

A Noncontact Method for Measuring Thermal Conductivity and Thermal Diffusivity of Anisotropic Materials

M. Fujii,^{1,2} S. C. Park,¹ T. Tomimura,¹ and X. Zhang¹

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A noncontact method for measuring the thermal conductivity and thermal diffusivity of anisotropic materials is proposed. This method is based on the fact that the surface temperature variation with time depends on the thermal properties of the material when its surface is heated locally. The three-dimensional transient heat conduction equation in the material is solved numerically. The dimensionless average surface temperature variations are obtained along each principal axis; that is, the x and y axes. The relation between the dimensionless temperature and the Fourier number is expressed by a polynomial equation and used as a master plot, which is a basic relation to be compared with measured temperature variation. In the experiments, the material surface is heated with a laser beam and the surface temperature profiles are measured by an infrared thermometer. The measured temperature variations with time are compared with the master plots to yield the thermal conductivity λ_x and thermal diffusivity α_x in the x direction and the thermal conductivity ratio $E_{xy}(=\lambda_y/\lambda_x)$ simultaneously. To confirm the applicability and the accuracy of the present method, measurements were performed on multilayered kent-paper, vinyl chloride, and polyethylene resin film, whose thermal properties are known. From numerical simulations, it is found that the present method can measure the thermophysical properties λ_x , α_x and E_{xy} within errors of ± 6 , ± 22 , and $\pm 5\%$, respectively, when the measuring errors of the peak heat flux, the heating radius, and the surface temperature rise are assumed to be within ± 2 , $\pm 3^\circ$, and ± 0.2 K, respectively. This method could be applied to the measurement of thermophysical properties of biological materials.

KEY WORDS: anisotropic material; noncontact measurement method; thermal conductivity; thermal diffusivity.

¹ Institute of Advanced Material Study, Kyushu University, Kasuga, Fukuoka 816, Japan.

² To whom correspondence should be addressed.

1. INTRODUCTION

As anisotropic materials whose thermal conductivity and thermal diffusivity depend on directions, there are insulation materials, multilayered materials, biological materials, etc. In the fields of medical engineering and biotechnology, it is particularly important to perform an *in vivo* measurement of thermophysical properties of biological materials. It is very difficult to obtain accurate thermophysical properties by the conventional *invasive* method, because a living body reacts sensitively on such foreign substances as probes.

To overcome this problem, the authors have proposed a noncontact measurement method using a laser and an infrared thermometer, and established the simultaneous measuring technique of thermal conductivity and thermal diffusivity of isotropic materials and, further, examined errors in the measurements [1, 2]. The method is based on the fact that the surface temperature variation with time after step heating depends on the thermal properties of the material, therefore, by combining the laser heating with the surface temperature measurement by an infrared thermometer, a noncontact (i.e., *noninvasive*) measurement of thermophysical properties becomes possible.

In this paper, the method proposed for isotropic materials is extended to measure anisotropic materials by performing three-dimensional transient numerical simulation and by measuring the two-dimensional surface temperature distributions. The thermophysical properties of anisotropic materials such as multilayered kent-paper, vinyl chloride, and polyethylene resin film are measured. As for the kent-paper, the measured results are compared to those obtained by Takegoshi et al. [3] using the transient hot-wire method. Further, the validity of the present measurement method is confirmed, and the effects of the measurement errors of the peak heat flux, the heating radius, and the surface temperature rise on the accuracy of the thermophysical property evaluation are investigated quantitatively.

2. NUMERICAL ANALYSIS AND MEASURING PRINCIPLE

2.1. Physical Model and Basic Equations

In the present measurement method, the thermal conductivity and thermal diffusivity of a material are estimated simultaneously by fitting the measured surface temperature variation to that obtained numerically. In this section, a numerical analysis is described which clarifies the relation between the surface temperature variation and its thermophysical properties.

Figure 1 shows the physical model (two-layered model) and the coordinate system. Corresponding to the test material shown in Section 3, a black paint layer coated on its surface is considered. The rectangular prism ($2y_s$ long, $2x_s$ wide, and z_s thick) of an anisotropic material is placed in still air of temperature T_∞ . The thermal conductivities and thermal diffusivities in the x , y , and z directions are λ_x , λ_y , and λ_z and α_x , α_y , and α_z , respectively. The thin black paint layer of thickness z_b has the thermal conductivity λ_b , the thermal diffusivity α_b , and the emissivity ϵ . The black paint surface is locally heated with a laser beam. Taking account of the Gaussian distribution of the beam intensity, the heat flux distribution $q(r)$ is given as

$$q(r) = q_p \exp\{-2.3(r/r^*)^2\} \tag{1}$$

where r^* is the heating radius, which is defined as the radius where the heat flux becomes 10% of the peak value at the beam center $q_p = q(0)$. Most part of the heat input is conducted through the paint layer and the material, and the rest is transferred back to the ambient by free convection (q_c) and radiation (q_r). The basic equations of three-dimensional transient

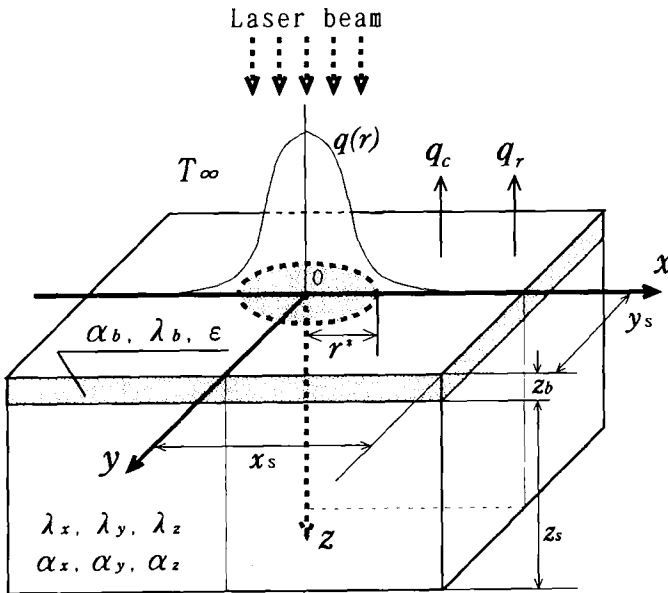


Fig. 1. Physical model and the coordinate system.

heat conduction for the black paint layer (subscript b) and the anisotropic material (subscript s) are written in dimensionless form as

$$\frac{\partial \Theta_b}{\partial \text{Fo}} = P_{sv} \left(\frac{\partial^2 \Theta_b}{\partial X^2} + \frac{\partial^2 \Theta_b}{\partial Y^2} + \frac{\partial^2 \Theta_b}{\partial Z^2} \right) \quad (2)$$

$$\frac{\partial \Theta_s}{\partial \text{Fo}} = \frac{\partial^2 \Theta_s}{\partial X^2} + E_{sv} \frac{\partial^2 \Theta_s}{\partial Y^2} + E_{sz} \frac{\partial^2 \Theta_s}{\partial Z^2} \quad (3)$$

The initial conditions are given by

$\text{Fo} = 0$:

$$\begin{aligned} 0 \leq X \leq X_s, \quad 0 \leq Y \leq Y_s, \quad 0 \leq Z \leq Z_b; \quad \Theta_b = 0 \\ 0 \leq X \leq X_s, \quad 0 \leq Y \leq Y_s, \quad Z_b \leq Z \leq Z_b + Z_s; \quad \Theta_s = 0 \end{aligned} \quad (4)$$

and the boundary conditions are given by

$\text{Fo} > 0$:

$$\begin{aligned} 0 < X < X_s, \quad 0 \leq Y \leq Y_s, \quad Z = 0; \\ -P_{sv} \frac{\partial \Theta_b}{\partial Z} &= \exp(-2.3R^2) - \text{Bi} \Theta_w - Rcc \Theta_w \\ 0 < X \leq X_s, \quad 0 < Y \leq Y_s, \quad Z = Z_b + Z_s; \quad \frac{\partial \Theta_s}{\partial Z} &= 0 \\ 0 < X \leq X_s, \quad Y = 0, Y_s, \quad 0 < Z \leq Z_b; \quad \frac{\partial \Theta_b}{\partial Y} &= 0 \\ 0 < X \leq X_s, \quad Y = 0, Y_s, \quad Z_b < Z \leq Z_b + Z_s; \quad \frac{\partial \Theta_s}{\partial Y} &= 0 \\ X = 0, X_s, \quad 0 < Y \leq Y_s, \quad 0 < Z \leq Z_b; \quad \frac{\partial \Theta_b}{\partial X} &= 0 \\ X = 0, X_s, \quad 0 < Y \leq Y_s, \quad Z_b < Z \leq Z_b + Z_s; \quad \frac{\partial \Theta_s}{\partial X} &= 0 \\ 0 < X \leq X_s, \quad 0 < Y \leq Y_s, \quad Z = Z_b; \\ P_{sz} \frac{\partial \Theta_b}{\partial Z} &= \frac{\partial \Theta_s}{\partial Z}, \quad \Theta_b = \Theta_s \end{aligned} \quad (5)$$

The dimensionless variables are defined by

$$\begin{aligned} \Theta &= \frac{T - T_s}{q_p r^* / \lambda_x}, & \text{Fo} &= \frac{\alpha_x t}{r^{*2}}, & X &= \frac{x}{r^*}, & Y &= \frac{y}{r^*}, & Z &= \frac{z}{r^*}, & R &= \frac{r}{r^*} \\ P_{\lambda_x} &= \frac{\lambda_b}{\lambda_x}, & P_{\alpha_x} &= \frac{\alpha_b}{\alpha_x}, & P_{\lambda_z} &= \frac{\lambda_b}{\lambda_z}, & E_{xy} &= \frac{\lambda_y}{\lambda_x}, & E_{xz} &= \frac{\lambda_z}{\lambda_x} \\ \text{Bi} &= \frac{hr^*}{\lambda_x}, & Rc &= \frac{4T_s^3 \varepsilon \sigma r^*}{\lambda_x} \end{aligned} \tag{6}$$

In the above equations, Θ is the dimensionless temperature, Θ_w [$=\Theta_b(\text{Fo}, R, 0)$] the dimensionless surface temperature, T the absolute temperature, t the time after step heating, and Fo the corresponding Fourier number, X , Y , Z , and R are the dimensionless coordinates, P_{λ_x} and P_{λ_z} are the ratios of thermal conductivity of the black paint to those of the material in the x and z directions, respectively, P_{α_x} is the similar ratio of thermal diffusivities, E_{xy} , and E_{xz} are the ratios of thermal conductivities in the respective directions, and Bi is the Biot number, h the heat transfer coefficient, Rc the conduction-radiation parameter, and σ the Stefan-Boltzmann constant.

2.2. Parameters

The basic equations are solved numerically by the finite difference method under the previous initial and boundary conditions. In the present calculations, the ranges of the dimensionless parameters P_{α_x} , E_{xy} , and E_{xz} in Eqs. (2) and (3), and P_{λ_x} , P_{λ_z} , Z_b , Bi , and Rc in the initial and boundary conditions Eqs. (4) and (5), are chosen as

$$\begin{aligned} P_{\lambda_x} &= 1, 6, 15, & P_{\alpha_x} &= 1, 5, 10, 15, & P_{\lambda_z} &= P_{\lambda_x} \\ E_{xy} &= 0.1, 0.2, 0.5, 1, 2, 4, 6, & E_{xz} &= 1 \\ \text{Bi} &= 0.02 \sim 0.11, & Rc &= (0.66 \sim 4.0) \times 10^{-2}, & Z_b &= 3.5 \times 10^{-2} \end{aligned}$$

where the ranges of P_{α_x} and P_{λ_x} are determined by considering the values of $\lambda_b = 1.89 \text{ W} \cdot \text{m}^{-1} \cdot \text{K}^{-1}$ and $\alpha_b = 1.09 \times 10^{-6} \text{ m}^2 \cdot \text{s}^{-1}$ obtained from the previous measurements [4], and also those of the present test materials and the living tissues [5], as

$$\lambda_x = 0.1 \sim 0.6 \text{ W} \cdot \text{m}^{-1} \cdot \text{K}^{-1}, \quad \alpha_x = (0.1 \sim 0.5) \times 10^{-6} \text{ m}^2 \cdot \text{s}^{-1}$$

The values of Bi , Rc , and Z_b are determined from the relevant values of $h = 15 \text{ W} \cdot \text{m}^{-2} \cdot \text{K}^{-1}$, $r^* = 0.71 \times 10^{-3} \text{ m}$, $T_s = 293 \text{ K}$, $\varepsilon = 0.98$, and

$z_b = 25 \times 10^{-6}$ m. It should be noted that only the case of $E_{vz} = 1$ is treated in the present paper.

2.3. Measuring Principle

To establish a simple procedure for evaluating the thermal conductivity and thermal diffusivity from the numerical results, a unique relation between Θ_w and Fo as a master plot which is independent of the parameters Bi and Rc should be obtained. For this purpose, similar to the previous study [1, 2], the following dimensionless surface temperature,

$$\Theta_{wn} = \frac{T_w - T_\infty}{q_n r^* / \lambda_x} = \frac{\Delta T_w}{q_n r^* / \lambda_x} = \Theta_w \frac{q_p}{q_n} \tag{7}$$

is introduced by replacing the peak heat flux q_p included in Θ_w with an average net heat flux defined by

$$q_n = \frac{1}{\pi r^{*2}} \int_0^{r^*} [q_p \exp\{-2.3(r/r^*)^2\} - h(T_w - T_\infty) - \epsilon\sigma(T_w^4 - T_\infty^4)] 2\pi r dr \tag{8}$$

where T_w is the surface temperature of black paint. The redefined dimensionless surface temperature Θ_{wn} is confirmed to be almost independent of Bi and Rc for the ranges shown in the previous section.

When a material is anisotropic, the isotherms on the surface show not concentric circle but ellipsoid. In this case, the X and Y axes are chosen to coincide with the long and short axes of the ellipsoid, respectively. Then the following dimensionless parameters are introduced.

$$I_{th-X} = \frac{1}{\pi R^{*2}} \int_0^{R^*} \Theta_{wn-X} 2\pi R dR \tag{9}$$

$$I_{th-Y} = \frac{1}{\pi R^{*2}} \int_0^{R^*} \Theta_{wn-Y} 2\pi R dR \tag{10}$$

Equations (9) and (10) are the superficial average temperatures which would be defined under the assumption that the respective temperature distribution in the X or Y direction is the same as that for an isotropic material.

Figure 2 shows an example of time variation of I_{th-X} and I_{th-Y} . The relations also are found to be independent of Bi and Rc in the ranges of the present calculations. The relations, therefore, can be used as the master plots for the present measurement method. Since the larger fraction of heat

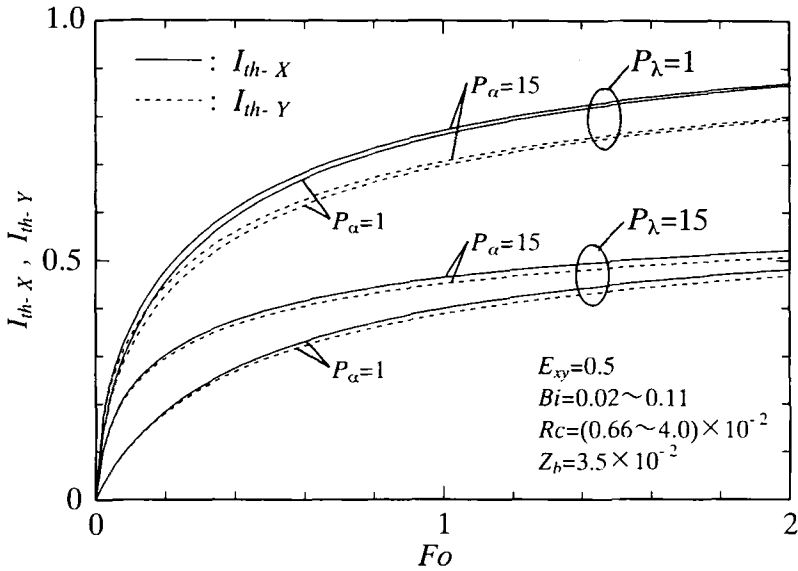


Fig. 2. Dimensionless mean surface temperature variations.

flows in the X direction for the case of $E_{xy}=0.5(\lambda_y=0.5\lambda_x)$ shown in Fig. 2, the average surface temperature I_{th-X} becomes high compared with I_{th-Y} . With increasing P_λ , however, the lateral heat flows through the black paint layer increases, and therefore, the difference between I_{th-X} and I_{th-Y} is reduced for $P_\lambda=15$.

The change of I_{th-Y} with time is approximated with the polynomial expression $I_{th-Y}=G(Fo)$. Here, the integration of the function $G(Fo)$ for $Fo=0 \sim Fo$ is defined as

$$A_{th-Y} = \int_0^{Fo} I_{th-Y} dFo = \int_0^{Fo} G(Fo) dFo \tag{11}$$

From the measurements, on the other hand, corresponding to Eqs. (9), (10), and (11), the following quantities are introduced from the measured surface temperature distributions along the x and y axes ΔT_{w-x} and ΔT_{w-y} , at time t .

$$i_{ex-x} = \frac{1}{\pi r^{*2}} \int_0^{r^*} \frac{\Delta T_{w-x}}{q_n} 2\pi r dr \tag{12}$$

$$i_{ex-y} = \frac{1}{\pi r^{*2}} \int_0^{r^*} \frac{\Delta T_{w-y}}{q_n} 2\pi r dr \tag{13}$$

Also, by integrating the polynomial approximation expression $i_{ex-y} = g(t)$ for $t = 0 \sim t$, the quantity

$$a_{ex-y} = \int_0^t i_{ex-y} dt \equiv \int_0^t g(t) dt \tag{14}$$

is defined.

The estimation method of the thermal conductivity and thermal diffusivity of anisotropic materials using Eqs. (9)–(14) is shown in Fig. 3. The process is as follows:

(1) By selecting a proper time after step heating t_0 , the values of i_{ex-x} and i_{ex-y} and its integration value a_{ex-y} at $t = t_0$ are calculated. The

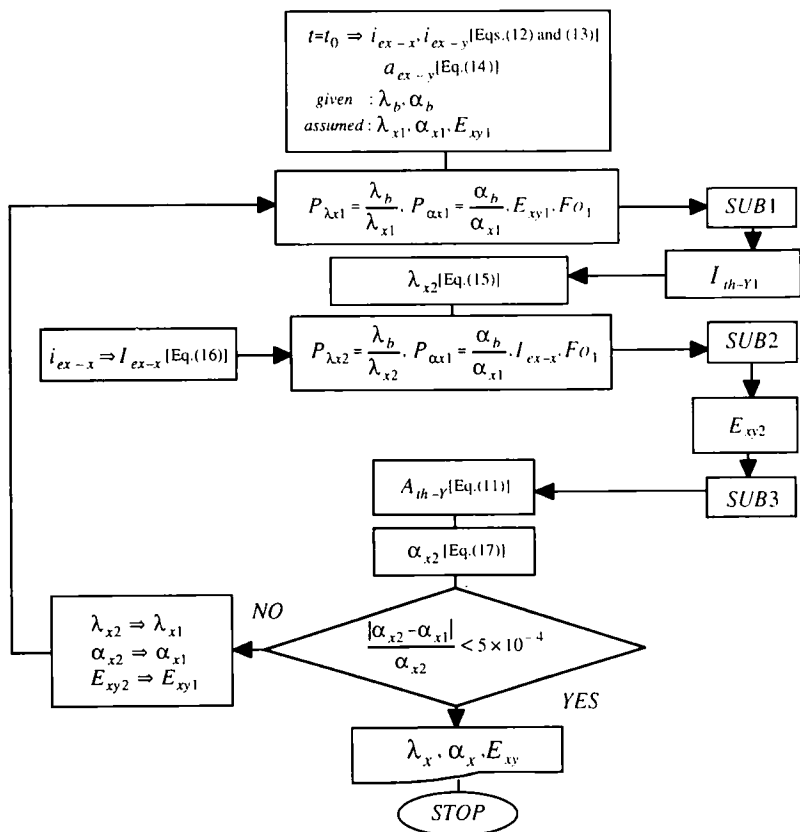


Fig. 3. Flowchart for evaluating thermal conductivity and thermal diffusivity of anisotropic materials.

thermal conductivity and thermal diffusivity in the x direction λ_{x1} , and α_{x1} and the ratio of thermal conductivities in the x and y direction E_{xy1} of the material are assumed.

(2) The thermal conductivity and thermal diffusivity ratios of the black paint to the material $P_{\lambda x1}$ and $P_{\alpha x1}$ are determined from the assumed values λ_{x1} and α_{x1} and the given thermophysical properties of the black paint layer λ_b and α_b . The Fourier number Fo_1 corresponding to α_{x1} and the time t_0 is also determined.

(3) Using the program SUB1 where the relations between I_{th-y} and Fo are given for each combination of the parameters P_{λ} , P_{α} , and E_{xy} , the dimensionless average surface temperature I_{th-y1} at $P_{\lambda x1}$, $P_{\alpha x1}$, E_{xy1} , and Fo_1 is obtained by interpolation.

(4) Thus the obtained I_{th-y1} and the measured value i_{cx-y} are substituted into the relation,

$$\lambda_{x2} = \frac{I_{th-y1} r^*}{i_{cx-y}} \tag{15}$$

to obtain the first approximate value of thermal conductivity λ_{x2} .

(5) Using λ_{x2} the measured value of i_{cx-x} is transformed into dimensionless form as

$$I_{cx-x} = \frac{i_{cx-x} \lambda_{x2}}{r^*} \tag{16}$$

(6) Similar to the third step, using the program SUB2 in which the relations between I_{th-x} and E_{xy} are given at the fixed values of $P_{\lambda x}$, $P_{\alpha x}$, and Fo , the first approximate value of the thermal conductivity ratio E_{xy2} is determined so that I_{th-x} coincides with I_{cx-x} .

(7) Through the program SUB3, the polynomial expression $I_{th-y} = G(Fo)$ for $P_{\lambda x2}$, $P_{\alpha x1}$, and E_{xy2} is obtained by interpolation. Then the integral A_{th-y} is calculated from Eq. (11) for $Fo = 0 \sim Fo$.

(8) The measured value a_{cx-y} and thus the obtained A_{th-y} are substituted into the relation,

$$\alpha_{x2} = \frac{r^{*3}}{a_{cx-y} \lambda_{x2}} A_{th-y} \tag{17}$$

to obtain the first approximate value of thermal diffusivity α_{x2} .

(9) Until the relative error in thermal diffusivity satisfies a given convergent criterion, the steps from 2 to 8 are repeated by replacing λ_{x1} ,

α_{x1} , and E_{xy1} with λ_{x2} , α_{x2} , and E_{xy2} . Finally, the thermal conductivity and thermal diffusivity in the y direction $\lambda_y (=E_{xy}\lambda_x)$, and $\alpha_y (=E_{xy}\alpha_x)$ are estimated from the converged values of λ_x , α_x , and E_{xy} .

3. MEASUREMENTS

A schematic diagram of the measuring system is shown in Fig. 4. The system consists of an argon laser (wavelength, 514.5 nm) (4) with a shutter (3), an infrared thermometer system (6–9), and a test material (1). The material surface is heated directly by the argon laser. The intensity distribution of the laser beam is confirmed to be Gaussian, and at the same time, the heating radius r^* is determined to be 0.71 mm by measuring the relative intensity through a 10- μm -diameter pin hole with a photodiode [3]. The laser output is adjusted by a power controller (5), and its values before and after a measurement are checked using a power meter (2) and a digital multimeter (10).

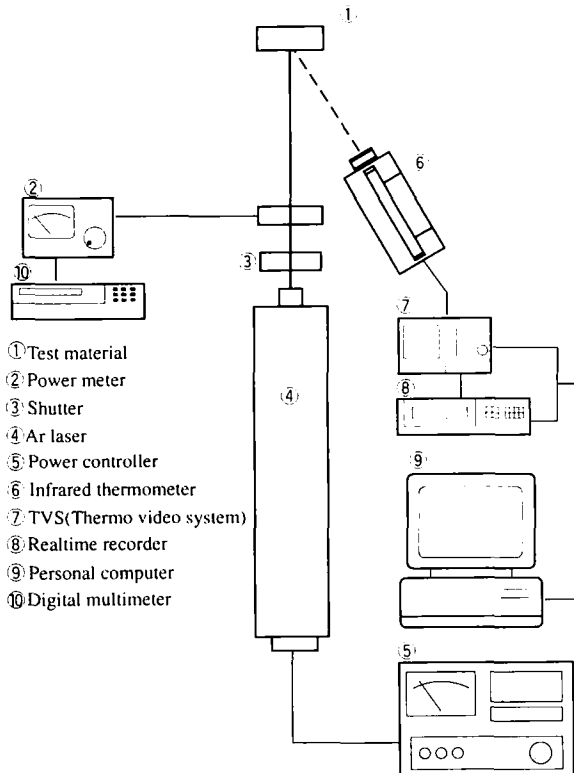


Fig. 4. Schematic diagram of the measuring system.

multimeter (10). The power meter has about 0.1-mW resolution. In the present measurement, the laser output is adjusted to about 20 mW, and under this condition, the surface central temperature rise at 3 s after step heating becomes 13–15 K. The variation of laser power before and after a measurement is within $\pm 1\%$.

The infrared thermometer system can measure the two-dimensional surface temperature distributions and, also, record the data every 1/30 s. The temperature range of 20–40°C is selected in the measurement. In this case, resolution of the temperature measurements is estimated to be about 0.1 K, however, some fluctuations of about 0.2 K are observed. The space resolution is estimated to be 0.1 mm, when the focal length of the thermometer is set to be 0.1 m as in the present case.

Figure 5 shows a schematic diagram of the sample as an anisotropic material. The samples used in the present study are multilayered sheets of kent-paper, vinyl chloride, and polyethylene resin film. The materials are 30 mm long, 20 mm high, and 13–16 mm thick. To minimize the effect of the laser beam reflection, a thin black paint layer is coated on the central part ($10 \times 10 \text{ mm}^2$) of the material surface. The layer is rubbed with a glass rod to have a $24 \pm 2\text{-}\mu\text{m}$ thickness. Such a degree of nonuniformity in thickness is confirmed to have no effect on the accuracy of the measurement. The absorptivity of the paint layer for argon laser beam is estimated to be 0.73 under such surface conditions [4]. The detail of the test materials is listed in Table I. The materials, Kent-a, -b, and -c, are composed of 60 sheets of kent-paper. As for Kent-d, each sheet is pasted together with a 50- μm -thick acrylic double-faced adhesive tape having the thermal conductivity $\lambda = 0.21 \text{ W} \cdot \text{m}^{-1} \cdot \text{K}^{-1}$ [6]. The materials, V.C. and P.E., are composed of 115 sheets of vinyl chloride resin film ($\lambda = 0.15\text{--}0.21 \text{ W} \cdot \text{m}^{-1} \cdot \text{K}^{-1}$ [6]) and 150 sheets of polyethylene resin film ($\lambda = 0.34\text{--}0.53 \text{ W} \cdot \text{m}^{-1} \cdot \text{K}^{-1}$ [6]), respectively.

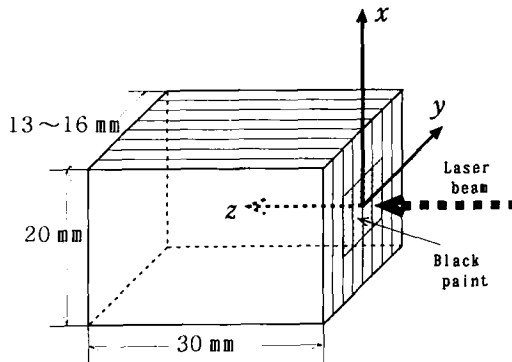


Fig. 5. Schematic diagram of the multilayered test material.

Table I. Parameters of the Multilayered Test Material

Test material	Thickness of paper or film (mm)	Number of sheets of paper or film	Bulk density ($\text{kg} \cdot \text{m}^{-3}$)
Kent-a	0.227 ± 0.003	60	911
Kent-b	0.224 ± 0.003	60	916
Kent-c	0.228 ± 0.003	60	915
Kent-d	0.226 ± 0.003	60	950
V.C.	0.115 ± 0.006	115	1300
P.E.	0.095 ± 0.002	150	970

The measurements were performed six times for each material at an ambient temperature of 22–23°C and a relative humidity of 70–80%. In the data processing, the coordinate system is taken as that shown in Fig. 5, that is, the x axis to be parallel to the paper or film edge direction.

4. RESULTS AND DISCUSSION

Figure 6 shows an example of the measured variations of $i_{\text{ex},x}$ and $i_{\text{ex},y}$ which are defined by Eqs. (12) and (13) as the variables corresponding to the average surface temperatures. As the larger fraction of heat flows along the paper or film edge direction, that is, in the x direction, $i_{\text{ex},x}$ is always higher than $i_{\text{ex},y}$. The fluctuations observed in the figure

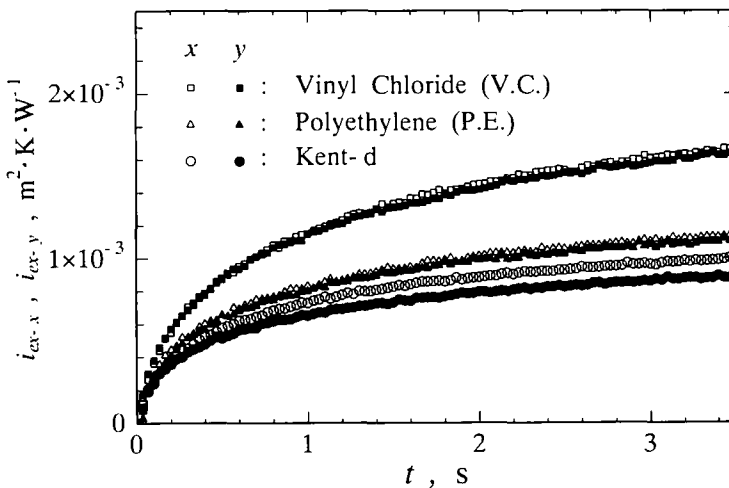


Fig. 6. Measured mean surface temperature variations.

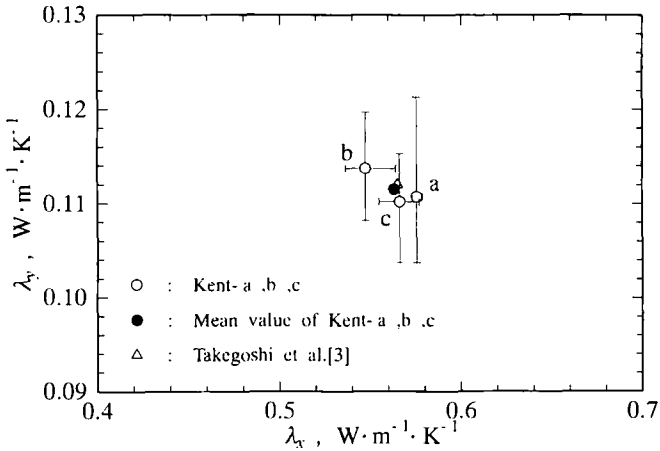


Fig. 7. Comparison of measured thermal conductivities with literature values. (○) Kent-b, -b, and -c; (●) mean value of Kent-a, -b, and -c; (△) value measured by Takegoshi et al. [3] with the transient hot-wire method.

are eliminated by approximating i_{ex-x} and i_{ex-y} with polynomial expressions of time t , and the expressions are used for evaluation process described previously.

In Fig. 7, the measured thermal conductivities λ_x and λ_y for Kent-a, -b, and -c are compared to those measured by Takegoshi et al. [3] with the transient hot-wire method. They used a sample composed of 60 sheets of kent-paper 200 mm long, 200 mm wide, and about 0.2 mm thick. The lines attached to each symbol show the range of scatter in each six measurements. As shown in Fig. 7, the mean value of Kent-a, -b and -c agrees well with those obtained by Takegoshi et al.

The measured results of λ_x , λ_y , α_x , α_y , and E_{xy} of the present multi-layered test materials are listed in Table II. Compared with Kent-a, -b, and -c, the thermal conductivity of Kent-d is lower in the x direction, and higher in the y -direction. The reason may be attributed to the effect of adhesive tape between the sheets of paper, and explained by considering the following equivalent thermal conductivities

$$\lambda_{x \dots eq} = \frac{\lambda_x \delta_s + \lambda_t \delta_t}{\delta_s + \delta_t} \tag{18}$$

$$\lambda_{y \dots eq} = \frac{\delta_s + \delta_t}{(\delta_x/\lambda_y) + (\delta_t/\lambda_t)} \tag{19}$$

Table II. Measured Thermophysical Properties^a

	λ_x	λ_y	α_x	α_y	E_{λ}
Kent-a	0.576	0.111	0.415	0.080	0.192
Kent-b	0.547	0.114	0.332	0.069	0.208
Kent-c	0.566	0.110	0.392	0.076	0.195
Mean value of Kent-a, -b, -c	0.563	0.111	0.380	0.075	0.198
Kent-d	0.497	0.155	0.222	0.069	0.311
V.C.	0.200	0.122	0.120	0.073	0.608
P.E.	0.342	0.275	0.141	0.113	0.803

^a Thermal conductivity, λ , in $\text{W} \cdot \text{m}^{-1} \cdot \text{K}^{-1}$ and thermal diffusivity, α , in $10^{-6} \text{m}^2 \cdot \text{s}^{-1}$.

These relations are derived from one-dimensional heat conduction model in each direction for the multilayered material where the kent-paper and the adhesive tape are piled up alternately. By using the values $\delta_t = 0.05 \times 10^{-3} \text{m}$ and $\lambda_t = 0.2 \text{W} \cdot \text{m}^{-1} \cdot \text{K}^{-1}$ for the adhesive tape, and the mean values $\delta_x = 0.226 \times 10^{-3} \text{m}$, $\lambda_x = 0.563 \text{W} \cdot \text{m}^{-1} \cdot \text{K}^{-1}$, and $\lambda_y = 0.111 \text{W} \cdot \text{m}^{-1} \cdot \text{K}^{-1}$ for kent-paper, the equivalent values of $\lambda_{x-cq} = 0.499 \text{W} \cdot \text{m}^{-1} \cdot \text{K}^{-1}$ and $\lambda_{y-cq} = 0.121 \text{W} \cdot \text{m}^{-1} \cdot \text{K}^{-1}$ are obtained from Eqs. (18) and (19), respectively. Although thus estimated values do not coincide perfectly with the measured values of Kent-d, the effect of the adhesive tape may be estimated well with these equations.

In order to examine the validity of the present method, the numerical simulations are performed again by using the measured thermophysical properties. The recalculated results are compared with the measured surface temperatures in dimensionless forms. Figure 8 shows the surface temperature profiles along the x and y axes of the kent-paper at 3 s after step heating. The variables I_{th} and I_{cx} are plotted against Fo in Fig. 9. In these figures, the lines and the symbols represent the recalculated and the measured results, respectively. It is seen that these recalculated and measured agree very well. The fact confirms that the present curve-fitting method is performed satisfactorily.

Assuming that the thermal conductivity λ_b , the thermal diffusivity α_b , and the thickness z_b of the black paint layer are exactly known, the accuracy of the present method is estimated and the errors are listed in Table III. These are obtained from numerical simulations for the effects of the measurement errors of the peak heat flux q_p , the heating radius r^* , and the surface temperature rise ΔT_w . The errors of q_p , r^* , and ΔT_w are

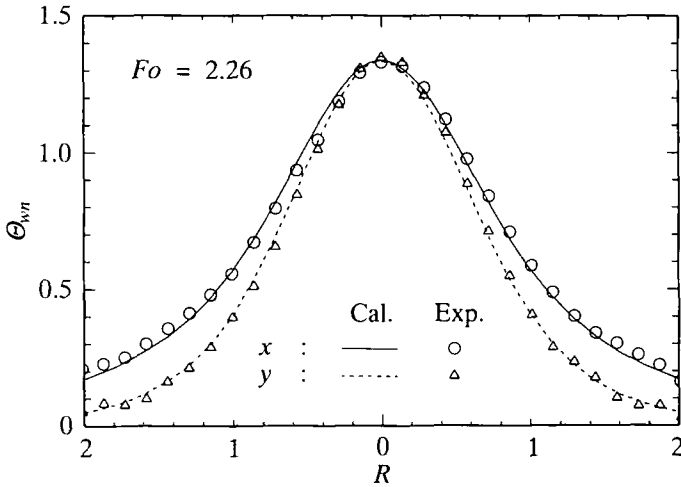


Fig. 8. Surface temperature profiles at 3 s after step heating.

estimated from the variations of the laser output before and after a measurements ($\pm 1\%$), the space resolution of the pin hole ($20\ \mu\text{m}$) and the temperature fluctuations of the infrared thermometer every $1/30\text{-s}$ measurement ($\pm 0.2\ \text{K}$), respectively. Therefore, the errors in r^* and ΔT_w are overestimated, especially for the latter, because the maximum fluctuation $0.2\ \text{K}$ is uniformly added to or subtracted from the surface temperature at

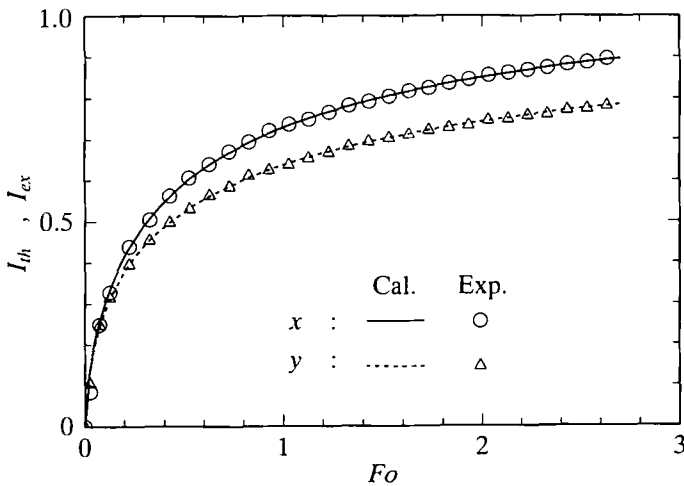


Fig. 9. Dimensionless mean surface temperature variations.

Table III. Estimates of Errors

Measured quantity	Measuring error	Error (%)		
		λ_v	α_v	E_{xy}
q_p ($W \cdot m^{-2}$)	2%	2.4 (2.4)	0.23 (0.28)	1.0 (1.0)
r^* (m)	3%	3.18 (3.14)	8.7 (8.7)	1.2 (1.2)
ΔT_u (K)	0.2 K	0 (0.1)	12.9 (45)	2.3 (5.0)
Total error (%)	-	6 (6)	22 (54)	5 (7)

every measurements. In each column of λ_v , α_v , and E_{xy} , the errors are indicated for two cases, that is, one for maximum temperature rise at 3 s after heating at $\Delta T_c \cong 13$ K and the other for $\Delta T_c \cong 5$ K (in parentheses). The latter is estimated for the purpose of the application of this method to the thermophysical property measurement of biological materials. The accuracy of λ_v depends largely on the accuracy of the measurement of q_p and r^* . As for α_v , the errors of r^* and ΔT_u are much more significant. In the case of $\Delta T_c \cong 5$ K, the accuracy of λ_v is the same for $\Delta T_c \cong 13$ K. The accuracy of α_v , however, remarkably reduces, since the relative error of ΔT_u to ΔT_c becomes large.

5. CONCLUSIONS

The noncontact method for measuring the thermal conductivity and thermal diffusivity proposed for isotropic materials is extended to anisotropic materials. The major conclusions are as follows:

(1) From the numerical simulations using the estimated thermophysical properties and comparisons with the measured surface temperatures, the validity of the present measurement method is confirmed. As for the kent-paper, the measured results agree well with those obtained by Takegoshi et al. [3] using the transient hot-wire method.

(2) It is clarified that the present method can measure the thermal conductivity λ_v , the thermal diffusivity α_v , and the anisotropic ratio E_{xy} within errors of ± 6 , ± 22 , and $\pm 5\%$ respectively, for the surface central temperature rise at 3 s after step heating at $\Delta T_c \cong 13$ K.

(3) From the error estimation for the case of $\Delta T_c \cong 5$ K, the method is found to be applicable at least for thermal conductivity measurements of biological materials.

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