LETTER TO THE EDITOR

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Osmolarity: A hidden factor in Nanotoxicology

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Abstract

In the field of drug delivery, long circulating nanocarriers in the blood have many advantages such as targeted drug delivery and sustained release. Based on our current knowledge, evaluation of the effect of long circulating nanocarriers in the blood stream on osmolarity of plasma has not been reported before. In this study, osmotic pressure developed by some commercially available nanocarriers was estimated based on Van't Hoff equation. It is noteworthy that theoretically, nanocarriers do not have any significant effect on osmolarity of plasma. However, it is worth being evaluated experimentally in order to be taken into account in future studies.

Keywords: Nanocarriers, Drug delivery, Osmotic pressure

Letters to the editor

An osmosis phenomenon is the net movement of solvent (usually water) through a semipermeable membrane from a region of high water concentration (hypoosmolar solution) to a lower water concentration (hyperosmolar solution). Adding a solute to pure water decreases water concentration in the solution. In such conditions, water molecules can diffuse from a region of low solute concentration to one with a high solute concentration [1]. The effect of diverse solutes (i.e., molecules or ions) on osmolarity is depended on the number of dissolved particles in a solution, and is not correlated to their mass. Consequently, in an equal mass ratio, macromolecules (e.g., proteins, nucleic acids, polysaccharides) have much less influence on the osmolarity of a solution in comparison with their monomeric components. For example, a gram of a polysaccharide comprised of 1,000 glucose units and a milligram of glucose have the identical effect on osmolarity. In order to prevent of an enormous increase in osmolarity inside the storage cell (e.g., hepatocyte), the fuel is stored in the form of polysaccharide (i.e., starch or glycogen) rather than glucose or other simple sugars by the cells [2].

In the field of drug delivery, long circulating nanocarriers in the blood have many advantages such as targeted drug delivery and sustained release [4–6]. Until now, numerous studies have been performed in the Nanotoxicology [7, 8]. Nevertheless, evaluation of the effect of long circulating nanocarriers in the blood stream on osmolarity of plasma has not been reported before.

For this reason, osmotic pressure developed by some commercially available nanocarriers per 1 L of plasma estimated based on Van't Hoff equation. Data shown in Table 1 illustrates that theoretically, nanocarriers do not have any significant effect on osmolarity of plasma. However, it is worth being evaluated experimentally in order to be taken into account in future studies.

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Similarly, increasing osmolarity of plasma leads to net reabsorption of fluid from interstitial fluid into the capillaries rather than net filtration. On the other hand, increasing osmolarity of plasma leads to anti-diuretic hormone secretion. Osmolarity of plasma is approximately 300 mOsm/L. Change as small as 1 % in osmolarity of plasma leads to increasing antidiure-tic hormone secretion significantly. This hormone decreases the excreted volume of fluid by the kidneys. Ultimately, Due to increased blood volume, arterial pressure increases [1, 3].

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Table 1 Estimation of osmotic pressure developed by some commercially available nanocarriers per 1 L of plasma

Trade name	Osmotic pressure (mm Hg)	Osmolarity (mOsm/L)
Doxil	7.0×10^{-5}	4.0×10^{-6}
Abraxane	2.0×10^{-2}	1.0×10^{-3}
AmBisome	3.2×10^{-4}	1.7×10^{-5}
Marqibo	3.2×10^{-5}	1.7×10^{-6}
DepoCyt	3.0×10^{-5}	2.3×10^{-6}

Competing interests

The authors declare that they have no competing interests.

Authors' contributions

SM conceived of the study and participated in its design and carried out the calculation. OM participated in design of study and drafted the manuscript. RD participated in design and supervision of study and revised the manuscript. All authors read and approved the final manuscript.

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