scientific reports



OPEN Two-dimensional biothermomechanical effects in a layer of skin tissue exposed to variable thermal loading using a fourth-order MGT model

Sami F. Megahid^{1,2⊠}

A key consideration in medical procedures like thermal therapy is the danger of thermal harm to skin tissues from exposure to fluctuating thermal loads. To maximize treatment effectiveness while safeguarding healthy tissues, it is crucial to accurately anticipate and manage this damage, especially in hyperthermia therapy. The fourth-order Moore-Gibson-Thompson (4MGT) idea is employed in this study to lay a theoretical foundation for bioheat analysis. The purpose of this work is to clarify how skin tissues respond biothermally to varying thermal loading. The model developed makes it easier to anticipate the thermal reactions that occur in human skin and to estimate the efficiency of biothermal transfer in biological tissues. For the suggested model, a two-dimensional skin layer is utilized. The analytical results for tissue temperature are obtained using the normal mode approach. Both the impact of the duration of heat loading exposure and thermal damage are examined. Furthermore, the accuracy of the suggested model is evaluated by contrasting the obtained analytical results with accepted theories. The findings show that when the thermal relaxation time constant is included, the modified Moore-Gibson-Thomson biothermal model forecasts a decrease in temperature compared to the Pennes model

Keywords Two-dimensional skin tissue, Fourth-order MGT thermoelastic model, Variable thermal loading, Normal mode method

Many thermal treatment methods are specifically designed to target injured skin tissues in the medical profession with the least amount of damage to surrounding undamaged tissue. Heat ablation has shown great promise in the medical field for laser procedures, freckle removal, wound healing, and cancer treatment. There are several methods for generating heat when it comes to thermal ablation. RF waves, infrared radiation, microwave radiation, thermoseeds with magnetic excitability, and other heat emitters are among them. It's crucial to keep in mind, nevertheless, that high temperatures can damage healthy cells because they can cause denaturation of proteins and rupture of cellular membranes2. The greatest possible therapeutic outcomes and patient health depend on the careful regulation of temperature gradients inside the tissues. The projected temperature distribution must be known before starting thermal treatments because it is challenging to monitor comprehensive real-time temperature charts. More research is required to advance our understanding of how humans' bodies respond to hyperthermia.

The heat effects of blood in biological tissues, including perfusion of blood and convection, should be considered to accurately characterize the mechanisms of thermal transfer in living things, especially with regard to heat therapeutic benefits. Physiological processes also need to be considered, especially the production of metabolic heat. First introduced by Pennes in 1948, the Pennes bioheat equation is a widely used mathematical model for studying heat transport in biological tissues, particularly skin³. It extends the standard heat equation by incorporating blood circulation and metabolic heat generation⁴. Numerous heat transport mechanisms in tissues are taken into account by the model, including conduction, metabolic heat production, and blood perfusion. The Pennes bioheat equation is used in many different contexts, such as thermal ablation, cryotherapy,

¹Department of Mathematics, Faculty of Science, Mansoura University, Mansoura 35516, Egypt. ²Department of Mathematics, Faculty of Science, New Mansoura University, New Mansoura City 35712, Egypt. □ email: samifikri@mans.edu.eg

and hyperthermia, to predict temperature distribution and potential thermal damage within tissues. However, the model has limitations, including neglecting the pulsatility and nonuniformity of blood flow and assuming constant temperature characteristics. Several changes and modifications have been proposed to improve the accuracy and applicability of the Pennes bioheat equation in various biological systems⁵. One important use of the Pennes bioheat equation is thermal therapy, which employs heat to kill malignant cells or tissues. The equation facilitates the optimization of treatment parameters and minimizes side effects by forecasting the temperature distribution within the tissue during treatment. Additionally, it can be utilized to predict heat damage in the tissue, which aids in assessing the efficacy of a treatment.

Fu et al.'s primary focus was on how tumor form affected the distribution of skin temperature during thermal therapy⁷. They used the Pennes model to evaluate the heat response in this scenario. Similar to this, Singh and Repaka⁸ investigated the effects of radiofrequency ablation on the thermal characteristics of breast tumors using a multilayered breast model that includes the effects of temperature-dependent components like electrical and thermal conductivities. Zhang et al.⁹ employed the Pennes bioheat transfer model in steady-state conditions to investigate the effect of skin tissue temperature on blood perfusion rate. Temperature was found to have a linear, quadratic, and exponential association with perfusion. The model was solved using the boundary reciprocity approach.

Despite being widely utilized for a wide range of technical and thermal challenges, the standard Fourier heat conduction model has limitations in several situations. It's interesting to note that experimental research has demonstrated that the Pennes bioheat transfer (parabolic) model can yield inaccurate results, particularly in situations with significant heat flow or extremely brief intervals¹⁰. This discrepancy arises from the Penne model's inability to take tissue thermal inertia into consideration, which can lead to serious errors in temperature prediction. The thermal wave model is predicated on the premise that the temperature gradient and energy transfer happen quickly, which may not always be the case, even if it can accurately replicate biological materials¹¹. Because biological tissues are naturally complex and non-uniform, there is a temporal delay in both the temperature gradient and energy transfer as well as the heat flux and temperature gradient. In order to account for thermal inertia, Cattaneo and Vernotte^{12,13} created the concept of thermal relaxation time. This number illustrates the time lag between the heat flux vector and the corresponding temperature gradient, particularly in processes requiring rapid heat transfer. Liu et al.¹⁴ acknowledged this limitation by proposing a novel bioheat transport model that uses a thermal wave approach. By incorporating the thermal relaxation period into the Pennes equation, this enhanced model offers a more accurate representation of unstable thermal transmission mechanisms in living organisms or in laboratory environments.

The thermal wave model accurately predicts the time response of heat transport on a small scale. However, it has a limited capacity to catch minute spatial thermal exchanges. Tzou et al. ¹⁵ created the dual-phase lag (DPL) model to address this problem and investigate rapid heat transport in nonuniform materials. This model incorporates phase delays due to thermal inertia and microstructural interactions. The thermal wave bio-heat transfer and DPL models were then used to analyze heat transfer processes in biological tissues, taking into account heat sources including lasers ^{16–18}, microwaves ¹⁹, and ultrasound ²⁰.

Generalized thermoelasticity theories, which assume a finite speed for thermal signals, have been of considerable interest for over 40 years $^{21-24}$. These theories employ a hyperbolic heat equation in contrast to the standard coupled thermoelasticity (CT) theory, which is predicated on a parabolic heat equation and infinite heat propagation speed. By substituting a new heat transfer model for the standard Fourier law, Lord and Shulman (LS) 25 were the first to generalize for isotropic materials, enabling the creation of a wave-type heat equation. A second generalization, the thermoelasticity theory (GL), which incorporates the rate of temperature change, was then presented by Green and Lindsay (GL) 26 . Green and Naghdi (GN) 27 extended the theoretical framework with the GN theory, which deviates from previous models by omitting thermal energy dissipation.

The Moore–Gibson–Thompson (MGT) equation has attracted a lot of research interest because of its practical applicability in a range of fields, including industry and healthcare²⁷. High-intensity ultrasound, which has numerous applications in lithotripsy, thermal therapy, and ultrasonic cleaning, has also been the subject of numerous investigations²⁷. In lithotripsy, the temperature distribution that emerges from simulating the passage of ultrasonic waves through the body can be predicted using the MGT equation and other generalized thermoelasticity theories^{28,29}. Quintanilla^{30,31} developed a novel approach to heat transmission in thermoelasticity by incorporating the MGT equation into the thermal conductivity equation. This model adds a relaxation component to the structure of the third-kind Green–Naghdi model³². Building on Quintanilla's work, the references^{33–35} examined several new thermoelastic systems and their applications. The thermomechanical response to various engineering challenges and the complexities of fluid dynamics have also been studied using the modified model^{36–38}.

The main goal of this work is to develop a more thorough and improved framework for thermoelasticity that better represents the relationship between temperature and elasticity than earlier models. This new model combines the Moore-Gibson-Thompson (MGT) equation with the Lord-Shulman and Green-Naghdi type III (GN-III) thermoelastic theories. Higher-order temporal derivatives of heat flow are incorporated into the updated model to accurately account for the intricate reactions of skin tissues to heating.

One of the study's main accomplishments is the model's superiority over previous generalized thermoelasticity models due to its ability to effectively depict the behavior of skin tissues. Furthermore, the suggested paradigm provides improved predictions and insights into the thermoelastic response of skin tissues by clarifying the complex link between temperature and elasticity.

Skin tissues subjected to varying heat stress are directly modeled using the fourth-order MGT thermal model. The analytical results for tissue are obtained using the normal mode approach. The behavior of the system is examined using a range of methodologies, including graphical and analytical approaches. The paper discusses important topics like the frequency of angular thermal loading and the consequences of thermal relaxation in

addition to thermal damage. When compared to experimental data, the results demonstrate that the proposed model is very accurate, emphasizing its ability to improve healthcare planning and execution.

Mathematical equations

Heat transport in skin tissues is governed by the well-known and widely used Fourier's law of heat transfer. According to Fourier's law, a material's internal temperature gradient controls how quickly heat passes through it. The following equation represents this association:

$$\overrightarrow{h}(\overrightarrow{p}, t) = -\mathcal{K}_t \nabla \theta (\overrightarrow{p}, t). \tag{1}$$

In this equation, \mathcal{K}_t stands for the skin tissue's thermal conductivity, whereas \overrightarrow{h} indicates the heat flux at a certain point in time t and place \overrightarrow{p} . The variable $\theta = T - T_b$ represents the temperature differential between arterial blood (T_b) and tissue (T).

Within living biological tissues, heat is transferred by three primary mechanisms: heat conduction, convective heat transfer, and blood perfusion. Fourier's law of heat conduction states that when thermal energy is transferred through a material due to a temperature gradient, heat conduction occurs inside the tissue. The blood's movement promotes convective heat transmission between the blood and the surrounding tissue, which in turn facilitates thermal energy exchange. Among the factors influencing this process are the blood vessel surface area, the temperature differential between the blood and tissue, and the blood flow rate. Blood perfusion, or the flow of blood through capillaries and other blood vessels within the tissue, also has a significant effect on the entire process of heat transmission. The interaction of blood flow with surrounding tissue affects the complex thermal dynamics observed in biological systems.

Among the models developed to answer Fourier's law are those put forth by Green and Naghdi^{27,32}. Some of the models that have been developed are these. There are several ways to display the augmented Fourier law from the GN-III model, including

$$\overrightarrow{h} = -\mathcal{K}_t \nabla \theta - \mathcal{K}_t^* \nabla \varnothing , \qquad (2)$$

with \mathcal{K}_t^* representing the rate of thermal conductivity. $\nabla \varnothing$ is regarded as a novel constitutive variable in Green and Naghdi's ideas. Additionally, \varnothing is read as the thermal-displacement function, which is the equivalent of the mechanical displacement in mechanical fields. $\frac{\partial \mathscr{D}}{\partial t} = \theta$ is satisfied by the thermal displacement function. Quintanilla³¹ created a new version of the GN-III heat conduction model inside the context of the MGT

equation by adding the relaxation parameter τ_h . The updated Fourier formula was as follows:

$$\left(1 + \tau_h \frac{\partial}{\partial t} + \frac{1}{2} \tau_h^2 \frac{\partial^2}{\partial t^2}\right) \overrightarrow{h} = -\mathcal{K}_t \nabla \theta - \mathcal{K}_t^* \nabla \varnothing.$$
(3)

The MGT model is based on the following: Viscoelastic Behavior: The material possesses both viscous and elastic qualities, which means that both the rate of deformation and the existing deformation determine how the material reacts to stress or strain and Nonlocal Effects: The model takes into account nonlocal interactions, which means that a point's behavior is dependent on both the local conditions and the conditions in the surrounding area. The modeling of distributed interactions and effects, like dispersive waves, is made possible by this incorporation.

Since it describes the connection between temperature, the heat transfer coefficient, and energy flow, the energy balance equation for heat transfer is among the most crucial basic equations that may be applied in the field of heat transfer analysis. The phrase for this is

$$\rho_{t} c_{t} \frac{\partial \theta}{\partial t} + T_{b} \gamma_{t} \frac{\partial e}{\partial t} + \nabla \cdot \overrightarrow{h} = S_{t}, \tag{4}$$

where γ_t is the thermal modulus, which is determined using the formula $\gamma_t = \alpha_t (3\lambda_t + 2\mu_t)$, c_t is the tissue specific heat, and ρ_t is the tissue density. In this case, the character α_t stands for the thermal expansion coefficient, whereas the symbols μ_t and λ_t represent Lame's constants. The symbol for the cubical dilatation, e, is

$$e = u_{i,i} = \operatorname{div} \overrightarrow{\mathbf{u}},$$
 (5)

where u_i stands for each of the displacement's constituent parts. S_t takes into consideration both external and interior heat sources, as indicated by ^{39,40}

$$S_t = S_b + S_{met} + S_{ext}. (6)$$

The amount of heat produced by tissue metabolism is represented by S_{met} , the heat energy from an external source by S_{ext} , and the heat source from blood circulation by S_b .

After finishing the time derivation process for Eqs. (3) and (4) and combining them, the fourth-order MGT thermal conductivity equation can be produced as follows:

$$\left[1 + \tau_{h} \frac{\partial}{\partial t} + \frac{1}{2} \tau_{h}^{2} \frac{\partial^{2}}{\partial t^{2}}\right] \left[\rho_{t} c_{t} \frac{\partial^{2} \theta}{\partial t^{2}} + T_{b} \gamma_{t} \frac{\partial^{2} e}{\partial t^{2}} - \frac{\partial}{\partial t} \left(S_{b} + S_{met} + S_{ext}\right)\right] \\
= \mathcal{K}_{t} \nabla^{2} \frac{\partial}{\partial t} + \mathcal{K}_{t}^{*} \nabla^{2} \theta. \tag{7}$$

In addition to the basic equation of thermoelasticity, equations of motion and constitutive relationships are crucial components when developing thermoelastic models. The following formula applies to the general versions of these equations 41,42

$$\rho_{t} \frac{\partial^{2} u_{i}}{\partial t^{2}} = \lambda_{t} u_{j,ij} + \mu_{t} (u_{j,ij} + u_{i,jj}) + \rho_{t} F_{i} - \gamma_{t} \theta_{,i}, \tag{8}$$

$$\sigma_{ij} = 2\mu_t e_{ij} + \delta_{ij} \left[\lambda_t e_{kk} - \gamma_t \theta \right], \tag{9}$$

$$2e_{ij} = u_{i,j} + u_{j,i}. (10)$$

The symbol σ_{ij} represents the stress tensor in the given framework, while the strain tensor is represented by the notation e_{ij} . The symbol F_i represents the vector components of the external body force. The Kronecker delta function is represented by the symbol δ_{ij} .

Mathematical formulation for the problem

The case of a 2D skin tissue in the interval $0 \le x \le L_x$ and $0 \le y \le L_y$ will be covered in this section (see Fig. 1). Cartesian coordinates (x, y, z) will be used in order to meet the needs of the challenge. In addition to the time variable t, the domain variables under study will be functions of x and y coordinates because the problem is 2D. The plane x=0 will serve as the free surface, and the x-axis will be plumb downward into the medium's depth.

By considering both thermal and mechanical boundary conditions, the proposed problem was resolved. The skin tissue's outside surface was believed to be heated in a time-dependent way and to be traction-free, whereas the inner surface was believed to be traction-free and insensitive to temperature changes. The following boundary conditions are therefore made possible:

$$\theta (x, y, t) = h(y, t), \ \sigma_{xx}(x, y, t) = \sigma_{xy}(x, y, t) = 0 \ at \ x = 0, \\ \theta (x, y, t) = \sigma_{xx}(x, y, t) = \sigma_{xy}(x, y, t) = 0 \ at \ x = L_x,$$
 (11)

where the heat load applied to the skin tissue's outer surface is denoted by $h\left(y,\,t\right)$.

The displacement components have the following form

$$u_x = u_x(x, y, t), u_y = u_y(x, y, t), u_z = 0.$$
 (12)

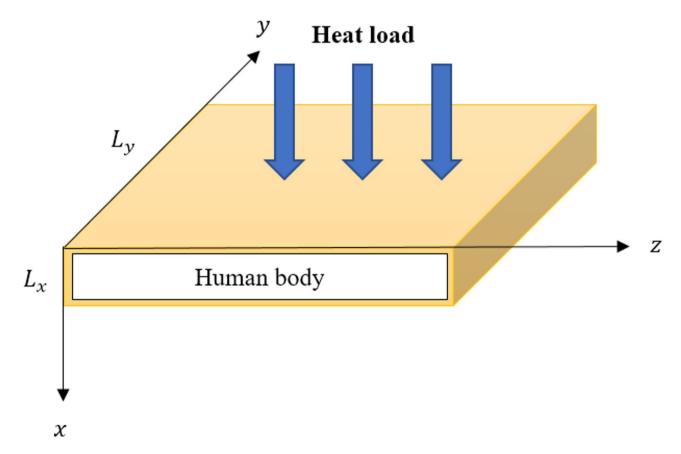


Fig. 1. Schematic diagram of skin tissue exposed to heat load.

Equations (10) and (12) can be used to derive the strain components listed below

$$e_{xx} = \frac{\partial u_x}{\partial x}, \ e_{yy} = \frac{\partial u_y}{\partial y}, \ e_{xy} = \frac{1}{2} \left(\frac{\partial u_x}{\partial y} + \frac{\partial u_y}{\partial x} \right), \ e_{xz} = e_{yz} = e_{zz} = 0.$$
 (13)

It is possible to obtain the stress components by applying Eqs. (9) and (13) as

$$\sigma_{xx} = (\lambda_t + 2\mu_t) \frac{\partial u_x}{\partial x} + \lambda_t \frac{\partial u_y}{\partial y} - \gamma_t \theta, \qquad (14)$$

$$\sigma_{yy} = (\lambda_t + 2\mu_t) \frac{\partial u_y}{\partial y} + \lambda_t \frac{\partial u_x}{\partial x} - \gamma_t \theta, \qquad (15)$$

$$\sigma_{xy} = \mu_t \left(\frac{\partial u_x}{\partial y} + \frac{\partial u_y}{\partial x} \right). \tag{16}$$

Equations (5) and (12) can be substituted into Eq. (8) to produce the following

$$(\lambda_t + \mu_t) \frac{\partial e}{\partial x} + \mu_t \nabla^2 u_x - \gamma_t \frac{\partial \theta}{\partial x} = \rho_t \frac{\partial^2 u_x}{\partial t^2}, \tag{17}$$

$$(\lambda_t + \mu_t) \frac{\partial e}{\partial y} + \mu_t \nabla^2 u_y - \gamma_t \frac{\partial \theta}{\partial y} = \rho_t \frac{\partial^2 u_y}{\partial t^2}.$$
 (18)

The current analysis makes use of the following blood perfusion term⁴³

$$S_b = -w_b \rho_b c_b \theta . {19}$$

The heat exchange between the tissue and the blood flow is described by this phrase, where w_b is the rate of blood perfusion, ρ_b is the blood mass density, and c_b is the blood specific heat.

The temperature of the tissue has an impact on the body's metabolic heat generation S_{met} , especially for skin tissue that contains blood vessels. The metabolic heat source S_{met} is represented as follows⁴⁴

$$S_{met} = S_{im}(1 + c_m \theta), \tag{20}$$

where S_{im} is the rate at which heat is generated by the baseline metabolic process, and c_m is the temperature coefficient that influences the rate of metabolic heat synthesis.

The heat conduction Eq. (7) can be written as follows when the heat energy coming from an external source is absent ($S_{ext} = 0$):

$$\left[1 + \tau_{h} \frac{\partial}{\partial t} + \frac{1}{2} \tau_{h}^{2} \frac{\partial^{2}}{\partial t^{2}}\right] \left[\rho_{t} c_{t} \frac{\partial^{2} \theta}{\partial t^{2}} + T_{b} \gamma_{t} \frac{\partial^{2} e}{\partial t^{2}} - (c_{m} S_{im} - w_{b} \rho_{b} c_{b}) \frac{\partial \theta}{\partial t}\right] \\
= \left(\mathcal{K}_{t} \frac{\partial}{\partial t} + \mathcal{K}_{t}^{*}\right) \nabla^{2} \theta. \tag{21}$$

The addition of dimensionless variables simplifies the problem's governing mathematical equations. Equations (17), (18), and (21) are made dimensionless by adding the following dimensionless values

By substituting the dimensionless variables from Eq. (31) into Eqs. (17), (18), and (21), we reduce the fundamental equations as follows (It removes the prime notation)

$$\left(\beta^2 - 1\right) \frac{\partial e}{\partial x} + \nabla^2 u_x - \beta^2 \frac{\partial \theta}{\partial x} = \beta^2 \frac{\partial^2 u_x}{\partial t^2},\tag{23}$$

$$\left(\beta^2 - 1\right) \frac{\partial e}{\partial y} + \nabla^2 u_y - \beta^2 \frac{\partial \theta}{\partial y} = \beta^2 \frac{\partial^2 u_y}{\partial t^2}, \tag{24}$$

$$\left(\epsilon + \frac{\partial}{\partial t}\right) \nabla^2 \theta = \left(1 + \tau_h \frac{\partial}{\partial t} + \frac{1}{2} \tau_h^2 \frac{\partial^2}{\partial t^2}\right) \left[\frac{\partial^2 \theta}{\partial t^2} - \epsilon_1 \frac{\partial \theta}{\partial t} + \epsilon_2 \frac{\partial^2 e}{\partial t^2}\right],\tag{25}$$

where

$$\beta^2 = \frac{\lambda_t + 2\mu_t}{\mu_t}, \ \epsilon = \frac{\mathcal{K}_t^*}{\mathcal{K}_t \zeta \rfloor^2}, \ \epsilon_1 = \frac{w_b \rho_b c_b - S_{im} c_m}{\rfloor^2 \zeta^2 \mathcal{K}_t}, \ \epsilon_2 = \frac{\gamma_t^2 T_b}{\rho_t c_t (\lambda_t + 2\mu_t)}. \tag{26}$$

The constitutive formulas diminish to

$$\sigma_{xx} = \left(\beta^2 - 2\right)e + 2\frac{\partial u_x}{\partial x} - \beta^2 \theta, \qquad (27)$$

$$\sigma_{yy} = (\beta^2 - 2) e + 2 \frac{\partial u_y}{\partial y} - \beta^2 \theta, \qquad (28)$$

$$\sigma_{xy} = \frac{\partial u_x}{\partial y} + \frac{\partial u_y}{\partial x}.$$
 (29)

After being differentiated with regard to x and y, respectively, Eqs. (33) and (34) are added

$$(\nabla^2 - \frac{\partial^2}{\partial t^2})e - \nabla^2 \theta = 0. \tag{30}$$

Normal mode solution and damage prediction

The following type of normal mode can be used to break down the solution of the many variables being studied

$$(\theta, u_x, u_y, \sigma_{ij}, e) = \left(\bar{\theta}, \bar{u_x}, \bar{u_y}, \bar{\sigma_{ij}}, \bar{e}\right)(x) \exp(\mathrm{i}wy + ft), \tag{31}$$

where f is the complex frequency constant, i is an imaginary unit, and w is the wave number in the y-direction. Equation (31), when inserted into Eqs. (25) and (30), yields

$$[(f+\epsilon)(D^2-w^2) - (1+\tau_h f + \frac{1}{2}\tau_h^2 f^2) f(f-\epsilon_1)] \bar{\theta} (x) = (1+\tau_h f + \frac{1}{2}\tau_h^2 f^2) \epsilon_2 f^2 \bar{e} (x),$$
(32)

$$(D^2 - w^2 - f^2) \bar{e}(x) - (D^2 - w^2) \bar{\theta}(x) = 0,$$
(33)

where $D = \frac{d}{dx}$.

The differential equation below is obtained by taking the variables θ (x) or e (x) out of Eqs. (32) and (33)

$$(D^4 - AD^2 + B) \{ \bar{e}(x), \bar{\theta}(x) \} = 0, \tag{34}$$

where

$$A = 2w^{2} + A_{1}, B = w^{4} + B_{1}w^{2} + B_{2},$$

$$A_{1} = f^{2} + \epsilon (f - \epsilon_{1} + \epsilon_{2}), B_{1} = f^{2} + \epsilon (f - \epsilon_{1} + f\epsilon_{2}),$$

$$B_{2} = \epsilon f^{2} (f - \epsilon_{1}), = \frac{f(1 + \tau_{h} f + \frac{1}{2}\tau_{h}^{2}f^{2})}{f + \epsilon}.$$
(35)

Equation (34) can be factored as follows

$$(D^{2} - k_{1}^{2}) (D^{2} - k_{2}^{2}) \{\bar{e}(x), \bar{\theta}(x)\} = 0,$$
(36)

where k_i^2 (j = 1, 2) is the following characteristic equation's root

$$k^4 - Ak^2 + B = 0. (37)$$

Therefore, Eq. (34)'s solution is provided by

$$\left\{ \bar{e}(x), \bar{\theta}(x) \right\} = \sum_{n=1}^{2} \left[\{R_n, M_n\} e^{k_n x} + \{R_{n+2}, M_{n+2}\} e^{-k_n x} \right], \tag{38}$$

where the parameters R_l and M_l (l = 1, 2, 3, 4) depend on w and f.

The following relationship is found when Eq. (38) is replaced into Eq. (33)

$$\{M_n, M_{n+2}\} = \frac{k_n^2 - w^2 - f^2}{k_n^2 - w^2} \{R_n, R_{n+2}\}.$$
(39)

Upon substituting Eq. (39) into Eq. (38), we obtain

$$\left\{ \bar{e}(x), \bar{\theta}(x) \right\} = \sum_{n=1}^{2} \left\{ 1, \frac{k_n^2 - w^2 - f^2}{k_n^2 - w^2} \right\} \left[R_n e^{k_n x} + R_{n+2} e^{-k_n x} \right]. \tag{40}$$

The displacement u_x is found by inserting Eqs. (31) and (40) into Eqs. (23) to get

$$(D^{2} - m^{2}) \bar{u_{x}}(x) = \sum_{n=1}^{2} \frac{k_{n} (k_{n}^{2} - w^{2} - \beta^{2} f^{2})}{k_{n}^{2} - w^{2}} \left[R_{n} e^{k_{n} x} - R_{n+2} e^{-k_{n} x} \right],$$
(41)

where

$$m^2 = w^2 + \beta^2 f^2. (42)$$

The solution to Eq. (41) is provided by

$$\bar{u_x}(x) = N_1 w^2 e^{mx} + N_2 w^2 e^{-mx} + \sum_{n=1}^{2} \frac{k_n}{k_n^2 - w^2} [R_n e^{k_n x} - R_{n+2} e^{-k_n x}],$$
(43)

where the parameters N_1 and N_2 are dependent on w and f.

Using the cubical dilatation and Eq. (31), we can determine that

$$\bar{u_y}(x) = -\frac{i}{w} \left(\bar{e}(x) - \frac{d \bar{u_x}(x)}{dx} \right). \tag{44}$$

Equations (40) and (43) may be swapped into Eq. (44) to yield

$$\bar{u_y}(x) = iw \left\{ N_1 m e^{mx} - N_2 m e^{-mx} + \sum_{n=1}^{2} \frac{1}{k_n^2 - w^2} \left[R_n e^{k_n x} + R_{n+2} e^{-k_n x} \right] \right\}.$$
 (45)

The stress components can be derived from Eqs. (27)-(29) using Eqs. (40), (43), and (45) as

$$\sigma_{xx}^{-}(x) = 2N_1 w^2 m e^{mx} - 2N_2 w^2 m e^{-mx} + \sum_{n=1}^{2} \frac{2w^2 + \beta^2 f^2}{k_n^2 - w^2} \left[R_n e^{k_n x} + R_{n+2} e^{-k_n x} \right], \quad (46)$$

$$\sigma_{yy}^{-}(x) = -2N_1 w^2 m e^{mx} + 2N_2 w^2 m e^{-mx} + \sum_{n=1}^{2} \frac{-2k_n^2 + \beta^2 f^2}{k_n^2 - w^2} \left[R_n e^{k_n x} + R_{n+2} e^{-k_n x} \right], \quad (47)$$

$$\sigma_{xy}^{-}(x) = iw \left\{ \left(w^2 + m^2 \right) \left(N_1 e^{mx} + N_2 e^{-mx} \right) + \sum_{n=1}^{2} \frac{2k_n}{k_n^2 - w^2} \left[R_n e^{k_n x} - R_{n+2} e^{-k_n x} \right] \right\}.$$
(48)

The normal mode analysis is used to find the solution in the Fourier-transformed domain. Assuming that the normal mode analysis for these functions can occur because all of the relations on the real line are smooth enough.

The following equations emerge from plugging the considered variable expressions into the boundary conditions (11)

$$\sum_{n=1}^{2} \frac{k_n^2 - w^2 - f^2}{k_n^2 - w^2} [R_n + R_{n+2}] = \bar{h}, \tag{49}$$

$$\sum_{n=1}^{2} \frac{k_n^2 - w^2 - f^2}{k_n^2 - w^2} [R_n e^{k_n L_x} + R_{n+2} e^{-k_n L_x}] = 0,$$
 (50)

$$\sum_{n=1}^{2} \frac{2w^2 + \beta^2 f^2}{k_n^2 - w^2} \left[R_n + R_{n+2} \right] + 2N_1 w^2 m - 2N_2 w^2 m = 0, \tag{51}$$

$$\sum_{n=1}^{2} \frac{2w^2 + \beta^2 f^2}{k_\pi^2 - w^2} \left[R_n e^{k_n L_x} + R_{n+2} e^{-k_n L_x} \right] + 2N_1 w^2 m e^{mL_x} - 2N_2 w^2 m e^{-mL_x} = 0, \quad (52)$$

$$\sum_{n=1}^{2} \frac{2k_n}{k_n^2 - w^2} \left[R_n - R_{n+2} \right] + \left(w^2 + m^2 \right) (N_1 + N_2) = 0, \tag{53}$$

$$\sum_{n=1}^{2} \frac{2k_n}{k_n^2 - w^2} \left[R_n e^{k_n L_x} - R_{n+2} e^{-k_n L_x} \right] + \left(w^2 + m^2 \right) \left(N_1 e^{mL_x} + N_2 e^{-mL_x} \right) = 0.$$
 (54)

It is possible to solve the above linear system and determine the integration constants R_l (l=1, 2, 3, 4), N_1 and N_2 .

The burn evaluation states that thermal injury occurs when the temperature of the basal layer, which acts as the interface between the epidermis and dermis, rises above 44 °C⁴⁵. The evaluation of thermal damage is essential for living tissue and its therapeutic uses in bioengineering. The first to introduce a numerical approach for evaluating thermal damage was Moritz and Henriques⁴⁶. In this assessment, the damage parameter is determined using the Arrenius equation⁴⁷. The damage rate can be determined by

$$\mathcal{D} = F_f \int_0^t e^{-\frac{E_a}{RT}} d\Box \tag{55}$$

where R is the universal gas constant, E_a is the energy of activation for the denaturation reaction, and F_f is the frequency factor⁴⁸.

The computed data and interpretation

This section examines thermal transmission and the relations between mechanical forces and temperature for skin tissue exposed to varying heat loading. The study makes use of the generalized fourth-order MGT thermoelastic model. The way that heat loading is supplied to the skin tissue's outer surface is interpreted as

$$h(y, t) = \theta_0 U(t_s - t) U(L_y - y) \cos\left(\frac{\pi}{2t_s}t\right), \tag{56}$$

where t_s is the thermal loading exposure duration, $\theta_0 > 0$ is the thermal loading strength, and U(.) is the unit step function.

The following numerical computations and comparisons will make use of the biological and thermal properties of skin tissue shown in Table 1^{39,48}.

Figures 2, 3, 4, 5, 6, 7, 8, 9, 10, 11 and 12 display the numerical outcomes for different locations throughout the x-axis and the thermal damage along the t-time. The study will assess how specific influential parameters, such as relaxation (delay) time, model order, and duration of heat loading exposure, interact with skin tissues. The figures' meticulous examination, sensitivities of parameters, and linkages to prior research will offer a thorough assessment of the fourth-order MGT thermoelastic model and its capacity to faithfully depict skin tissue's mechanical and thermal reactions exposed to the varying thermal loading.

The primary objective of improving bio-thermal models is to resolve the intrinsic conflict in the Pennes model for bio-thermal transmission, which makes the assumption that thermal waves move at infinite rates. Therefore, a comparison of bioheat thermoelastic models will be covered in this initial scenario. The fourth-order Moore–Gibson–Thompson bio-thermal transfer model (4MGT-PH), the third-order Moore–Gibson–Thompson bio-thermal transmission model (3MGT-PH), and the conventional Pennes bio-thermal transmission parabolic model (P-PH) are the three different biothermal models used in this study to compare the locations of different physical variables inside skin tissues.

The physical field variables that were investigated at different distances x across several thermal models are shown in Figs. 1, 2, 3, 4 and 5. The parameters y=0.03 m, $t_s=20$ s, and t=10 sare employed in numerical computations. The findings show a distinct difference between the 4MGT-PH, 3MGT-PH, and P-PH models' outputs. The various field distributions are significantly influenced by the model's order as well as the thermal coefficient τ _h.

Figure 2 shows the thermodynamic temperature T changes as a function of distance x for the three thermoelasticity models. The graph shows that the P-PH model has higher temperature values than the 4MGT-PH and 3MGT-PH models. Interestingly, heat waves move more slowly in the 4MGT-PH and 3MGT-PH models than in the P-PH version. This discrepancy is caused by the thermal relaxation periods (τ_h) that are present in both updated models. This finding further supports the accuracy of the results generated by the suggested model, which offers a more accurate representation of thermal waves propagating through skin tissues.

In addition to the previously reported results, the 4MGT-PH model outperforms the P-PH and 3MGT-PH models in terms of prediction abilities because of incorporating additional higher-order terms. The enhanced efficiency of the 4MGT-PH model is ascribed to its capability to disperse heat more quickly and efficiently. Additionally, the behavior of the 3MGT-PH and 4MGT-PH models is rather similar. This result is in line with

Parameter	Value	Parameter	Value
λ_t	$8.27 \times 10^8 \text{ kg/ (m s}^2)$	μ_t	$3.446 \times 10^7 \text{ kg/ (m s}^2)$
α_t	$1 \times 10^{-4} 1/\mathrm{K}$	\mathcal{K}_t	0.628 J/ (s m K)
\mathcal{K}_t^*	$0.0628 \text{ J/} \left(\text{s}^2 \text{ m K}\right)$	c_t	4187 J/ (kg K)
S_{im}	$1.19 \times 10^{3} \mathrm{W/m^{3}}$	c_m	0.1 1/K
ρ_b	$1060 \mathrm{kg/m^3}$	c_b	3860 J/ (kg K)
w_b	0.00187 1/s	θ_{0}	5 K
F_f	$3.1 \times 10^{98} 1/s$	E_a	$6.28 \times 10^5 \text{ J/mol}$
R	8.313 J/ (mol K)	ρ_t	$1000 \mathrm{kg/m^3}$
L_x	0.03 m	L_y	0.03 m
T_b	310 K	w	10 1/m
f	15 1/s	τ_h	8 s

Table 1. Thermophysical parameters of skin tissue.

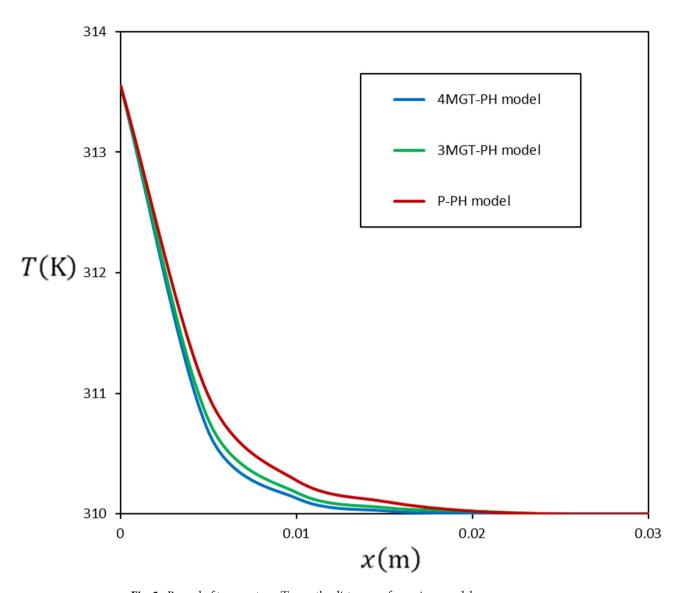


Fig. 2. Record of temperature $\,T\,$ over the distance $\,x\,$ for various models.

previous research^{49,50}, which demonstrates that the inclusion of a relaxation period lowers the thermal wave properties characteristics and produces behaviors akin to Fourier diffusion.

The issue of thermal waves is resolved by generalized models, which correctly predict that thermal signals move at bounded speeds. When approximating heat conduction in temperature fields that change dynamically, these models are especially helpful. The suggested MGT generalized thermoelasticity model overcomes the heat wave problem by offering a more realistic representation of thermal conductivity and guaranteeing the precision of the numerical results.

The displacements (u_x, u_y) and stresses $(\sigma_{xx}, \sigma_{yy})$ change with distance (x), as shown in Figs. 3, 4, 5 and 6. The skin tissue reacts to temperature changes more slowly when thermal relaxation periods (τ_h) are introduced. The time evolution of the skin tissue's stress and deformation is significantly impacted by this delay. Compared to traditional models, the MGT models' displacements (u_x, u_y) show a delayed heat response, leading to smoother and more dispersed deformation. Furthermore, the MGT models reveal fewer steep gradients in the thermal stress $(\sigma_{xx}, \sigma_{yy})$.

The fourth-order model provides additional enhancements by adding more higher-order terms, as can be seen when comparing the third-order and fourth-order MGT models. This development results in more accurate stress and deformation predictions. A more progressive thermal and mechanical response is also shown by the smoother distributions of all physical variables produced by the fourth-order MGT model. The study's findings support previous research⁵¹⁻⁵³, indicating the precision and dependability of the numerical method used.

The importance of the duration of heat loading exposure in skin tissues is covered in the second scenario. The thermal loading exposure period (t_s) plays a major role in determining the rate and velocity at which heat is injected into the skin tissue. Both the transient heat transmission process and the thermomechanical reaction of the skin tissue are directly impacted by this characteristic. By looking at numerical data related to significant

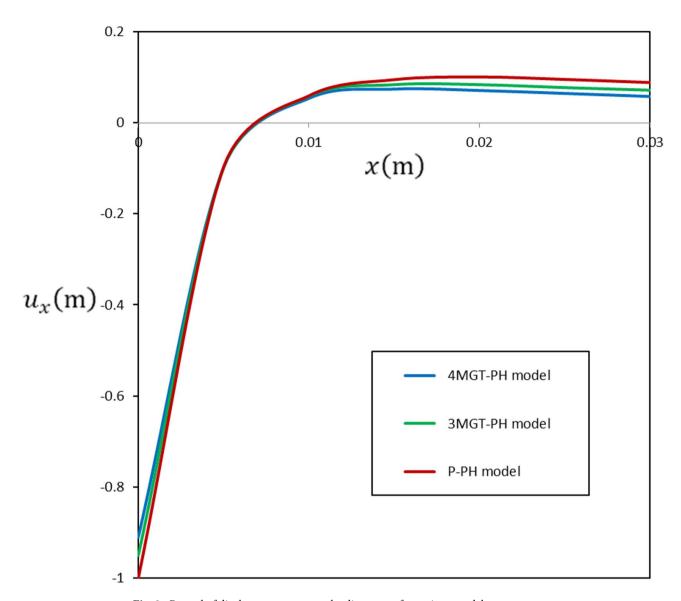


Fig. 3. Record of displacement u_x over the distance x for various models.

variables like temperature, displacement, and stress, one can gain a comprehensive understanding of how rapid heating affects skin tissue.

Figures 7, 8, 9, 10 and 11 show how t_s affects the temperature profile (T), displacements (u_x, u_y) , and thermal stress components $(\sigma_{xx}, \sigma_{yy})$. To investigate the effects of t_s , numerical simulations were performed for three different thermal loading exposure lengths $(t_s = 20 \text{ s}, 40 \text{ s}, \text{and } 60 \text{ s})$. The parameters y = 0.01 m and t = 10 s are used simultaneously in numerical computations. The discussion that follows provides a detailed analysis of the results.

The temperature profile investigation indicates that a more pronounced and localized temperature increase inside the skin tissue occurs with a longer thermal loading exposure length (t_s) (see Fig. 7). Increased temperature gradients and peak temperatures result in a sharp thermal response. Conversely, a shorter t_s promotes a gradual and consistent increase in temperature, lowering peak temperatures and producing smoother gradients over the skin tissue.

A longer t_s , which is linked to a higher rate of heat deposition, physically produces faster thermal responses. The concentrated heating creates steeper thermal gradients, increasing the likelihood of temperature overshoots and short-term variations. Conversely, a shorter t_s allows the temperature to spread more evenly across the skin tissue and reduces the rate of heat deposition. This lowers peak temperatures and prolongs the thermal impacts of the skin tissue.

Longer t_s results in concentrated and rapid thermal expansions and contractions, which create abrupt displacement shifts when examining displacements (u_x , u_y), as seen in Figs. 8 and 9. Conversely, a shorter t_s results in displacement fields that are more uniform and smooth, with slower transient deformations that progressively get closer to equilibrium. According to the scientific understanding of this phenomena, rapid temperature changes together with a longer t_s amplifying thermal expansion rates lead to significant transient

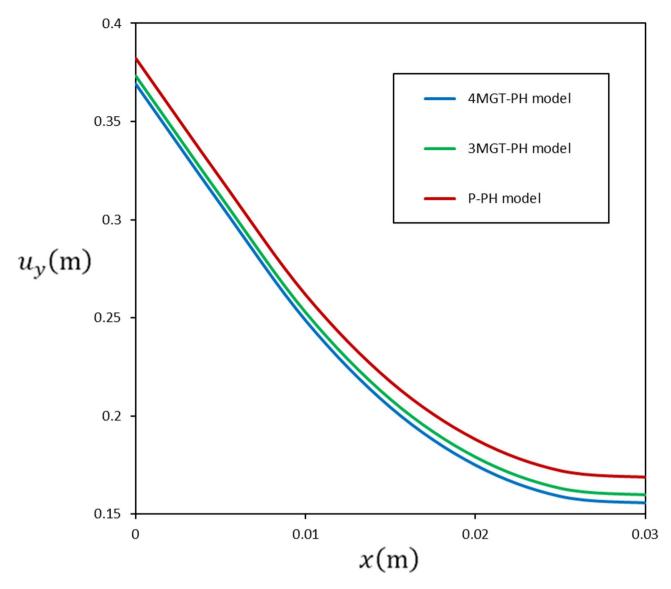


Fig. 4. Record of displacement u_y over the distance x for various models.

deformations concentrated near the heat source. Conversely, a slower temperature rise is indicated by a shorter t_s , which slows down the rate of thermal expansion and produces smoother displacement reactions and more gradual deformation.

Figures 10 and 11 demonstrate that as t_s increases, transient stresses (σ_{xx} , σ_{yy}) increase due to rapid temperature gradients and uneven expansion. A shorter t_s efficiently reduces the intensity of stresses (σ_{xx} , σ_{yy}), leading to more gradual stress response and smaller stress gradients.

The behaviors of thermomechanical coupling and heat transfer dynamics exhibit notable variations under different thermal loading exposure durations (t_s). This variation highlights how skin tissue reactions rely on the duration of thermal loading exposure by demonstrating that shorter durations encourage a more stable and consistent thermal response, whereas longer durations may result in more rapid alterations. Understanding these differences is crucial because they directly affect performance in real-world situations. By varying the duration of heat loading exposure, physicians and researchers can improve the performance skin tissues in a range of applications, including thermal therapy in biological tissues.

The extent of heat damage is covered in the final scenario. Extended exposure to extremely high temperatures can harm the cellular and structural components of the skin by thermally damaging the epidermal tissues. Thermal injury can occur via blistering from hot liquids, connection to heated bodies, fire damage, and even medical operations. The degree of thermal damage is dependent on the duration and intensity of temperature exposure, as well as the inherent susceptibility of the individual.

The assessment of skin tissue burns is essential in the field of medicine. Current studies have shown that heat damage starts when the temperature of the base stratum reaches 44 °C and increases by 0.53 °C at that point^{54–56}. The thermal damage to skin tissue following exposure to a heating load for t_s =20 seconds at x = 0.01 m and y = 0.01 m is calculated in this work using the composite trapezoidal rule. The findings in Fig. 12 clearly

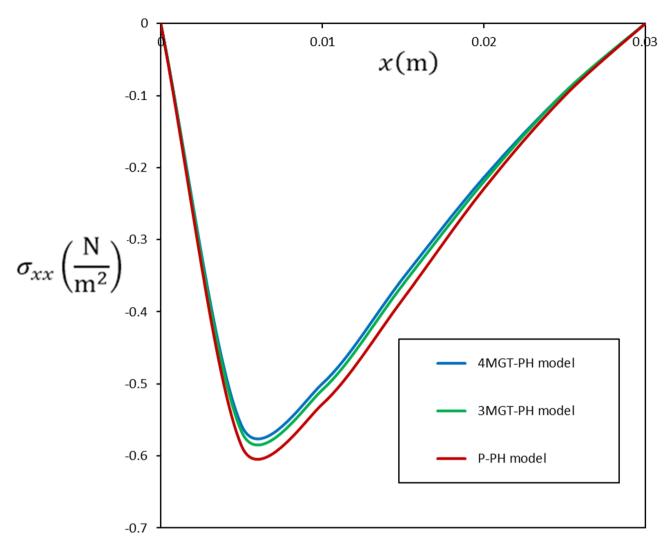


Fig. 5. Record of stress σ_{xx} over the distance x for various models.

demonstrate a statistically significant variation in the anticipated burn times between the bio-thermal models. Specifically, when applied to varying heating loads, the 4MGT-PH model exhibits lower burn periods than the P-PH model. Consequently, when the Pennes model is unable to adequately capture thermal transmission dynamics, the 4MGT-PH and 3MGT-PH models provide more precise forecasts in high power and rapid heating scenarios.

Conclusion

By introducing a novel Moore–Gibson–Thomson (MGT) thermoelasticity model, this study sought to advance the subject of thermoelasticity. This novel model provides a more thorough framework for studying thermoelastic events by combining the Moore–Gibson–Thomson equation with the relaxation period of the heat flux vector and higher-order derivatives. The thermo-elastic transient response of two-dimensional skin tissues subjected to varying thermal loads was examined, with thermophysical fields graphically displayed, in order to validate the model. The study examined thermal damage, the effect of the duration of heat loading exposure on the behavior of thermophysical fields, and a detailed comparison of classical and non-classical thermoelasticity models.

The following were some of the key findings:

- The inclusion of relaxation times in the heat conduction equation significantly improves the convergence in solutions for the 4MGT-PH and 3MGT-PH models. The accuracy of the model has improved by better capturing the non-local effects related to thermal and mechanical processes.
- Higher-order derivatives and relaxation periods reduce the propagation of mechanical and thermal waves within biological tissues, according to experimental data. The complex interaction between heat dissipation and mechanical responsiveness over time is suggested by this attenuation.
- Compared to temperature profiles produced by other models, the 4MGT-PH model exhibits a more pronounced and rapid decline in temperature as one moves farther away from the heat source. This implies that

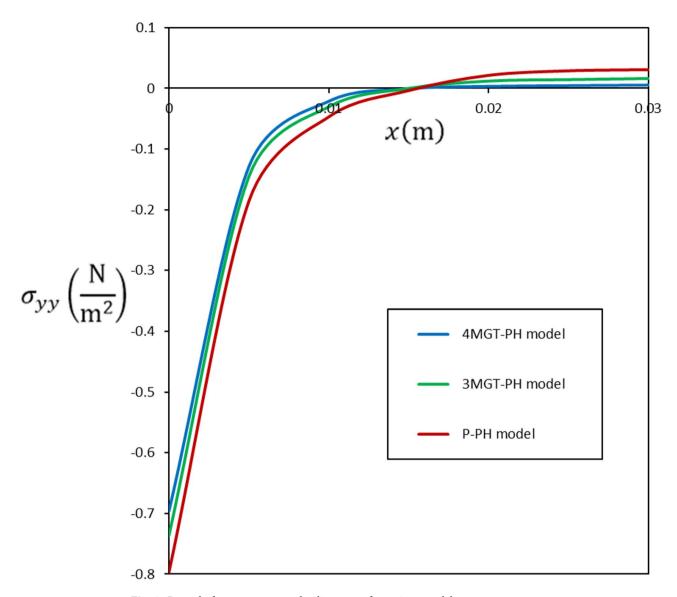


Fig. 6. Record of stress σ_{yy} over the distance x for various models.

the 4MGT-PH model more fully and accurately captures the underlying thermal phenomena within the skin tissue.

- More precisely, a longer heat loading exposure time leads to a faster and more concentrated temperature
 increase, which in turn produces higher peak temperatures, more pronounced temperature gradients, and
 stronger transitory effects. Conversely, a shorter heat loading exposure time results in slower, smoother thermal responses, which lessen excessive thermal and mechanical fluctuations.
- The Pennes thermal transmission model (P-PH) and the 3MGT-PH model produce considerably different estimates of temperature and thermal damage than the current bioheat model (4MGT-PH).
- The duration of the burn differs for each of the three heat transfer models when skin tissue is exposed to varying heating loads.
- This modeling technique may eventually improve the precision of heat modeling in biological fields, such as
 tissue diagnosis and the development of heat therapeutics, by better accounting for the intricate and dynamic
 structure of thermal transmission in skin tissue.

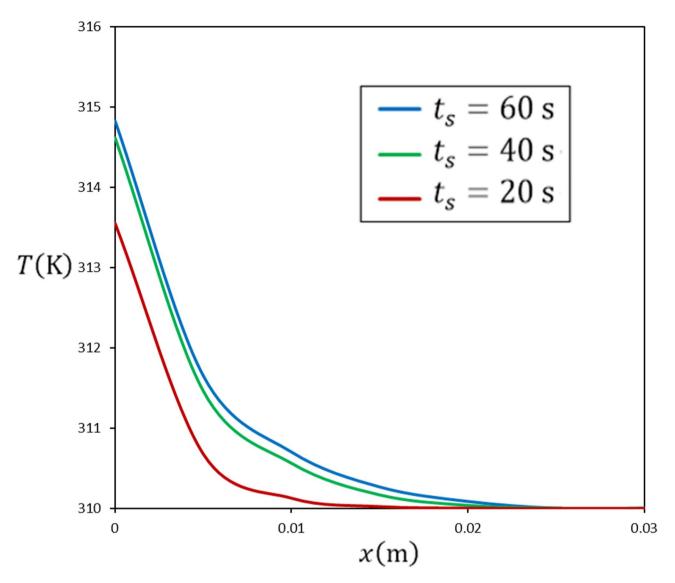


Fig. 7. The impact of the heat loading exposure duration t_s over temperature T.

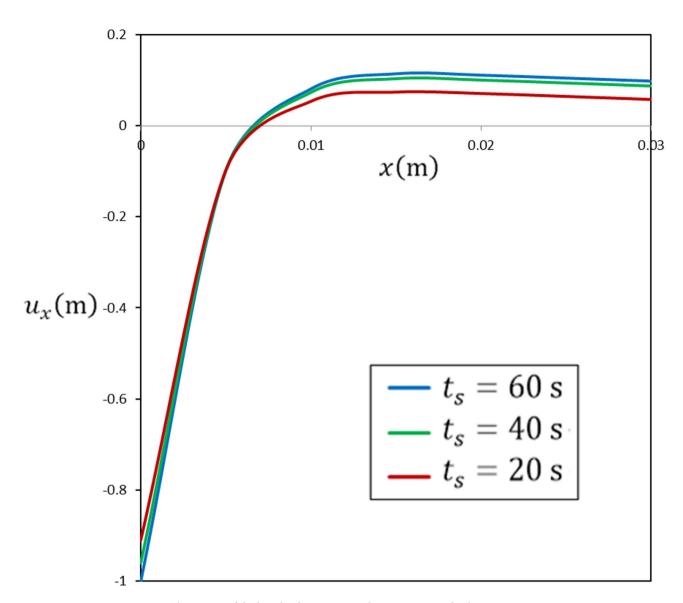


Fig. 8. The impact of the heat loading exposure duration t_s over displacement u_x .

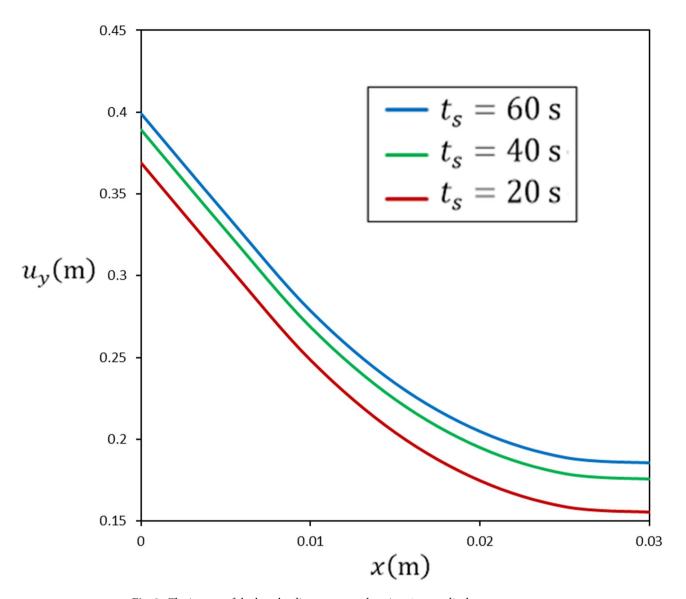


Fig. 9. The impact of the heat loading exposure duration t_s over displacement u_y .

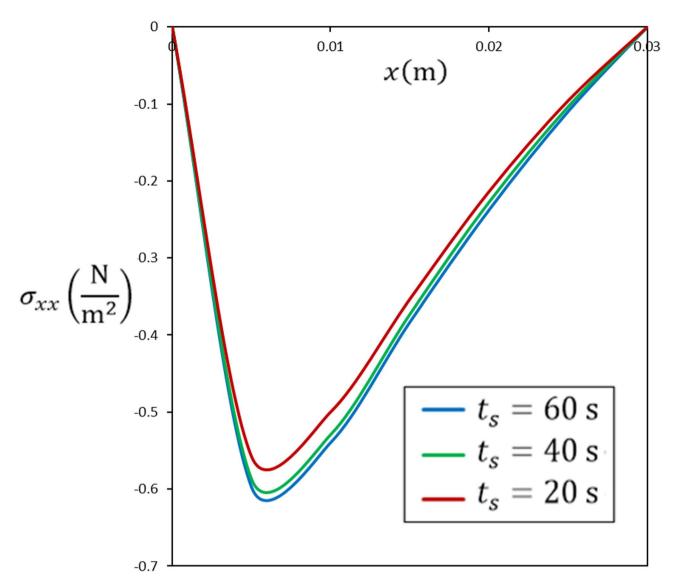


Fig. 10. The impact of the heat loading exposure duration t_s over stress σ_{xx} .

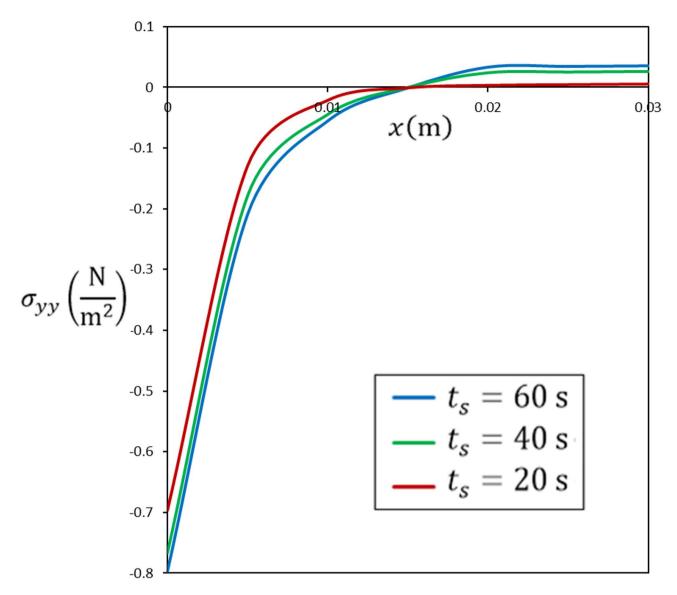


Fig. 11. The impact of the heat loading exposure duration t_s over stress σ_{yy} .

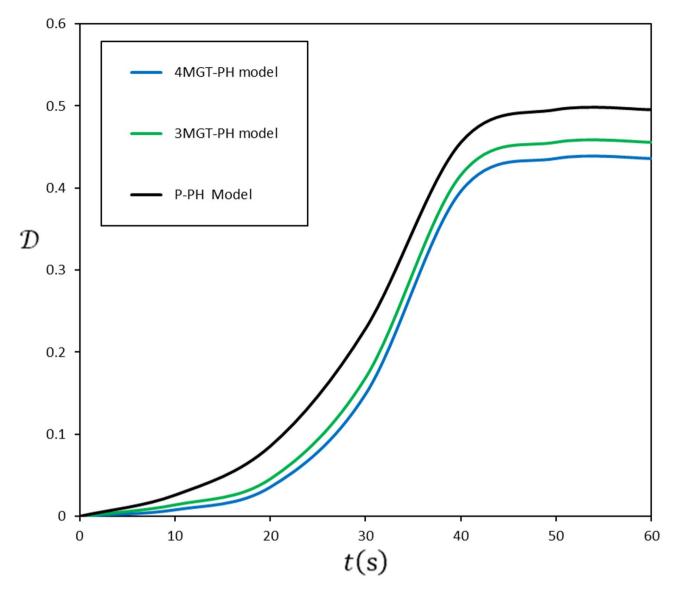


Fig. 12. Comparison between the models for the evolution of thermal damage \mathcal{D} .

Data availability

All available data are present in the manuscript. All data and models generated or used during the study appear in the submitted article.

Received: 3 February 2025; Accepted: 8 May 2025

Published online: 16 May 2025

References

- 1. Kumar, D. & Rai, K. N. A study on thermal damage during hyperthermia treatment based on DPL model for multilayer tissues using finite element legendre wavelet Galerkin approach. *J. Therm. Biol.* **62**, 170–180 (2016).
- 2. Huang, X., Jain, P. K., El-Sayed, I. H. & El-Sayed, M. A. Plasmonic photothermal therapy (PPTT) using gold nanoparticles. *Lasers Med. Sci.* 23, 217–228 (2008).
- 3. Pennes, H. H. Analysis of tissue and arterial blood temperatures in the resting human forearm. J. Appl. Physiol. 1 (2), 93–122 (1948).
- 4. Tung, M. M., Trujillo, M., Molina, J. L., Rivera, M. J. & Berjano, E. J. Modeling the heating of biological tissue based on the hyperbolic heat transfer equation. *Math. Comput. Modell.* **50** (5–6), 665–672 (2009).
- Tunc, M., Camdali, U., Parmaksizoğlu, C. & Cikrikci, S. The bio-heat transfer equation and its applications in hyperthermia treatments. Eng. Comput. 23 (4), 451–463 (2006).
- 6. Andreozzi, A., Brunese, L., Iasiello, M., Tucci, C. & Vanoli, G. P. Modeling heat transfer in tumors: a review of thermal therapies. Ann. Biomed. Eng. 47, 676–693 (2019).
- 7. Fu, Z. J., Xi, Q., Ling, L. & Cao, C. Y. Numerical investigation on the effect of tumor on the thermal behavior inside the skin tissue. Int. J. Heat. Mass. Transf. 108, 1154–1163 (2017).
- 8. Singh, S. & Repaka, R. Effect of different breast density compositions on thermal damage of breast tumor during radiofrequency ablation. *Appl. Therm. Eng.* 125, 443–451 (2017).

- 9. Zhang, Z. W., Wang, H. U. I. & Oin, O. H. Method of fundamental solutions for nonlinear skin bioheat model, I. Mech. Med. Biol. 14 (04), 1450060 (2014).
- 10. Mitra, K., Kumar, S., Vedevarz, A. & Moallemi, M. K. Experimental evidence of hyperbolic heat conduction in processed meat. J. Heat. Transf. 117 (3), 568-573 (1995).
- 11. Hooshmand, P., Moradi, A. & Khezry, B. Bioheat transfer analysis of biological tissues induced by laser irradiation. Int. J. Therm. Sci. 90, 214-223 (2015).
- 12. Cattaneo, C. Sur Une forme de L' equation de La Chaleur elinant Le paradoxe D' Une propagation instantance. C R Acad. Sci. III. 247, 431-432 (1958).
- 13. Vernotte, M. P. Les paradoxes de La theorie continue de l'equation de La Chaleur. C R Acad. Sci. III. 246, 3154-3155 (1958).
- 14. Liu, J., Chen, X. & Xu, L. X. New thermal wave aspects on burn evaluation of skin subjected to instantaneous heating. IEEE Trans. Biomed. Eng. 46 (4), 420-428 (1999).
- 15. Tzou, D. Y. A unified field approach for heat conduction from macro- to micro-scales. J. Heat. Transf. 117 (1), 8-16 (1995).
- 16. Du, B., Xu, G., Xue, D. & Wang, J. Fractional thermal wave bio-heat equation based analysis for living biological tissue with non-Fourier neumann boundary condition in laser pulse heating. Optik 247, 167811 (2021).
- 17. Abouelregal, A. E., Alharb, R. A., Yaylacı, M., Mohamed., B. O. & Megahid, S. F. Analysis of temperature changes in living tissue using the modified fractional thermal conduction model under laser heat flux on the skin surface. Continuum Mech. Thermodyn. https://doi.org/10.1007/s00161-024-01343-y (2024).
- 18. Sukumar, S. & Kar, S. P. Numerical investigation of a novel method of laser assisted cryopreservation of biological tissue considering non-Fourier heat conduction. Lasers Med. Sci. 38 (1), 98 (2023).
- 19. Kublanov, V. S., Borisov, V. I. & Babich, M. V. Simulation the distribution of thermodynamic temperatures and microwave radiation of the human head. Comput. Methods Programs Biomed. 190, 105377 (2020).
- 20. Richards, N. et al. Evaluation of acoustic-thermal simulations of in vivo magnetic resonance guided focused ultrasound ablative therapy. Int. J. Hyperth. 41 (1), 2301489 (2024).
- 21. Megahid, S. F., Abouelregal, A. E. & Sedighi, H. M. Modified Moore-Gibson-Thompson Pennes' bioheat transfer model for a finite biological tissue subjected to harmonic thermal loading. Mech. Time Depend. Mater. https://doi.org/10.1007/s11043-023-09647-3
- 22. Lamba, N. K. Impact of memory-dependent response of a thermoelastic Thick solid cylinder. J. Appl. Comput. Mech. 9 (4), 1135– 1143 (2023)
- 23. Safaei, B., Onyibo, E. C. & Hurdoganoglu, D. Thermal buckling and bending analyses of carbon foam beams sandwiched by composite faces under axial compression. Facta Universitatis Ser.: Mech. Eng. 20 (3), 589-615 (2022).
- 24. Zenkour, A. M., Saeed, T. & Al-Raezah, A. A. Fractional order of refined Lord-Shulman model for a 1D thermoelastic response of skin tissue due to ramp-type heating. Arch. Appl. Mech. 94, 989-1003 (2024).
- 25. Lord, H. W. & Shulman, Y. A generalized dynamical theory of thermoelasticity. J. Mech. Phys. Solids. 15 (5), 299-309 (1967).
- 26. Green, A. E. & Lindsay, K. Thermoelast. J. Elast. 2 (1), 1-7 (1972).
- 27. Green, A. E. & Naghdi, P. Thermoelasticity without energy dissipation. J. Elast. 31 (3), 189-208 (1993).
- 28. Hu, Y., Zhang, X. Y. & Li, X. F. Thermoelastic analysis of biological tissue during hyperthermia treatment for moving laser heating using fractional dual-phase-lag bioheat conduction. Int. J. Therm. Sci. 182, 107806 (2022).
- Jangid, K., Singh, B. & Mukhopadhyay, S. Legendre wavelet collocation method for investigating thermo-mechanical responses on biological tissue during laser irradiation. Math. Comput. Simul. 219, 404-423 (2024).
- 30. Quintanilla, R. Moore-Gibson-Thompson thermoelasticity. Math. Mech. Solids. 24 (12), 4020-4031 (2019).
- 31. Quintanilla, R. Moore-Gibson-Thompson thermoelasticity with two temperatures. Appl. Eng. Sci. 1, 100006 (2020).
- 32. Green, A. E. & Naghdi, P. A re-examination of the basic postulates of thermomechanics. Proc. R Soc. Lond. Math. Phys. Sci. 432(1885), 171-194 (1991).
- Abouelregal, A. E., Megahid, S. F. & Sedighi, H. M. Thermal effects of electromagnetic radiation on the skin tissue by considering fourth-order MGT bioheat model. ZAMM-J Appl. Math. Mech. 104 (7), e202301085 (2024).
- 34. Megahid, S. F., Abouelregal, A. E., Ahmad, H., Fahmy, M. A. & Abu-Zinadah, H. A generalized Moore-Gibson-Thomson heat transfer model for the study of thermomagnetic responses in a solid half-space. Results Phys. 51, 106619 (2023).
- 35. Megahid, S. F., Abouelregal, A. E., Askar, S. S. & Marin, M. Study of thermoelectric responses of a conductive semi-solid surface to variable thermal shock in the context of the Moore-Gibson-Thompson thermoelasticity. Axioms 12 (7), 659 (2023).
- 36. Pellicer, M. & Quintanilla, R. Continuous dependence and convergence for Moore-Gibson-Thompson heat equation. Acta Mech. 234, 3241-3257 (2023).
- 37. Chen, W. & Takeda, H. Asymptotic behaviors for the Jordan-Moore-Gibson-Thompson equation in the viscous case. Nonlinear Anal. 234, 113316 (2023).
- 38. Ming, S., Fan, X., Ren, C. & Su, Y. Blow-up dynamic of solution to the semilinear Moore-Gibson-Thompson equation with memory terms. AIMS Math. 8 (2), 4630-4644 (2023).
- Ezzat, M. A. Bio-thermo-mechanics behavior in living viscoelastic tissue under the fractional dual-phase-lag theory. Arch. Appl. Mech. 91 (9), 3903-3919 (2021).
- 40. Shah, N. A., Almutairi, B., Vieru, D., Lee, B. & Chung, J. D. Bioheat transfer with thermal memory and moving thermal shocks. Fractal Fract. 7 (8), 629 (2023).
- 41. Hu, Y., Zhang, X. Y. & Li, X. F. Thermoelastic response of skin using time-fractional dual-phase-lag bioheat heat transfer equation. J. Therm. Stresses. 45 (7), 597–615 (2022).
- 42. Abbas, I. A., El-Bary, A. A. & Mohamed, A. O. Generalized thermomechanical interaction in two-dimensional skin tissue using eigenvalues approach. J. Therm. Biol. 119, 103777 (2024).
- Meena, B. S. & Kumar, S. Temperature response in skin tissue during hyperthermia based on three-phase-lag bioheat model using RBF Meshfree method. Numer. Heat. Transf. Part. A: Appl. 85 (2), 287-305 (2024).
- 44. Singh, J., Gupta, P. K. & Rai, K. N. Solution of fractional bioheat equations by finite difference method and HPM. Math. Comput. Modell. 54 (9-10), 2316-2325 (2011).
- 45. Arpaci, V. C. Conduction Heat Transfer (Addisson-Wesley, 1966).
- 46. Moritz, A. R. & Henriques, F. C. Study of thermal injuries II. The relative importance of time and source temperature in the causation of cutaneous burns. Am. J. Pathol. 23 (5), 695-720 (1947).
- 47. Welch, A. J. The thermal response of laser irradiated tissue. IEEE J. Quantum Electron. 20 (12), 1471-1481 (1984).
- 48. Hobiny, A. D. & Abbas, I. A. Nonlinear analysis of dual-phase lag bio-heat model in living tissues induced by laser irradiation. J. Therm. Stresses. 43 (4), 503-511 (2020).
- Ezzat, M. A., El-Karamany, A. S. & El-Bary, A. A. Thermoelectric viscoelastic materials with memory dependent derivative. Smart Struct. Syst. 19, 539-551 (2017).
- 50. Ziaei, P. H., Moosavi, H. & Moradi, A. Analysis of the dual phase lag bio-heat transfer equation with constant and time-dependent heat flux conditions on skin surface. Therm. Sci. 20 (5), 1457-1472 (2016).
- 51. Deng, Z. S. & Liu, J. Analytical study on bioheat transfer problems with Spatial or transient heating on skin surface or inside biological bodies. J. Biomech. Eng. 124 (6), 638-649 (2002). Kim, H. J. et al. Laser-tissue interaction simulation considering skin-specific data to predict photothermal damage lesions during
- laser irradiation. J. Comput. Des. Eng. 10 (3), 947-958 (2023).
- 53. Liu, K. C. Thermal propagation analysis for living tissue with surface heating. Int. J. Therm. Sci. 47 (5), 507–513 (2008).

- 54. Ahmadikia, H., Fazlali, R. & Moradi, A. Analytical solution of the parabolic and hyperbolic heat transfer equations with constant and transient heat flux conditions on skin tissue. *Int. Commun. Heat. Mass. Transf.* 39 (1), 121–130 (2012).
- 55. Xu, F, Lu, T. J. & Seffen, K. A. Biothermomechanical behavior of skin tissue. Acta Mech. Sin. 24 (1), 1-23 (2008).
- 56. Xu, F., Seffen, K. A. & Lu, T. J. Non-Fourier analysis of skin biothermomechanics. Int. J. Heat. Mass. Transf. 51 (9–10), 2237–2259 (2008).

Author contributions

Sami F. Megahid did all the contributions.

Funding

Open access funding provided by The Science, Technology & Innovation Funding Authority (STDF) in cooperation with The Egyptian Knowledge Bank (EKB).

Competing interests

The authors declare no competing interests.

Ethical approval and consent to participate

The submitted work is original, and the manuscript has not been submitted to another journal for simultaneous consideration.

Consent for publication

The authors declare that they consent to publish the article.

Additional information

Correspondence and requests for materials should be addressed to S.F.M.

Reprints and permissions information is available at www.nature.com/reprints.

Publisher's note Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

Open Access This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons licence, and indicate if changes were made. The images or other third party material in this article are included in the article's Creative Commons licence, unless indicated otherwise in a credit line to the material. If material is not included in the article's Creative Commons licence and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this licence, visit https://creativecommons.org/licenses/by/4.0/.

© The Author(s) 2025