## Nanochemistry and biotechnology

## Fullerene C<sub>60</sub> penetration into leukemic cells and its photoinduced effect on protein phosphotyrosine status

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Pristine fullerene  $C_{60}$  is nontoxic, able to accommodate inside hydrophobic regions of membranes [1], possesses photosensitizing potential with ability to produce reactive oxygen species and could be perspective for photodynamic therapy. Yet, still little is known about  $C_{60}$  interaction with cells of different types and the ways of its influence on cells signaling systems.

The aim was to study fullerene  $C_{60}$  penetration into cancer cells and its photocytotoxic effect against leukemic cells by estimation of protein tyrosine residues phosphorylation, high level of which is the marker of cell transformation.

With the use of fluorescent-labeled fullerene  $C_{60}$ –RITC the time-dependent accumulation of nanoparticles in cancer cells was confirmed by confocal microscopy. The decrease of the fluorescent signal of TMRE probe sensitive to the mitochondrial membrane polarization in leukemic cells treated with  $C_{60}$  was shown. To maximize the efficiency of fullerene  $C_{60}$  photoexcitation by visible light we use light-emitting diode lamp ( $\lambda_{max}=450$ , 500-600 nm). A comparative study showed that photoexcited  $C_{60}$  as well as  $H_2O_2$  (inducer of oxidative apoptosis) or inhibitor of tyrosine proteinkinases staurosporine (STS) decreased the level of proteins tyrosine phosphorylation, but the pattern of dephosphorylated proteins was different. Photoexcitation of fullerene  $C_{60}$  was followed by dephosphorylation of proteins, which still conserved high phosphotyrosine level after treatment with  $H_2O_2$  or STS. It is assumed that photoexcited  $C_{60}$  could be used for combined treatment with cytotoxic agents to block the elevation of protein phosphotyrosine level in leukemic cells.

Levi N., Hantgan R., Lively M. C<sub>60</sub>-Fullerenes: detection of intracellular photoluminescence and lack of cytotoxic effects// J Nanobiotechnology. – 2006. – 4. – P. 14–25.

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