimica et Biophysica Acta (BBA)-Molecular Basis of Disease* 2013; **1832**:121-127.
24. Li, Q., *et al.* Identification by shape-based virtual screening and evaluation of new tyrosinase inhibitors. *PeerJ* 2018; **6**:e4206.
25. Chen, Y., *et al.* Identification of 4-aminoquinoline core for the design of new cholinesterase inhibitors. *PeerJ* 2016; **4**:e2140.
26. Kang, C. & Hu, K. Impact of hypoxia in the expression and regulation of the TASK-1 potassium channel in cardiac myocytes. *The FASEB Journal* 2016; **30**:lb598-lb598.
27. Fan, S., *et al.* Integrative analysis with expanded DNA methylation data reveals common key regulators and pathways in cancers. *NPJ genomic medicine* 2019; **4**:2.
28. Kang, C. *Ion channels, protein kinase C and caveolae in cardioprotection*, (The Ohio State University, 2015).
29. HE, L.-y., *et al.* Isolation and Characterization of Cadmium-Resistant Endophytic and Rhizobacteria From *Solanum nigrum* in Orefield [J]. *Journal of Ecology and Rural Environment* 2011; **6**.
30. Yung, B.C., *et al.* Lipid nanoparticles composed of quaternary amine-tertiary amine cationic lipid combination (QTsome) for therapeutic delivery of AntimiR-21 for lung cancer. *Molecular pharmaceutics* 2016; **13**:653-662.
31. Cheng, X., *et al.* Lipid Nanoparticles Loaded with an Antisense Oligonucleotide Gapmer Against Bcl-2 for Treatment of Lung Cancer. *Pharmaceutical research* 2017; **34**:310-320.
32. Fan, S. & Chi, W. Methods for genome-wide DNA methylation analysis in human cancer. *Brief Funct Genomics* 2016; **15**:432-442.
33. Kang, C. & Hu, K. Modulation of the two-pore domain potassium channel TASK-1 by caveolin-3. *The FASEB Journal* 2015; **29**:845.814.
34. Davis, M.E., Chen, Z.G. & Shin, D.M. Nanoparticle therapeutics: an emerging treatment modality for cancer. *Nat Rev Drug Discov* 2008; **7**:771-782.
35. Kang, C., Sun, Y., Wang, M. & Cheng, X. Nanosized camptothecin conjugates for single and combined drug delivery. *European Journal of BioMedical Research* 2016; **2**:8-14.
36. Qiao, H., *et al.* Orally delivered polycurcumin responsive to bacterial reduction for targeted therapy of inflammatory bowel disease. *Drug Delivery* 2017; **24**:233-242.
37. Liu, F., Sun, Y., Kang, C. & Zhu, H. Pegylated Drug Delivery Systems: From Design to Biomedical Applications. *Nano LIFE* 2016; **6**:1642002.
38. Sun, Y., Kang, C., Yao, Z., Liu, F. & Zhou, Y. Peptide-Based Ligand for Active Delivery of Liposomal Doxorubicin. *Nano Life* 2016; **6**:1642004.
39. Yeh, C.Y., Hsiao, J.K., Wang, Y.P., Lan, C.H. & Wu, H.C. Peptide-conjugated nanoparticles for targeted imaging and therapy of prostate cancer. *Biomaterials* 2016; **99**:1-15.
40. Fan, S., Huang, K., Ai, R., Wang, M. & Wang, W. Predicting CpG methylation levels by integrating Infinium HumanMethylation450 BeadChip array data. *Genomics* 2016; **107**:132-137.
41. Qiao, H., *et al.* Redox-triggered mitoxantrone prodrug micelles for overcoming multidrug-resistant breast cancer. *Journal of drug targeting* 2018; **26**:75-85.
42. Kang, C., Qin, J., Osei, W. & Hu, K. Regulation of protein kinase C-epsilon and its age-dependence. *Biochemical and Biophysical Research Communications* 2017; **482**:1201-1206.
43. Sun, Y., *et al.* RGD Peptide-Based Target Drug Delivery of Doxorubicin Nanomedicine. *Drug development research* 2017; **78**:283-291.
44. Kang, C. & Hu, K. Role of caveolin-3 in adenosine-induced increase in mitochondrial PKCε. *The FASEB Journal* 2013; **27**:1191.1197-1191.1197.
45. Cheng, X. & Lee, R.J. The role of helper lipids in lipid nanoparticles (LNPs) designed for oligonucleotide delivery. *Adv Drug Deliv Rev* 2016; **99**:129-137.
46. Sun, Y. & Kang, C. Self-Assembly of Peptides into Hydrogel. *Journal of Organic & Inorganic Chemistry* 2016; **2**:5.
47. Yao, Z., Sun, Y. & Kang, C. Structure and self-assembly of multicolored Naphthalene Diimides Semiconductor. *Nano LIFE* 2016; **6**:1642007.
48. Gorkin, D.U., Williams, B.A., Trout, D. & Amrhein, H. Systematic mapping of chromatin state landscapes during mouse development. (2017).
49. Zhang, K., Wang, M., Zhao, Y. & Wang, W. Systems-level identification of transcription factors critical for mouse embryonic development. *bioRxiv*, 167197 (2017).
50. Cheng, X., *et al.* T7 Peptide-Conjugated Lipid Nanoparticles for Dual Modulation of Bcl-2 and Akt-1 in Lung and Cervical Carcinomas. *Molecular pharmaceutics* 2018; **15**:4722-4732.
51. Zhong, X., Sun, Y., Kang, C. & Wan, G. The theory of dielectrophoresis and its applications on medical and materials research. *European Journal of BioMedical Research* 2017; **2**:7-11.