

Femtosecond Laser Microscopy of Stem Cells

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ABSTRACT:

Femtosecond laser multiphoton microscopy is an already well-established method for *in vivo* imaging of biological cells. The use of innovative femtosecond laser techniques for imaging and nanoprocessing in stem cell research is the subject of this talk. The multiphoton-FLIM tomograph MPTflex was used to investigate stem cells' autofluorescence originated from the metabolic coenzymes NAD(P)H and FAD/flavoproteins. Significant differences on autofluorescence lifetime signatures have been identified in variety of stem cells as well as their differentiating counterparts¹. Moreover, metabolic activity of individual cells can also be detected within the complex culture of artificially generated induced pluripotent stem cell (iPS) colonies². Furthermore, femtosecond-laser based transfection found to be a superior method to realize contamination-free delivery of foreign molecules into single cells of interest³. In order to achieve transfection of a large cell population, a software-aided automatic and a continuous flow femtosecond laser-assisted cell transfection systems were realized^{4,5}. Such fs laser transfection systems are shown to be beneficial in optical cell reprogramming to generate iPS cells^{6,7}.

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References

1. Uchugonova, A and Hoffman, R.M., „Non-invasive single-photon and multi-photon imaging of stem cells and cancer cells in mouse models“. In König, K. (Ed.), Multiphoton Microscopy and Fluorescence Lifetime Imaging. Applications in Biology and Medicine. Berlin, Boston: De Gruyter, pp 411-421 (2018).
2. Uchugonova, A. "Multiphoton Autofluorescence Lifetime Imaging of Induced Pluripotent Stem Cells", J. Biomed. Opt. submitted, Paper Number: JBO 170125P (2017)
3. Tirlapur, UK., and König, K., "Targeted transfection by femtosecond laser", Nature 418(6895), 290-1 (2002)
4. Breunig, H.G., Uchugonova, A., König, K., „High-throughput continuous flow femtosecond laser-assisted cell optoporation and transfection“, J. Microsc. Techn., 77 (12): 974-979 (2014).
5. Breunig, H. G., Uchugonova, A., A. Batista., König, K., "Software-aided automatic cell optoporation with femtosecond laser pulses", Scientific Reports 5, 11185 (2015).
6. Uchugonova, A., Breunig, H.G., Batista, A., König, K., "Optical Reprogramming of human somatic cells with near infrared femtosecond laser pulses", J. Biomed. Opt. 20(11), 115008 (2015)
7. Uchugonova, A., Breunig, H.G., Batista, A., König, K., "Optical reprogramming of human cells with an ultrashort femtosecond laser microfluidic transfection platform", J Biophotonics 9(9), 942-7 (2016)