A model to simulate laser ablation in vascularized tumor based on dynamic photothermal coupling interaction

Xiyang Zhang, Zuoran Li, Zhifang Li*, and Hui Li College of Photonic and Electronic Engineering, Fujian Normal University, Fujian Provincial Key Laboratory of Photonic Technology,

Key Laboratory of Optoelectronic Science and Technology for Medicine, Ministry of Education,



Fuzhou, Fujian 350007, China *lizhifang@fjnu.edu.cn

J.CN Fujian Normal U

Introduction:

- •In laser immunotherapy, the photothermal effect induces immune responses by destroying and interrupting tumor cells through temperature elevation in target tissue.
- •Tumor is rich in vascularture, and plays a critical part in photothermal effect.
- •We focus on the effect of the large vessel around tumor and the mircovessel in tumore on photothermal therapy.

Computational Methods:

•Photon propagation from laser light was described by diffusion approximation:

$$\frac{n}{c} \frac{\partial \varphi(x, y, z, t)}{\partial t} + \nabla \cdot (-D\nabla \varphi(x, y, z, t)) + u_a \times \varphi(x, y, z, t) = Q(x, y, z, t), \qquad (1)$$

•The bioheat equation is the realizing of the principle of conservation of the energy applied to tissue volume:

$$\rho C_{\rho} \frac{\partial T}{\partial t} + \rho C_{\rho} \mathbf{u} \cdot \nabla T + \nabla \cdot \mathbf{q} = Q + Q_{\text{bio}}$$

$$Q_{\text{bio}} = \rho_{\text{b}} C_{\rho,\text{b}} \omega_{\text{b}} (T_{\text{b}} - T) + Q_{\text{met}}$$

•3D base model of liver tissue consist of a cube of 10*10*10cm³ with the central in point(0mm,0mm,0mm) shown in Fig.1. The cube contains a tumor with a diameter 20mm and a blood vessel approximated by the cylindrical shape.

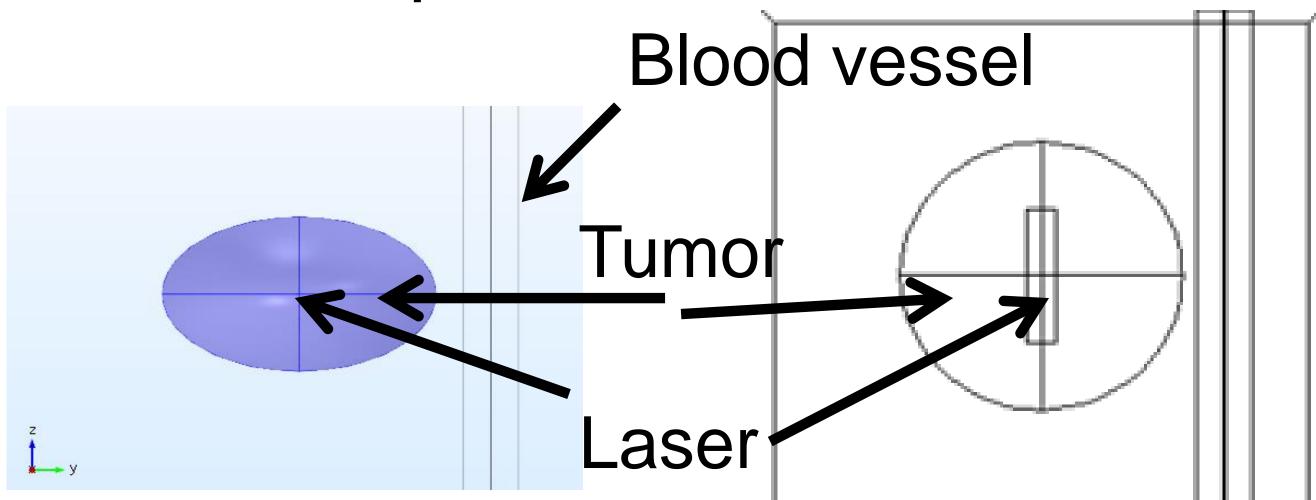


Figure 1. Tissue model

Results:

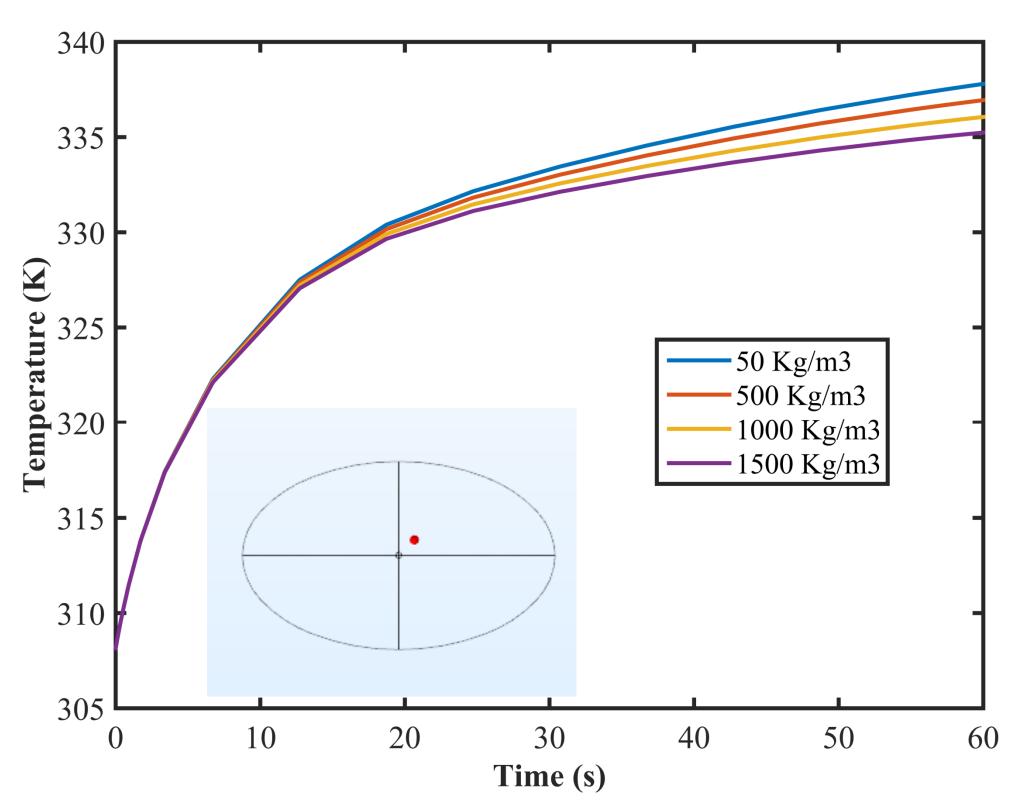


Figure 2. Temperature at point (0 0.05 0.05) vs time at the different blood densities

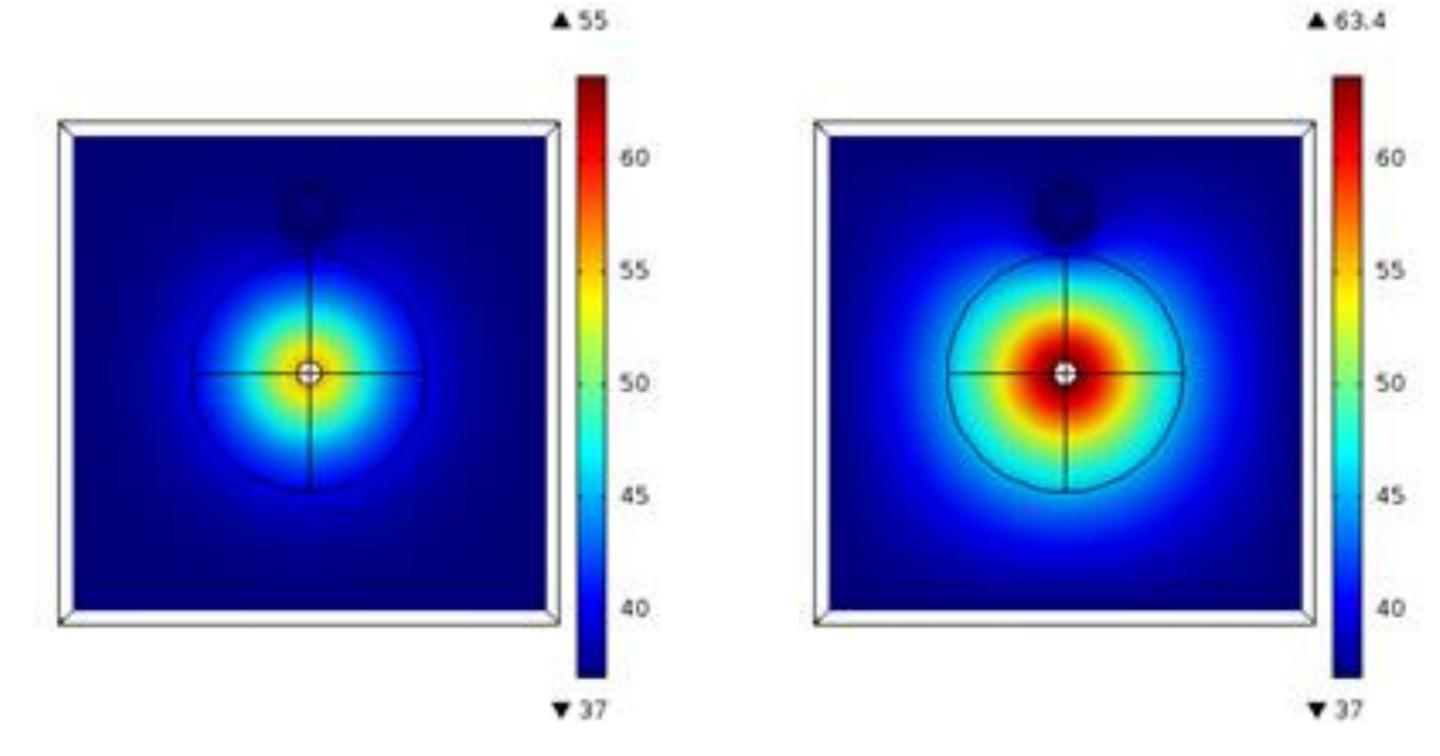


Figure 3. Cross-section images of the temperature distribution varying with irradiation time of (a) 150s; (b) 600s. The white parts denote light sources, big black circles are tumor regions, the small circles are blood vessels.

Conclusions:

Vascularized tumour grown within in a vascularized tissue displays a characteristic compartmentalization into essentially three regions: highly vascularized tumour, well-vascularized tumour and poorly vascularized tumour. These region could be described by blood density. Furthermore, the large blood vessel has the cooling effect. Finally, we should consider the sizes of tumor and light beam.

