

Docetaxel-loaded solid lipid nanoparticles as a basis for a targeted and dose-sparing personalized breast cancer treatment strategy

Natalia V Danilova^{1,2}

Zhomart R Kalzhanov³

Nina A Nefedova²

Pavel G Mal'kov²

Ioannis P Kosmas^{1,4}

Marina Y Eliseeva^{1,5}

Ospan A Mynbaev^{1,5,6}

¹International Translational Medicine and Biomodeling Research Team, MIPT Center for Human Physiology, Laboratory of Cellular and Molecular Technologies, Moscow Institute of Physics and Technology, State University,

²Department of Physiology and Basic Pathology, Faculty of Fundamental Medicine, Lomonosov Moscow State University, Moscow, Russia;

³Department of Human Metabolism, Academic Unit of Reproductive and Developmental Medicine, Sheffield University, Sheffield, UK; ⁴Department of Obstetrics and Gynecology, Ioannina State General Hospital G Chatzikosta, Ioannina, Greece;

⁵Department of Obstetrics, Gynecology and Reproductive Medicine, Peoples' Friendship University of Russia, ⁶Laboratory of Immunology, Moscow State University of Medicine and Dentistry named after AI Evdokimov, Moscow, Russia

Correspondence: Ospan A Mynbaev
Moscow Institute of Physics and Technology, State University,
Dolgoprudny, 9 Institutskii Lane,
Dolgoprudny, Moscow 141700, Russia
Email ospanmynbaev@hotmail.com

Dear editor

The long-term survival rate of patients with breast cancer was improved by the application of systemic adjuvant chemotherapy,¹ although the primary breast cancer treatment strategy consists of mastectomy with lymphadenectomy and radiotherapy followed by breast reconstruction.²⁻⁵ Unfortunately, most adjuvant chemotherapeutic agents trigger major side effects.^{1,6} Therefore, we have read with great interest an article in the *International Journal of Nanomedicine* on the design of docetaxel-loaded solid lipid nanoparticles (DSNs) aimed at reducing the systemic toxicity of standardized docetaxel treatment.⁷

Our congratulation to the authors⁷ for their clear demonstration of the reduced cytotoxicity of DSNs and significantly decreased myelosuppressive toxicity by recovering the proliferation and differentiation of bone marrow progenitor cells, while triggering more apoptosis in MCF-7 cells at a low dose compared with the commercial formula of docetaxel by an arrested cell cycle progression in the G2/M stage.

The acute necessity for such state-of-the-art studies is linked to a high worldwide incidence of breast cancer; in the World Health Organization Fact sheet, its increased metastatic potency is listed as one of the most common causes of cancer death.⁸

The incidence of breast cancer is high in Western European countries, ie, about 89.7 per 100,000 women.⁸ The same high incidence applies to other developed countries. For our part, we have analyzed the statistical data concerning breast cancer in the Russian Federation.⁹

According to the statistical report by the Federal Research Institute for Health Organization and Informatics of the Ministry of Health of the Russian Federation between 2003 and 2012 (Table 1), the 10-year breast cancer incidence rates average 68.99 per 100,000 women (an average of 52,647 women annually), of which 95.1% cases were histologically verified. The average 5-year survival rate was 56.2% among women with breast cancer followed up in state and municipal cancer institutions. The average annual mortality of patients with verified breast cancer was 10.0% among individuals who were followed up in cancer institutes. Subsequently, between 2003 and 2012, about 25,697 women per year died of breast cancer in the Russian Federation.

Our investigation, as proved by time series analysis (JMP7 software), revealed that the incidence of breast cancer had been rising continually, with increasing trends from 2003 to 2012 while at the same time the number of cancer deaths has been steadily decreasing (Table 1). Overall, the same trends exist for all cancer patients, in the same time.

Table 1 Modified data from a statistical report for 2012 by the Federal Research Institute for Health Organization and Informatics of the Ministry of Health of the Russian Federation

Statistical parameters	Years											10-year average m ± 95% confidence interval
	2003	2004	2005	2006	2007	2008	2009	2010	2011	2012		
1. Incidence rates for breast cancer per 100,000 women	60.3	64.4	65.1	65.7	67.9	68.8	71.2	74.8	74.9	76.8	69.0	
2. Histologically verified diagnosis of breast cancer (%)	93.7	94.0	94.4	94.4	95.4	95.2	95.6	95.7	96.6	96.3	95.1	
3. Five-year survival of patients with breast cancer followed up in cancer institutions (%)	54.1	55.1	55.8	55.9	56.3	56.0	56.7	57.0	57.6	57.9	56.2	
4. Annual mortality rate in women with verified breast cancer and followed up in cancer institutes (from all deaths due to cancer, %)	12.1	11.5	10.9	10.3	10.1	9.7	9.5	9.1	8.7	8.3	10.0	
5. Patients having died of any cancer among individuals who were followed up in cancer institutes (n)	264,972	263,088	259,456	257,655	260,926	258,006	257,114	255,811	249,398	243,308	256,973.4	
6. Women having died of breast cancer among individuals who were followed up in cancer institutes (n)	32,061	30,255	28,281	26,538	26,353	25,026	24,425	23,279	21,697	20,194	25,697.0	

Note: Adapted from: Federal Research Institute for Health Organization and Informatics of the Ministry of Health of the Russian Federation. Annual Statistics, 2012. Socially important diseases in the Russian population. Moscow, 2013. Available from: <http://www.mednet.ru/en.html>. Accessed November 14, 2014.⁹

This phenomenon can be explained first by improvements in the Russian health care system. Government programs were set up applying new diagnostic technologies for early breast cancer screening, and preventive medical strategies were encouraged. Second, the rise in incidence of breast cancer might also be associated with gradual lifestyle changes. Many mothers in Russia decline breastfeeding and, in addition, worldwide environmental changes are reflected in an increased incidence of breast cancer, including in countries with a low prevalence of the disease.

This brief analysis demonstrates that our modern community calls for new therapeutic approaches in the treatment of breast cancer. We believe that further studies could show the application of DSNs to be a basic compound for a targeted and dose-sparing personalized breast cancer treatment strategy.

Acknowledgment

This work was supported by the Russian Science Foundation (grant 14-31-00024). All authors are members of the International Translational Medicine and Biomodeling Research Team (<http://mathbiomed.crec.mipt.ru>).

Author contributions

All authors contributed to the discussion regarding the original study by Yuan et al,⁷ and revising the final manuscript, and agree to be accountable for all aspects of the work.

Disclosure

The authors report no conflicts of interest in this work.

References

1. Rampurwala MM, Rocque GB, Burkard ME. Update on adjuvant chemotherapy for early breast cancer. *Breast Cancer (Auckl)*. 2014; 8:125–133.
2. Kurian AW, Lichtensztajn DY, Keegan TH, Nelson DO, Clarke CA, Gomez SL. Use of and mortality after bilateral mastectomy compared with other surgical treatments for breast cancer in California, 1998–2011. *JAMA*. 2014;312:902–914.
3. Burki TK. Increases in bilateral mastectomy for breast cancer. *Lancet Oncol*. 2014;15:e480.
4. Halsted CP, Benson JR, Jatoi I. A historical account of breast cancer surgery: beware of local recurrence but be not radical. *Future Oncol*. 2014;10:1649–1657.
5. Berbers J, van Baardwijk A, Houben R, et al. Reconstruction: before or after postmastectomy radiotherapy? A systematic review of the literature. *Eur J Cancer*. 2014;50:2752–2762.
6. Hugenoltz-Wamsteker W1, Robbeson C, Nijs J, Hoelen W, Meeus M. The effect of docetaxel on developing oedema in patients with breast cancer: a systematic review. *Eur J Cancer Care (Engl)*. October 27, 2014. [Epub ahead of print].
7. Yuan Q, Han J, Cong W. Docetaxel-loaded solid lipid nanoparticles suppress breast cancer cells growth with reduced myelosuppression toxicity. *Int J Nanomedicine*. 2014;9:4829–4846.
8. World Health Organization. Cancer Fact sheet 297. Key facts. Available from: <http://www.who.int/mediacentre/factsheets/fs297/en/>. Accessed November 14, 2012.
9. Federal Research Institute for Health Organization and Informatics of the Ministry of Health of the Russian Federation. Annual Statistics, 2012. Socially important diseases in the Russian population. Moscow, 2013. Available from: <http://www.mednet.ru/en.html>. Accessed November 14, 2014.

Authors' reply

Qing Yuan¹

Jing Han^{1,2}

Wenshu Cong¹

Ying Ge³

Dandan Ma^{1,3,4}

Zhaoxia Dai^{3,4}

Yaping Li⁵

Xiaolin Bi^{1,3,4}

¹CAS Key Laboratory for Biological Effects of Nanomaterials and Nanosafety, Institute of High Energy Physics, Chinese Academy of Sciences, Beijing, People's Republic of China; ²School of Life Sciences, Anhui University, Hefei, People's Republic of China; ³Cancer Center, Institute of Cancer Stem Cell, ⁴Graduate School, Dalian Medical University, Dalian, People's Republic of China; ⁵Shanghai Institute of Materia Medica, Chinese Academy of Sciences, Shanghai, People's Republic of China

Correspondence: Xiaolin Bi

CAS Key Laboratory for Biological Effects of Nanomaterials and Nanosafety, Institute of High Energy Physics, Chinese Academy of Sciences, Beijing 100049, People's Republic of China
Tel +86 10 8823 6709
Fax +86 10 8823 6456
Email bixl@ihep.ac.cn

Dear editor

As we all know, breast cancer is the most significant cause of mortality among females both in developing and developed countries recent years.^{1,2} In 2008, the percentage of breast cancer was 23% in total new cancer cases and death by breast cancer was 14% in total cancer death.² We now know that the breast cancer incidence is high in the Russian Federation from the analysis by Danilova et al. Based on their data, Danilova et al found that incidence of breast cancer was increased yearly while the mortality had been reduced continually in Russian Federation from 2003 to 2012.

The incidence rates and mortality rates of breast cancer between developing and developed countries showed some differences. In 2008, about 50% of the breast cancer cases occurred in developing countries and the percentage of cancer deaths is 60%.² While developed countries have the relative high incidence rates and the death rates have been decreasing continually. In contrast, incidence and mortality rates have been rising in many African and Asian countries such as Uganda and India.¹ These data indicated that breast cancer patients especially those in developing countries need more efficient and affordable therapy.

Systemic adjuvant chemotherapy can improve outcomes after surgery for breast cancer patients.³ Unfortunately, most adjuvant chemotherapeutic agents, including docetaxel-one of the most effective adjuvant therapy drugs, cause serious side

effect.³⁻⁶ Lots of efforts has been put into reducing side effects and enhancing antitumor activity, such as using nanotechnology to improve the formulations.⁷⁻¹⁴ Lipids are safe materials with good biocompatibility for drug formulations, some lipid-based formulations of anticancer drugs have been approved by the US Food and Drug Administration, such as Doxorubicin liposomal (New Drug Application (NDA) number 050718). To date, lipid-based nanoparticles have been proved to be one of the most promising drug-delivery candidates.^{15,16}

We prepared the docetaxel-loaded solid lipid nanoparticles (DSNs) which can significantly reduce the side-effect of docetaxel.^{14,17} Moreover, DSNs have lots of advantages compared with other nanoformulations, such as component safety, easier preparation, better stability, and controlled release etc that will promote its clinical application.^{14,18} These features make DSNs a potential economical adjuvant chemotherapeutic drug for breast cancer therapy with higher efficacy, especially affordable for patients in developing countries.

We appreciate Danilova et al proposed DSNs as a basic compound for a targeted and dose-sparing personalized breast cancer treatment strategy. There are a lot of problems that need to be overcome, such as large-scale production, targeting conjugation and on-demand release before DSNs' clinical use. We believe with the development of nanotechnology and pharmaceuticals, the more effective docetaxel formulations based on DSNs will be developed and applied in the future.

Acknowledgments

This work was supported by grants from National Basic Research Program of China (973 Program grant numbers 2010CB934004 and 2010CB934003), National Natural Science Foundation of China (grant number 31271480), Program of Changjiang Scholar and Innovative Research Team in University (IRT13049) and CAS Knowledge Innovation Program to Xiaolin Bi.

Disclosure

The author reports no conflicts of interest in this communication.

References

1. Jemal A, Bray F, Center MM, Ferlay J, Ward E, Forman D. Global cancer statistics. *CA: a cancer journal for clinicians*. 2011;61(2):69–90.
2. Ferlay J, Shin HR, Bray F, Forman D, Mathers C, Parkin DM. Estimates of worldwide burden of cancer in 2008: GLOBOCAN 2008. *Int J Cancer*. 2010;127(12):2893–2917.
3. Rampurwala MM, Rocque GB, Burkard ME. Update on adjuvant chemotherapy for early breast cancer. *Breast cancer (Auckl)*. 2014;8:125–133.

4. Saloustros E, Mavroudis D, Georgoulas V. Paclitaxel and docetaxel in the treatment of breast cancer. *Expert Opin Pharmacother*. 2008;9(15):2603–2616.
5. Crown J, O'Leary M, Ooi WS. Docetaxel and paclitaxel in the treatment of breast cancer: a review of clinical experience. *Oncologist*. 2004;9 Suppl 2:24–32.
6. Baker J, Ajani J, Scotte F, et al. Docetaxel-related side effects and their management. *Eur J Oncol Nurs*. 2009;13(1):49–59.
7. Hwang HY, Kim IS, Kwon IC, Kim YH. Tumor targetability and antitumor effect of docetaxel-loaded hydrophobically modified glycol chitosan nanoparticles. *J Control Release*. 2008;128(1):23–31.
8. Esmaili F, Dinarvand R, Ghahremani MH, Ostad SN, Esmaily H, Atyabi F. Cellular cytotoxicity and in-vivo biodistribution of docetaxel poly(lactide-co-glycolide) nanoparticles. *Anticancer Drugs*. 2010;21(1):43–52.
9. Yanasarn N, Sloat BR, Cui Z. Nanoparticles engineered from lecithin-in-water emulsions as a potential delivery system for docetaxel. *Int J Pharm*. 2009;379(1):174–180.
10. Immordino ML, Brusa P, Arpicco S, Stella B, Dosio F, Cattel L. Preparation, characterization, cytotoxicity and pharmacokinetics of liposomes containing docetaxel. *J Control Release*. 2003;91(3):417–429.
11. Elsabahy M, Perron ME, Bertrand N, Yu GE, Leroux JC. Solubilization of docetaxel in poly(ethylene oxide)-block-poly(butylene/styrene oxide) micelles. *Biomacromolecules*. 2007;8(7):2250–2257.
12. Liu J, Zahedi P, Zeng F, Allen C. Nano-sized assemblies of a PEG-docetaxel conjugate as a formulation strategy for docetaxel. *J Pharm Sci*. 2008;97(8):3274–3290.
13. Zheng D, Li D, Lu X, Feng Z. Enhanced antitumor efficiency of docetaxel-loaded nanoparticles in a human ovarian xenograft model with lower systemic toxicities by intratumoral delivery. *Oncol Rep*. 2010;23(3):717–724.
14. Gao Y, Yang R, Zhang Z, Chen L, Sun Z, Li Y. Solid lipid nanoparticles reduce systemic toxicity of docetaxel: performance and mechanism in animal. *Nanotoxicology*. 2011;5(4):636–649.
15. Puri A, Loomis K, Smith B, et al. Lipid-based nanoparticles as pharmaceutical drug carriers: from concepts to clinic. *Crit Rev Ther Drug Carrier Syst*. 2009;26(6):523–580.
16. Porter CJ, Trevaskis NL, Charman WN. Lipids and lipid-based formulations: optimizing the oral delivery of lipophilic drugs. *Nat Rev Drug Discov*. 2007;6(3):231–248.
17. Yuan Q, Han J, Cong W, et al. Docetaxel-loaded solid lipid nanoparticles suppress breast cancer cells growth with reduced myelosuppression toxicity. *Int J Nanomedicine*. 2014;9:4829–4846.
18. Zhang P, Chen L, Zhang Z, Lin L, Li Y. Pharmacokinetics in rats and efficacy in murine ovarian cancer model for solid lipid nanoparticles loading docetaxel. *J Nanosci Nanotechnol*. 2010;10(11):7541–7544.

International Journal of Nanomedicine

Publish your work in this journal

The International Journal of Nanomedicine is an international, peer-reviewed journal focusing on the application of nanotechnology in diagnostics, therapeutics, and drug delivery systems throughout the biomedical field. This journal is indexed on PubMed Central, MedLine, CAS, SciSearch®, Current Contents®/Clinical Medicine,

Submit your manuscript here: <http://www.dovepress.com/international-journal-of-nanomedicine-journal>

Dovepress

Journal Citation Reports/Science Edition, EMBase, Scopus and the Elsevier Bibliographic databases. The manuscript management system is completely online and includes a very quick and fair peer-review system, which is all easy to use. Visit <http://www.dovepress.com/testimonials.php> to read real quotes from published authors.