

The *in vivo* photothermal treatment of gold nanorod in the mouse ear model

Bruce Yao Wen Liu^{1,2}, Cheng-Lung Chen^{*1}, Shin-Yu Lee¹, Fu-Hsiung Chang³,
Win-Li Lin⁴, Chih-Ta Chia² and Yang-Yuan Chen^{**1,5}

¹*Institute of Physics, Academia Sinica, Taipei, Taiwan.*

²*Department of Physics, National Taiwan Normal University, Taipei, Taiwan*

³*Institute of Biochemistry and Molecular Biology, National Taiwan University, Taipei, Taiwan*

⁴*Institute of Biomedical Engineering, National Taiwan University, Taipei, Taiwan*

⁵*Graduate Institute of Applied Physics, National Chengchi University, Taipei, Taiwan*

(Received January 17, 2014, Revised March 18, 2014, Accepted March 18, 2014)

Abstract. Gold nanorod's exceptional light to heat transduction is a robust phenomenon that has been extensively verified. The phenomenon is a trait from which many novel applications across disciplines have been proposed. In this investigation, the feasibility of utilizing heat harvested from such photothermal method to combat cancer is presented. Using non-invasive laser methods, an *in vivo* study is conducted on mouse ear tumors administered with gold nanorods (Au NRs). An emphasis is placed on monitoring the tumor developments after photothermal treatments, over time. The findings reveal significant tumor growth suppression at a threshold laser power of 0.6 W/cm² lasting 2 minutes; this energy also brought about dramatic size reduction in treated tumors. Furthermore, the apparent formation of an eschar over the laser treated region indicates extensive hemorrhagic necrosis of the tumor tissue; a phenomenon implicative to the inhibition of angiogenesis.

Keywords: gold nanorod; photothermal; tumor; laser; cancer; necrosis

1. Introduction

Nano photothermal cancer therapeutics, the novel practice of heat treating malignant tumors using energy harvested from light activated nanomaterials, have received considerable interest for a while now; the method is advocated as, at the least, a strategically beneficial candidate for the next generation of cancer treatments (Menon *et al.* 2013). Au NRs in particular, with its biocompatibility and fascinating optical properties, have been adopted extensively in photothermal therapeutic studies (Ekici *et al.* 2008). However, further studies beyond petri dishes—i.e., studies that clinical or posing *in vivo* relevance at the least—are rather limited. Specifically, investigations into the variation in tumor sizes and their corresponding pathological characteristics subsequent to photothermal treatment should prove to be insightful. For *in vivo* studies that have been reported

*Corresponding author, Ph.D., E-mail: aabbss@phys.sinica.edu.tw

**Ph.D., E-mail: cheny2@phys.sinica.edu.tw

- Pourtier-Manzanedo, A., Vercamer, C. and Van-Belle E. (2003), "Expression of an Ets-1 dominant-negative mutant perturbs normal and tumor angiogenesis in a mouse ear model", *Oncogene*, **22**(12), 1795-1806.
- Pelz, J., Mollwitz, M. and Stremmel, C. (2004), "The impact of surgery and mild hyperthermia on tumor response and angiogenesis of malignant melanoma in a rat perfusion model", *Bmc. Cancer*, **4**(52), 1-9.
- Tong, L., Zhao, Y. and Huff, T.B. (2007), "Gold nanorods mediate tumor cell death by compromising membrane integrity", *Adv. Mater.*, **19**(20), 3136-3141.
- Zetter, B.R. (1998), "Angiogenesis and tumor metastasis", *Annu. Rev. Med.*, **49**(1), 407-424.
- Zharov, V.P., Galitovskaya, E.N. and Johnson, C. (2005), "Synergistic enhancement of selective nanophotothermolysis with gold nanoclusters: potential for cancer therapy", *Laser. Surg. Med.*, **37**(3), 219-226.

YC