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Numerical study on the three-dimensional temperature distribution according to laser conditions in photothermal therapy of periimplantitis

Jeeyong Paik¹, Donghyuk Kim¹, Hyunjung Kim^{1*} and Hee-Sun Kim^{2*}

Abstract

Purpose Dental implants have been successfully implemented as a treatment for tooth loss. However, periimplantitis, an inflammatory reaction owing to microbial deposition around the implant, can lead to implant failure. So, it is necessary to treat peri-implantitis. Therefore, this numerical study is aimed at investigating conditions for treating peri-implantitis.

Methods Photothermal therapy, a laser treatment method, utilizes photothermal effect, in which light is converted to heat. This technique has advantage of selectively curing inflamed tissues by increasing their temperature. Accordingly, herein, photothermal effect on peri-implantitis is studied through numerical analysis with using Arrhenius damage integral and Arrhenius thermal damage ratio.

Results Through numerical analysis on peri-implantitis treatment, we explored temperature changes under varied laser settings (laser power, radius, irradiation time). We obtained the temperature distribution on interface of artificial tooth root and inflammation and determined whether temperature exceeds or does not exceed 47°C to know which laser power affects alveolar bone indirectly. We defined the Arrhenius thermal damage ratio as a variable and determined that the maximum laser power that does not exceed 47°C at the AA' line is 1.0 W. Additionally, we found that the value of the Arrhenius thermal damage ratio is 0.26 for a laser irradiation time of 100 s and 0.50 for 500 s.

Conclusion The result of this numerical study indicates that the Arrhenius thermal damage ratio can be used as a standard for determining the treatment conditions to help assisted laser treatment for peri-implantitis in each numerical analysis scenario.

Keywords Dental implants, Peri-implantitis, Photothermal therapy, Inflammation, Numerical analysis, Computerassisted, Computer simulation

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Introduction

In patients with tooth loss, despite the extensive and demanding procedures as well as the considerable time and expense involved, the successful completion of implant placement surgery and the installation of the superstructure necessitates ongoing maintenance for the longevity and healthy use of the implant [1]. Peri-implantitis is an inflammation involving bone loss around the implant tissue owing to microbial infection, analogous to periodontitis in natural teeth [2–6]. Initially, the accumulation of biofilm on the implant surface leads to biofilm proliferation. This results in the formation of plaques on the implant surface, and as the gap between the gum and the implant starts to widen, inflammation occurs in the surrounding gum tissue. If this issue is unresolved, the biofilm can cause an inflammatory response that extends from the surrounding tissue to the inner bone, leading to the progressive destruction of the bone around the implant [7-10]. Peri-implantitis is known as the major cause of late-stage implant failure; while the treatment of the biofilm is essential for treatment, an ideal method for sterilizing the implant surface and restoring the health of the surrounding tissue has not yet been established [11-14].

Various methods, such as physical treatment using plastic curettes, local or systemic administration of antibiotics, local antiseptic treatments using chlorhexidine, and the use of air-abrasive devices, have been employed to treat the biofilm around implants [15-18]. However, mechanical treatment alone often fails to eliminate the biofilm [14, 19], and the use of chemical agents or antibiotics can be minimally effective or raise concerns regarding side effects [20]. Therefore, recent studies have focused on the use of dental lasers to sterilize and purify implant surfaces [21-23]. The laser could be used in either secondary prevention or raising levels of prevention for peri-implantitis [24]. This approach has advantages over traditional methods, including a lower likelihood of bleeding, faster recovery, and greater penetration depth, which facilitate the treatment of inflammation deep within the gingiva.

Photothermal therapy (PTT) is a laser-based method that treats inflammation by increasing the temperature through the heat generated by the laser. PTT operates based on the principle of the photothermal effect [25, 26]. The photothermal effect refers to the phenomenon in which light energy is absorbed by a specific substance and converted into heat [27]. When PTT is used to treat inflammation, the pattern of inflammatory destruction is determined by the location of the inflammation [28]. When inflammation is in direct contact with body tissues, it is necessary to differentiate between apoptosis and necrosis. However, if the inflammation is not in contact with body tissues, the focus can be solely on destruction within the inflammation itself, determining whether the damage is reversible or irreversible. However, if the temperature of the implant surface excessively increases, the heat transferred through the implant can cause thermal damage to the alveolar bone; thus, the temperature of the alveolar bone must be maintained below 47 $^{\circ}$ C [29, 30]. Irreversible tissue damage can be quantitatively assessed using the Arrhenius damage integral [31, 32].

Various studies have been conducted on the changes in the implant surface temperature when a laser is employed [33-35]. Matthias et al. [33] used a continuous-wave 809-nm diode laser to measure the temperature around the implant. The laser was positioned 0.5 mm from the implant surface and irradiated laser power at ranging from 0.5 to 2.5 W in increments of 0.5 W. After irradiation, the time required for the temperature to rise more than 10 °C from the initial temperature was measured Alessandro et al. [34] applied 810-nm and 980-nm diode lasers in both, pulsed- and continuous-wave modes to measure temperature changes. In this experiment, the laser was positioned at the upper part of the implant and irradiated (3-mm distance) in laser power 2.0 W and measured temperature changes after an irradiation period of 60 s were compared with before the laser irradiation. Deppe et al. [35] used a 445-nm diode laser to measure the temperature changes of five different types of implants and six irradiation conditions were used. After irradiation, the temperature effects of the 445-nm laser on five different types of implants were assessed. The results showed that, under the same laser intensity and irradiation conditions, the temperature changes varied depending on the type of implant. Although all these studies experimentally measured the temperature of the implant surface, they did not measure the temperature of the inflamed areas of the implant.

Most studies have focused solely on measuring the changes in the implant surface temperature, with insufficient research on the temperature of the inflammation itself [21, 36]. Therefore, in this numerical study, we investigated the effectiveness of various laser irradiation techniques in the adjunctive treatment of peri-implantitis, specifically focusing on identifying laser intensities that ensure the temperature of the surrounding alveolar bone does not exceed 47 °C. Following this, we employed the Pennes bioheat equation, to accurately calculate the temperature distribution within a cylindrical inflammation area around the implant. Also, using the Arrhenius damage integral, we quantitatively assessed the extent of irreversible inflammation damage and evaluated Arrhenius thermal damage ratio (ϕ_{Arrh}) values to find an appropriate value of laser power, radius and irradiation time. Our goal was to ensure an effective approach to peri-implantitis adjunctive treatment by photothermal therapy, while seeking scenarios where the ϕ_{Arrh} values were maximized without causing harm to the alveolar bone.

Methods

Heat transfer model

In this numerical study, after laser irradiation, the Pennes bioheat equation was used to determine the temperature of the surrounding tissues and inflammation around the implant [37]. This equation assumes that the heat generated by blood circulation and metabolic processes is uniformly distributed within the biological tissues. This equation is expressed as Eq. (1): Here, $k, \rho, \text{and} c_p$ represent the thermal conductivity, density, and specific heat, respectively. The term q_b denotes the heat transfer by the blood, whereas q_m represents the metabolic heat source. q_b is expressed in Eq. (2), and is calculated based on each material's blood perfusion rate (ω_b), blood density (ρ_b), blood specific heat ($C_{p,b}$), and the temperature difference between the initial blood temperature (T_b) and the current temperature. Moreover, as there is no blood flow or metabolic activity in the implant, the values of q_b and q_m in Eq. (1) become zero, reducing it to a simple heat-diffusion Eq.

$$\rho c_p \frac{\partial T}{\partial t} = k \nabla^2 T + q_b + q_m \tag{1}$$

$$q_b = \rho_b \omega_b c_{p,b} (T_b - T) \tag{2}$$

In this numerical study, as heat is applied to inflamed tissues through a 630 nm diode laser, the Pennes bioheat equation in Eq. (1) can be expressed in the form of Eq. (3), which includes a heat source term, q_l , owing to the laser. q_l is expressed in Eq. (4).

$$\rho c_p \frac{\partial T}{\partial t} = k \nabla^2 T + q_b + q_m + q_l \tag{3}$$

$$q_{l} = \mu_{a} \frac{P_{l}}{\pi r_{l}^{2}} e^{-\mu_{tot}z} \cdot e^{-\frac{x^{2}+y^{2}}{r_{l}^{2}}} (\mu_{tot} = \mu_{a} + \mu_{s}')$$
(4)

Equation (4) is suitable for use when the direction of the laser irradiation aligns with the *z*-direction. However, if the direction of the laser irradiation is inclined, Eq. (4) must be transformed via rotation to account for the angle. The treatment of peri-implantitis involves laser irradiation between the implant and the gingiva, indicating that an inclined laser is more realistic. Therefore, in this numerical study, to formulate the situation of an inclined laser, the positions of *x* and *z* were interchanged, and the concept of rotational transformation, along with differential lengths dx and dz, was used to consider the angle and position. Additionally, considering the change in the irradiated area owing to the laser's reflectivity and angle, Eq. (5) was derived in the form presented in [38].

$$q_{l} = (1 - R_{t}) \cdot \mu_{a} \frac{P_{l} \cdot \cos\theta}{\pi r_{l}^{2}} e^{-\mu_{tot}(-(x+dx)\cos\theta - (z-dz)\sin\theta)}$$

$$\cdot e^{-\frac{(-(x+dx)\sin\theta + (z-dz)\cos\theta)^{2} + y^{2}}{r_{l}^{2}}} (\mu_{tot} = \mu_{a} + \mu_{s}')$$
(5)

$$R_{t} = R_{1} + R_{2}, R_{1} = \left[\frac{\sqrt{n_{2}^{2} - (n_{1}sin\theta)^{2}} - n_{1}cos\theta}}{\sqrt{n_{2}^{2} - (n_{1}sin\theta)^{2}} + n_{1}cos\theta}\right]^{2}$$
(6)

Here, q_l represents the amount of heat applied to periimplantitis by the laser, and R_t , μ_a , P_l , r_l , and θ correspond to the reflectivity, light absorption coefficient, laser power, laser radius, and irradiation angle, respectively. The attenuation coefficient μ_{tot} is sum of μ_a and the reduced scattering coefficient μ'_s . The value of R_t is the sum of the specular reflection value R_1 and the diffuse reflection value R_2 . The value for specular reflection is expressed using Eq. (6), and the diffuse reflection value for inflammation at 630 nm was measured as 0.28 [39]. n_1 and n_2 represent the refractive indices of air and inflammation, respectively [40, 41]. Ultimately, by substituting Eq. (5) in Eq. (3), the temperature distribution within the medium (peri-implantitis and implant) can be calculated.

Arrhenius damage integral & variable

In this numerical study, the final goal of PTT was to maximize the irreversible damage caused by inflammation. Accordingly, the Arrhenius damage integral was utilized to evaluate the extent of damage [28]. This is expressed as presented in Eq. (7). Here, Ω represents the extent of damage to the inflammation, A is the frequency factor, which is a probabilistic variable indicating the likelihood of a reaction, E_a is the activation energy required to initiate the reaction, R is the ideal gas constant, and T(t)represents the temperature of the inflammation at time t [42, 43].

$$\Omega\left(t\right) = \int_{0}^{t} A e^{-\frac{E_{a}}{RT(t)}} dt$$
(7)

The Arrhenius damage integral is calculated using the temperature values (K) at each time point, the frequency factor of inflammation $A = 2.84 \times 10^{99} s^{-1}$, the activation energy of inflammation $E_a = 6.19 \times 10^5 J/mol$, and the ideal gas constant 8.314 $J/mol \cdot K$, after which the value of Ω is obtained [44, 45]. A value of Ω greater than 1 is considered to indicate irreversible damage. In numerical analysis, the Ω value can be obtained for each grid in all calculated areas, allowing for a quantitative extraction of



Fig. 1 Schematic of the numerical model



Fig. 2 3D model with the grid of the numerical model

the extent of irreversible damage within the peri-implantitis. To this end, the volume ratio of the part that has been irreversibly cured compared with the total inflammation is denoted as ϕ_{Arrh} , and this is defined as the Arrhenius thermal damage ratio.

$$\phi_{Arrh} = \frac{Inflammation \ Volume \ at \ \Omega(t) > 1}{Total \ Inflammation \ Volume}$$
(8)

Numerical geometry and properties

In this numerical study, PTT for inflammation caused by implants was evaluated through a numerical analysis. Figure 1 presents a schematic of the numerical model, with the image on the left showing the shape of the 3D model when viewed from the XZ plane and the image on the right showing the same from the YZ plane. Figure 2 illustrates the 3D model with the grid formed in the numerical analysis model. The entire calculation area was set as a rectangular prism encompassing air, gum, implant, and peri-implantitis, with a total height of 37 mm, a width of 30 mm, and a depth of 20 mm; the peri-implantitis was modeled as a cylindrical-shaped inflammation with a height of 2 mm and a diameter of 2 mm, attached to the artificial tooth root and abutment part, as shown in Fig. 2. The model consisted of the crown, gingiva, inflammation, artificial tooth root, abutment connecting the artificial tooth root and crown, alveolar bone, and air region. The wavelength of the irradiated laser was set to 630 nm, and the initial temperature (T_b) of the medium was set at 37 °C. The thermal and optical properties of each material are summarized in Table 1 [46].

Table 2 summarizes the numerical conditions. To depict PTT under various conditions, a numerical analysis was conducted across 765 cases, with the laser power (P_l) ranging from 0 to 2.0 W in increments of 0.04 W, the laser radius (r_l) ranging from 0.1 to 0.2 mm in increments of 0.05 mm, and the laser irradiation time (t) ranging from 100 to 500 s in intervals of 100 s. Additionally, the angle of the laser (θ) was fixed at 60°, and the laser was set to operate in the continuous-wave mode.

	ρ (kg/m³)	с (J/kgK)	<i>k</i> (W/mK)	<i>μα</i> (1/cm)	μ′ (1/cm)	ω _b (1/s)	<i>qm</i> (W/m³)
Crown (Zirconia)	6080	450	2.80	0.10	20.43	-	-
Gingival	1000	4200	0.63	0.530	3.817	0.0076	1091
Inflammation	1080	3500	0.48	2.16	17.03	0.009	65,400
Artificial tooth root (Ti-6Al-4 V)	4420	546	7.00	789,500	pprox 0	-	-
Abutment (Zirconia)	6080	450	2.80	0.10	20.43	-	-
Alveolar Bone	2060	1260	0.38	0.596	22.97	0.00369	-
Air	1.205	1.006	0.0256	0	0	-	-
Blood	1000	4200	-	-	-	-	-

Table 1 Various properties of the implant and surroundings [28, 47–56]

Table 2 Parameters of the numerical model

Parameter	Case	Number	Remarks
Laser power (P_l)	0 to 2.0 W	51	Intv. 0.04 W
Laser radius (r_l)	0.1 to 0.2 mm	3	Intv. 0.05 mm
Laser irradiation time (t)	100 to 500 s	5	Intv. 100 s

Results

Numerical analysis validation

The analysis tool used in this numerical study was COM-SOL Multiphysics, a powerful simulation software based on the finite element method. The backward differentiation formula method was applied to discretize the Pennes Bioheat equation, enabling the calculation of transient temperature distributions over time. Additionally, to validate the numerical analysis model propose herein, a grid independence test and convergence test based on the number of iterations were performed under the laser power of 1.0 W, the laser radius of 0.2 mm, and the irradiation time of 500 s, focusing on the temperature results in the area where the laser first contacts the inflammation. As observed in Fig. 3, when the number of grids reached 1,328,831 or more, the temperature change converged to less than 10^{-3} °C, setting the final number of grids at 1,328,831. Moreover, the error value in relation to the number of iterations dropped below $10^{-3}\%$ after 35 iterations, confirming convergence and validating the effectiveness of the numerical analysis model proposed in this numerical study.

Laser irradiation time and power behavior analysis in inflammation

In this numerical study, laser irradiation was focused on the central part of the inflammation (x=12.3 mm, y=0 mm, z=6 mm) to observe the effects of PTT depending on the laser intensity, radius, and irradiation time. To reiterate, the essence of PTT is to increase the temperature of inflammation to maximize irreversible damage. Although the temperature field was calculated in the numerical analysis for the entire model, to focus on the temperature rise in the inflammation owing to the laser, a plot was drawn for the area of x=11 mm to x=14 mm, y=-2 mm to y=2 mm, z=4 mm to z=8 mm, as shown in Fig. 4. Figure 4a shows a 3D schematic of the numerical analysis model showing the laser irradiation of the implant and inflamed parts. Figure 4b and c are schematics of the cross-sections in the XZ and YZ directions, respectively, at the inflammation center point (x=12.8 mm, y=0, z=6 mm). In Fig. 4b, the line AA'





Fig. 3 Mesh validation of the numerical model and iteration error in percentage



Fig. 4 Implant model in various directions

represents the tangent to the implant surface where the inflammation, abutment, and artificial tooth root meet.

First, with the laser radius fixed at 0.15 mm, the laser intensity was set to 0.6 W, 1.0 W, and 1.4 W, and the irradiation time was set to 100 s, 300 s, and 500 s. The temperature distribution under these settings and the corresponding thermal damage (irreversible damage) of the inflammation are depicted at the inflammation center point on the XZ plane (Fig. 5) and the YZ plane (Fig. 6).

In both Figs. 5 and 6, the black line represents where the value of the Arrhenius damage integral (Ω) equals 1, indicating that irreversible damage occurs where Ω exceeds 1. As observed in these figures, under the same laser power (Figs. 5 and 6 (a, b,c), (d, e,f), (g, h,i)), an increase in the laser irradiation time resulted in an increase in the temperature of the inflammation at the same location, and an expansion of the area where irreversible damage occurred.

Examining the temperature distribution within the inflammation, it was observed that the temperature spread in a slightly distorted shape owing to the influence of the laser irradiation angle. Temperature diffusion is a phenomenon that propagates in all directions. In the case of inflamed tissues, which differ from typical solids in having relatively lower light absorption coefficients, laser light penetrates from the surface to the interior of the inflamed tissue. Consequently, the temperature also appeared to spread in the direction of laser irradiation, as calculated accurately. Under the same laser irradiation time (Figs. 5 and 6 (a, d,g), (b, e,h), (c, f,i)), as the laser power increased, the temperature of the inflammation increased and the area suffering thermal damage

expanded. This indicated that, as the laser irradiation time and power increased, the extent of irreversible damage in inflammation increased. This can be interpreted as an increase in the total heat applied by the laser with longer irradiation times, allowing more time for heat to spread within the inflammation, thereby increasing the area of inflammation that undergoes irreversible damage. Additionally, as the laser power increased, the amount of heat applied per unit area to the inflammation increased, leading to an increase in the irreversible damage of inflammation. However, while inducing the irreversible damage of the inflammation is essential for the treatment of peri-implantitis, from the perspective of normal tissues, irreversible damage leads to tissue damage, and thus, an excessive temperature increase should be avoided.

The implant surface temperature is directly related to the inflammation's temperature. While laser energy absorption is localized within the inflammation, thermal conduction from the inflammation to implant can raises the implant surface temperature. The heat energy absorbed from the inflammation is transferred to the implant surface. If the temperature within the inflammation rises excessively due to overly intense laser power or prolonged irradiation times, it may result in the implant surface temperature exceeding the critical threshold of 47 °C. This scenario poses a risk of thermal damage to the alveolar bone. Consequently, it is necessary to examine the temperature on the implant surface, which is the hottest part of the implant (AA' line). As shown in Fig. 7, under all the laser radii and irradiation time conditions, the temperature at point AA, where the



Fig. 5 Temperature and Ω distribution with laser irradiation at various laser powers and laser irradiation times (XZ plane direction) for a laer radius of 0.15 mm

implant physically contacts the inflammation and is the hottest, was examined. Below 1.0 W, even after t=300 s, the implant surface temperature generally did not reach 47 °C, except in some cases. However, above 1.0 W, the temperature of a significant portion of the implant surface exceeded 47 °C; thus, caution is warranted during treatment.

Laser radius and power behavior analysis in inflammation

In Sect. 3.2, we analyzed the degree of irreversible damage and the temperature distribution of the inflammation according to the laser irradiation time and intensity in various planar directions (XZ direction, YZ direction). In Sect. 3.3, we fixed the laser intensity at 0.8 W and set the laser irradiation radius to 0.1 mm, 0.15 mm, and 0.2 mm, and the irradiation time to 100 s, 300 s, and 500 s. Subsequently, we plotted the temperature distribution and the corresponding results under these conditions.

Figure 8 shows the temperature distribution results and the corresponding thermal damage to the inflammation when the shape in Fig. 4a is cut along the XZ plane at the inflammation center point. Figure 9 shows the temperature distribution results and the thermal damage to the inflammation when the shape in Fig. 4a is cut along the YZ plane at the inflammation center point. As mentioned in Sect. 3.2, the black line represents where the Arrhenius damage integral value (Ω) is 1, and if Ω exceeds 1, it indicates irreversible damage. Observing Figs. 8 and 9, when the laser intensity was fixed, a smaller laser radius increased the intensity per unit area, concentrating the temperature increase at the center of the inflammation. Conversely, as the laser radius increased, the area covered by the laser also increased, showing a similar trend in the extent of irreversible damage of inflammation across different laser radii (Figs. 8 and 9 (a, b,c), (d, e,f), (g, h,i)).

However, as will be discussed in Sect. 3.4, with laser powers above 0.8 W, the Arrhenius thermal damage ratio (ϕ_{Arrh}) varied depending on the radius as the laser power increased. Additionally, at the same laser power, an increase in the irradiation time led to an increase in the area of irreversible damage of inflammation for all laser radii, as calculated. As shown in Fig. 7, the temperature difference at AA' for each radius is negligible,



Fig. 6 Temperature and Ω distribution with laser irradiation at various laser powers and laser irradiation times for a laser radius of 0.15 mm (YZ plane direction)

and at 0.8 W, there is no impact on the alveolar bone. However, as previously described, although the ϕ_{Arrh} value increases with increasing laser intensity, caution is required during treatment because of its potential impact on the alveolar bone.

Arrhenius thermal damage ratio (ϕ_{Arrh}) in inflammation

In Sect. 3.2 and 3.3, we examined the extent of reversible damage and temperature distribution in the XZ and YZ planes inside the inflammation. In Sect. 3.4, we assessed the extent of irreversible damage throughout the entire volume of inflammation. As mentioned in Sect. 2.2, to evaluate the extent of damage across the entire inflammation, we used the Arrhenius thermal damage ratio (ϕ_{Arrh}), which, as Eq. (8) shows, represents the ratio of the volume in which the value of the Arrhenius damage integral exceeds 1 to the total volume of the inflammation.

(a) laser irradiation time: 100 s, (b) laser irradiation time: 200 s, (c) laser irradiation time: 300 s, (d) laser irradiation time: 400 s, and (e) laser irradiation time: 500 s.

Figure 10 shows the values of ϕ_{Arrh} for different laser radii as the laser irradiation time increases, with the red

line representing the point at 1.0 W. For all the laser radii, the ϕ_{Arrh} value increased as both the irradiation time and laser intensity increased. Notably, as the laser intensity increased, the variation in ϕ_{Arrh} values depending on the laser radius also became larger. In addition, at t=100s, the ϕ_{Arrh} value was approximately 0.75 for the maximum laser intensity (2.0 W) at a laser radius of 0.2 mm. This indicated that approximately 75% of the inflammation was eradicated. At t=500 s, with the maximum laser intensity (2.0 W), the ϕ_{Arrh} value was close to 1, indicating that almost all the inflammation was eradicated. It is necessary to examine the ϕ_{Arrh} values at laser intensities below 1.0 W, where the temperature at the surface (AA' line) where the inflammation and the implant meet approaches 47 °C (as shown in Fig. 7). Thus, the ϕ_{Arrh} value obtained at a laser irradiation time of 100 s was approximately 0.26, and it reached approximately 0.5 when the irradiation time was extended to 500 s.

In Fig. 11, the effect of the laser radius on ϕ_{Arrh} is depicted. With increasing laser radius, the ϕ_{Arrh} value increased for the same laser irradiation duration. Furthermore, the variation in ϕ_{Arrh} values for each



Fig. 7 Temperature distribution of line AA' with various radii and irradiation times of the laser



Fig. 8 Temperature and Ω distribution with laser irradiation at various laser irradiation radii and laser irradiation times (XZ plane direction) for a laser power of 0.8 W

irradiation time became more pronounced at higher laser intensities, and this amplification was notably greater than that caused by changes in the laser radius.

The values of ϕ_{Arrh} are similar to the results shown in Fig. 10, where conditions close to ϕ_{Arrh} being 1 were obtained at the laser intensity of 2.0 W, the irradiation time of 500 s, and the laser radii of 0.15 mm and 0.2 mm. In addition, at a laser intensity less than 1.0 W, a maximum ϕ_{Arrh} of approximately 0.5 was achieved. Note that, as shown in Fig. 7, the AA' line (the line where the inflammation contacts the implant) is more than 2 mm away from the alveolar bone. Therefore, the temperature rise along the AA' line undergoes a conduction process from the artificial tooth root to the alveolar bone, and the temperature at the contact area between the alveolar bone and the artificial tooth root is lower. Further analysis could lead to the discovery of laser intensities higher than that (1.0 W) set in Figs. 10 and 11 to avoid thermal damage to the alveolar bone, potentially achieving higher ϕ_{Arrh} values without causing thermal damage to the alveolar bone.

Discussion

This numerical study identifies optimal laser irradiation conditions that effectively treat inflammation around implants while minimizing damage to the alveolar bone, through a numerical analysis of the effects of laser intensity, radius, and irradiation time on temperature distribution. By understanding the impact of these variables on temperature distribution, the numerical study lays a foundation for managing the risk of thermal damage, crucial in implant therapy.

The temperature distribution in the biological tissues and implants was determined using the Pennes bioheat equation, and the heat from the laser was modeled by modifying the existing laser heat formula to consider tilted laser irradiation. The wavelength of the laser was 630 nm, and the angle of the laser was fixed at 60°. Additionally, by utilizing the Arrhenius damage integral, which was calculated from the temperature distribution to assess the PTT effect quantitatively, the analysis differentiated between areas that underwent irreversible damage and those that did not. Finally, using the Arrhenius thermal damage ratio (ϕ_{Arrh}), the extent of irreversible



Fig. 9 Temperature and Ω distribution with laser irradiation at various laser irradiation radii and laser irradiation times (YZ plane direction) for a laser power of 0.8 W

damage throughout the entire inflammation was calculated under all the analysis conditions.

From the analysis results, at a laser intensity of 0.8 W or less, the variation in ϕ_{Arrh} values based on the laser radius was minimal. However, at laser intensities above 0.8 W, a greater deviation in ϕ_{Arrh} values were observed as the laser radius increased. Additionally, the increase in ϕ_{Arrh} values was quantitatively assessed as the laser irradiation time increased. The temperature distribution on the surface (AA' line) where the inflammation contacts the implant was also analyzed to consider the thermal damage to the alveolar bone. Based on the analysis of the implant surface (AA' line) temperature, it was confirmed that the thermal conduction process from the artificial tooth root causes no impact on the alveolar bone, which is situated more than 2 mm away. Consequently, conditions that could completely eradicate the inflammation without affecting the alveolar bone were identified. Moreover, the critical observation that the AA' line's temperature (the interface between the implant and inflammation) does not reach the temperature threshold when using specific laser conditions emphasizes the precision with which laser therapy can be applied. Specifically, our findings at laser intensities below 1.0 W, which prevented the temperature from exceeding the critical threshold of 47 $^{\circ}$ C, highlight the potential for optimizing laser conditions to maximize therapeutic effects while safeguarding alveolar bone.

By exploring the potential benefits of photothermal therapy as a non-surgical treatment method for periimplantitis, our findings offer promising insights into the application of photothermal therapy in clinical settings. By identifying laser irradiation conditions that effectively treats inflammation without thermal damage to the alveolar bone, the results of this numerical study suggest that more rigorous treatments can be performed in practical peri-implantitis laser therapy. Specifically, by establishing laser irradiation guidelines that prevent alveolar bone damage while effectively treating inflammation through precise thermal management, the numerical study represents significant progress in the field of dental implants. One limitation of this numerical study is that the numerical analysis model may not fully implement the complexity of actual biological conditions. The parameters



(e)



1.5

1.0

Laser Power(W)



Fig. 10 Arrhenius thermal damage ratio for various laser irradiation times

0.5

0.0



Fig. 11 Arrhenius thermal damage ratio for various laser irradiation radii (a) laser irradiation radius: 0.1 mm, (b) laser irradiation radius: 0.15 mm, and (c) laser irradiation radius: 0.2 mm

assumed in the modeling process might not consider all the variables present in practical clinical setting. Future studies could be of significant value, including investigations of thermal conduction through the artificial tooth root, the distance between the alveolar bone and the laser irradiation site, changes in laser irradiation angle, various shapes of inflammation, and expanding the analysis of irreversible eradication conditions to not only inflammation, but also surrounding tissues, such as bones and gingiva.

Conclusions

This research establishes a fundamental comprehension of the optimization of photothermal therapy for effective treatment of peri-implantitis, with a concurrent focus on preserving the alveolar bone integrity against thermal damage. As the results, the Arrhenius thermal damage ratio (ϕ_{Arrh}), the extent of irreversible damage throughout the entire inflammation was calculated under various the laser intensity, laser radius and laser irradiation times. Our findings highlight the promise of non-surgical photothermal therapy in dental implantology, which provides a precise, regulated way for treating peri-implant inflammation. It is

expected that the results and numerical analysis model presented in this numerical study will aid in establishing a method for the thermal adjunctive treatment of peri-implantitis.

Nomenclature

- frequency factor (1/s) Α
- $\overset{c_{\rho}}{E}_{a}$ specific heat (J/kgK)
- activation energy (J/mol)
- k thermal conductivity (W/mK)
- P intensity of laser (W)
- qvolumetric heat source (W/m³)
- r_l laser radius (mm)
- R_t reflectivity
- ideal gas constant (J/molK) R
- temperature (K)
- laser irradiation time (s) t

Greek symbols

- θ irradiation angle
- reduced optical coefficient (1/cm) μ'
- density (kg/m³) ρ
- ϕ_{Arrh} Arrhenius thermal damage ratio
- blood perfusion rate (1/s) ω_h
- Ω Arrhenius damage integral value
- μ optical coefficient (1/cm)

Subscripts

- absorption а
- b blood
- laser
- metabolic m
- scattering
- total tot
- notation of direction X, Y, Z

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Authors' contributions

Conceptualization: J. P., D. K., H-S. K, and H. K.; data curation: J. P.; formal analysis: J. P. and D. K.; funding acquisition: H. K.; investigation: J. P. and D. K.; methodology: J. P. and D. K.; project administration: H-S. K, and H. K.; Resources: H. K.; Software: J. P.; Supervision: H-S. K, H. K.; validation, J. P.; visualization, J. P.; writing—original draft, J. P.; writing—review and editing, H-S. K, H. K. All the authors have read and approved the final version of the manuscript.

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Data availability

All data generated or analyzed in this study are included in this published article.

Declarations

Ethics approval and consent to participate Not applicable.

Consent for publication

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Competing interests

The authors declare that they have no competing interests.

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References

- Adell R, Lekholm U, Rockler BRÄNEMARK, Brånemark PI. A 15-year study of osseointegrated implants in the treatment of the edentulous jaw. Int J Oral Sura. 1981:10(6):387-416.
- Larsson L, Kavanagh NM, Nguyen TV, Castilho RM, Berglundh T, Giannobile 2. WV. Influence of epigenetics on periodontitis and peri-implantitis pathogenesis. Periodontol 2000. 2022;90(1):125-37.
- Kormas I, Pedercini C, Pedercini A, Raptopoulos M, Alassy H, Wolff LF. Peri-3 implant diseases: diagnosis, clinical, histological, microbiological characteristics and treatment strategies. A narrative review. Antibiotics. 2020;9(11):835.
- Prathapachandran J, Suresh N. Management of peri-implantitis. Dent Res J. 4 2012;9(5):516.
- Lindhe J, Meyle J, Group D of the European Workshop on Periodontology. 5 Peri-implant diseases: consensus report of the sixth European workshop on periodontology. J Clin Periodontol. 2008;35:282-5.
- 6 Froum SJ, Rosen PS. A proposed classification for peri-implantitis. Int J Periodontics Restor Dentistry. 2012;32(5):533.
- Lasserre JF, Brecx MC, Toma S. Oral microbes, biofilms and their role in peri-7. odontal and peri-implant diseases. Materials. 2018;11(10):1802.
- Belibasakis GN. Microbiological and immuno-pathological aspects of peri-8. implant diseases. Arch Oral Biol. 2014;59(1):66-72.
- 9. Fragkioudakis I, Tseleki G, Doufexi AE, Sakellari D. Current concepts on the pathogenesis of peri-implantitis: a narrative review. Eur J Dentistry. 2021;15(02):379-87.
- 10. Salvi GE, Cosgarea R, Sculean A. Prevalence and mechanisms of peri-implant diseases. J Dent Res. 2017;96(1):31-7.
- 11. Berglundh T, Persson L, Klinge B. A systematic review of the incidence of biological and technical complications in implant dentistry reported in prospective longitudinal studies of at least 5 years. J Clin Periodontol. 2002;29:197-212.
- 12. Atieh MA, Alsabeeha NH, Faggion Jr CM, Duncan WJ. The frequency of peri-implant diseases: a systematic review and meta-analysis. J Periodontol. 2013:84(11):1586-98.
- 13. Esposito M, Grusovin MG, Worthington HV. (2012). Interventions for replacing missing teeth: treatment of peri-implantitis. Cochrane database of systematic reviews. (1).
- 14. Roos-Jansåker AM, Renvert S, Egelberg J. Treatment of Peri-implant infections: a literature review. J Clin Periodontol. 2003;30(6):467-85.
- 15. Muthukuru M, Zainvi A, Esplugues EO, Flemmig TF. Non-surgical therapy for the management of peri-implantitis: a systematic review. Clin Oral Implants Res. 2012;23:77-83.
- 16. Schwarz F, Becker K, Renvert S. Efficacy of air polishing for the non-surgical treatment of Peri-implant diseases: a systematic review. J Clin Periodontol. 2015;42(10):951-9.
- 17. Kotsovilis S, Karoussis IK, Trianti M, Fourmousis I. Therapy of peri-implantitis: a systematic review. J Clin Periodontol. 2008;35(7):621-9.
- 18. Ntrouka VI, Slot DE, Louropoulou A, Van der Weijden F. The effect of chemotherapeutic agents on contaminated titanium surfaces: a systematic review. Clin Oral Implants Res. 2011;22(7):681-90.
- 19. Schwarz F, Sculean A, Romanos G, Herten M, Horn N, Scherbaum W, Becker J. Influence of different treatment approaches on the removal of early plaque biofilms and the viability of SAOS2 osteoblasts grown on titanium implants. Clin Oral Invest. 2005;9:111-7.
- 20. Schou S, Holmstrup P, Jørgensen T, Skovgaard LT, Stoltze K, Hjørting-Hansen E, Wenzel A. Implant surface preparation in the surgical treatment of experimental peri-implantitis with autogenous bone graft and ePTFE membrane in cynomolgus monkeys. Clin Oral Implants Res. 2003;14(4):412-22.
- 21. Leja C, Geminiani A, Caton J, Romanos GE. Thermodynamic effects of laser irradiation of implants placed in bone: an in vitro study. Lasers Med Sci. 2013;28:1435-40.
- 22. Rios FG, Viana ER, Ribeiro GM, González JC, Abelenda A, Peruzzo DC. Temperature evaluation of dental implant surface irradiated with high-power diode laser. Lasers Med Sci. 2016;31:1309-16.
- 23. Fenelon T, Bakr M, Walsh LJ, George R. Effects of lasers on titanium dental implant surfaces: a narrative review. Lasers Dent Sci. 2022;6(3):153-67.

- Sanz M, Herrera D, Kebschull M, Chapple I, Jepsen S, Berglundh T, Wennström J. Treatment of stage I–III periodontitis—the EFP S3 level clinical practice guideline. J Clin Periodontol. 2020;47:4–60.
- 25. Jiao J, Guo Z. Thermal interaction of short-pulsed laser focused beams with skin tissues. Phys Med Biol. 2009;54(13):4225.
- Sahoo N, Ghosh S, Narasimhan A, Das SK. Investigation of non-fourier effects in bio-tissues during laser assisted photothermal therapy. Int J Therm Sci. 2014;76:208–20.
- Singh R, Das K, Mishra SC, Okajima J, Maruyama S. Minimizing tissue surface overheating using convective cooling during laser-induced thermal therapy: a numerical study. J Therm Sci Eng Appl. 2016;8(1):011002.
- Ren Y, Qi H, Chen Q, Ruan L. Thermal dosage investigation for optimal temperature distribution in gold nanoparticle enhanced photothermal therapy. Int J Heat Mass Transf. 2017;106:212–21.
- 29. Bashutski JD, D'Silva NJ, Wang HL. Implant compression necrosis: current understanding and case report. J Periodontol. 2009;80(4):700–4.
- Huang CC, Liu YC, Chen LW, Chen YC. Temperature rise of alveolar bone during dental implant drilling using the finite element simulation. Life Sci J. 2010;7(1):68–72.
- 31. Pearce JA. Comparative analysis of mathematical models of cell death and thermal damage processes. Int J Hyperth. 2013;29(4):262–80.
- 32. Pearce JA. Relationship between Arrhenius models of thermal damage and the CEM 43 thermal dose. Energy-based Treatment of Tissue and Assessment V. Volume 7181. SPIE; 2009, February. pp. 35–49.
- Kreisler M, Schoof J, Langnau E, Haj A, H., d'Hoedt B. Temperature elevations in endosseous dental implants and the peri-implant bone during diodelaser-assisted surface decontamination. Lasers in Dentistry VIII. Volume 4610. SPIE; 2002, June. pp. 21–30.
- Geminiani A, Caton JG, Romanos GE. Temperature change during non-contact diode laser irradiation of implant surfaces. Lasers Med Sci. 2012;27:339–42.
- Deppe H, Ahrens M, Behr AV, Marr C, Sculean A, Mela P, Ritschl LM. Thermal effect of a 445 nm diode laser on five dental implant systems: an in vitro study. Sci Rep. 2021;11(1):20174.
- Kreisler M, Haj A, H., d'Hoedt B. Temperature changes induced by 809-nm GaAlAs laser at the implant–bone interface during simulated surface decontamination. Clin Oral Implants Res. 2003;14(1):91–6.
- Falahatkar S, Nouri-Borujerdi A, Mohammadzadeh A, Najafi M. Evaluation of heat conduction in a laser irradiated tooth with the three-phase-lag bio-heat transfer model. Therm Sci Eng Progress. 2018;7:203–12.
- Zhang YL, Wang YM, Huang AP, Hu X. Study on reflection characteristics of underwater target and laser echo power. Optoelectron Lett. 2020;16(2):137–42.
- Einstein G, Udayakumar K, Aruna PR, Koteeswaran D, Ganesan S. Diffuse reflectance spectroscopy for monitoring physiological and morphological changes in oral cancer. Optik. 2016;127(3):1479–85.
- Li J, Xu J, Lian Z, Yu Z, Yu H. Fabrication of antireflection surfaces with superhydrophobic property for titanium alloy by nanosecond laser irradiation. Opt Laser Technol. 2020;126:106129.
- Gowda RB, Saara K, Sharan P. Detection of oral cancerous cells using highly sensitive one-dimensional distributed Bragg's Reflector Fabry Perot Microcavity. Optik. 2021;244:167599.

- 42. Gas P, Kurgan E. (2018, June). Evaluation of thermal damage of hepatic tissue during thermotherapy based on the arrhenius model. In *2018 Progress in Applied Electrical Engineering (PAEE)* (pp. 1–4). IEEE.
- Wang SL, Qi H, Ren YT, Chen Q, Ruan LM. Optimal temperature control of tissue embedded with gold nanoparticles for enhanced thermal therapy based on two-energy equation model. J Therm Biol. 2018;74:264–74.
- 44. He X, Bhowmick S, Bischof JC. (2009). Thermal therapy in urologic systems: a comparison of arrhenius and thermal isoeffective dose models in predicting hyperthermic injury.
- 45. Sapareto SA. (1982). The biology of hyperthermia in vitro. *Physical aspects of hyperthermia*, (8).
- 46. Ghalichia F, Behniab S, Sadigh Radb E. (2004, February). Calculations for Ultrasonic Transducer Design and Temperature Tracking for Hyperthermia in Cervical Cancer. In Proceedings of the Iranian Conference on Biomedical Engineering, Tehran, Iran (Vol. 11).
- Jaglarz J, Szewczenko J, Marszałek K, Basiaga M, Marszałek M, Gaweł R. Nonstandard optical methods as a tool for rough surface analysis. Mater Today: Proc. 2015;2(7):4046–52.
- Abreu AI, Lima TP, Meireles AB, Souza BE, Torres LAG. Evaluation of surface temperature of tongue for screening of patients with suspected oral cancer. Sci. 2019;7:1–13.
- Fernandez-Oliveras A, Rubiño M, Pérez MM. (2013). Scattering and absorption properties of biomaterials for dental restorative applications. *Journal of the European Optical Society-Rapid publications*, 8.
- 50. Addit Manuf. 2019;26:166–179. https://doi.org/10.1016/j.addma.2019.02.001.
- Известия Саратовского университета. Новая серия. Серия: Физика, 20(4), 258–267.
- Rabbani Arshad S, Oskui Z, I., Hashemi A. Thermal analysis of dental implants in mandibular premolar region: 3D FEM study. J Prosthodont. 2018;27(3):284–9.
- 53. Bender R, Gross M, Luo W, Tess E. Thermal modeling of tissue ablation during pulsed CO2 laser gingivectomy. for Gum Reshaping and Reduction; 2014.
- Bargo PR, Prahl SA, Goodell TT, Sleven RA, Koval G, Blair G, Jacques SL. In vivo determination of optical properties of normal and tumor tissue with white light reflectance and an empirical light transport model during endoscopy. J Biomed Opt. 2005;10(3):034018–034018.
- Dawood SN, Al-Zahawi AR, Sabri LA. Mechanical and thermal stress behavior of a conservative proposed veneer preparation design for restoring misaligned anterior teeth: a 3D finite element analysis. Appl Sci. 2020;10(17):5814.
- Bashkatov AN, Genina EA, Kochubey VI, Tuchin VV. Optical properties of human cranial bone in the spectral range from 800 to 2000 nm. Saratov fall Meeting 2005: Optical technologies in Biophysics and Medicine VII. Volume 6163. SPIE; 2006, July. pp. 306–16.

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