

PULSE METHOD OF MEASURING THE THERMAL DIFFUSIVITY OF SPHERICAL SAMPLES

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Consideration is given to a version of the pulse method of measurement of the thermal diffusivity of spherical samples with the use of laser heating. The method is based on solution of the heat-conduction equation in a spherical coordinate system. The computerized experimental setup used is described. Measurement results for the thermal diffusivity of Zr, Ni, Fe, Al are reported. The measurement error is no more than 5%.

In [1-3] a pulse method of determination of the thermal diffusivity of liquid metals in a hemispherical form is considered where the heat source is at the center of the flat surface and the delay time of the temperature signal is measured at a point located some distance from the source of the thermal disturbance. High-temperature investigations of thermal diffusivity using this version of the method have demonstrated the difficulty of introducing corrections for heat transfer, especially for a flat surface. In connection with this it is reasonable to use a sample in the form of a small-diameter drop, which in calculations can be approximated by a sphere of radius R with a point source at a pole.

The temperature T at the point with the spherical coordinates (r, θ, φ) caused by the action of a single instantaneous point source, located at the pole $(R; 0; 0)$ at the initial moment of time, at $d^2 = R^2 + r^2 - 2Rr \cos \theta$ is of the form [4]

$$T(d; t) = \frac{Q}{(4\pi at)^{3/2} \rho c_p} \exp\left(-\frac{d^2}{4at}\right), \quad (1)$$

Expression (1) satisfies the equation

$$\frac{\partial T(r; \theta; t)}{\partial t} = a \left[\frac{\partial^2 T(r; \theta; t)}{\partial r^2} + \frac{2}{r} \frac{\partial T(r; \theta; t)}{\partial r} + \frac{1}{r^2 \sin \theta} \frac{\partial}{\partial \theta} \left(\sin \theta \frac{\partial T(r; \theta; t)}{\partial \theta} \right) \right] \quad (2)$$

and is the solution of the problem of the temperature distribution in an unbounded medium.

Using the image method with account for the boundary conditions

$$T(r; \theta; 0) = 0; \quad -\lambda \left. \frac{\partial T}{\partial r} \right|_{r=R} = \alpha T; \quad 0 \leq r \leq R; \quad 0 \leq \theta \leq \pi$$

and expression (1), one can obtain the distribution of the temperature field in a sphere

$$T(r; \theta; t) = \frac{2Q}{\rho c_p} \frac{\exp\left(-\frac{d^2}{4at}\right)}{(4\pi at)^{3/2}} \left[1 - \frac{\alpha}{\lambda} \sqrt{\pi at} \times \right.$$

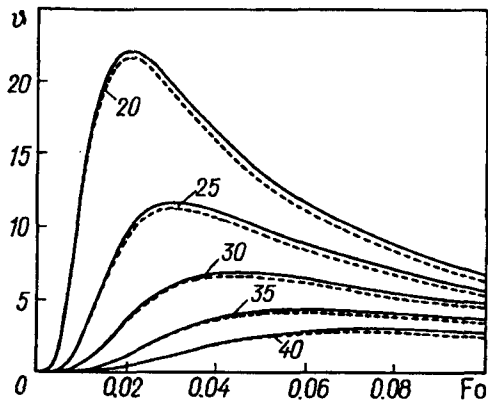


Fig. 1. Relative-temperature (ϑ) distribution as a function of dimensionless time (Fo) (the figures at the curves are values of θ , the solid curves refer to $Bi = 0$, the dashed curves refer to $Bi = 0.1$).

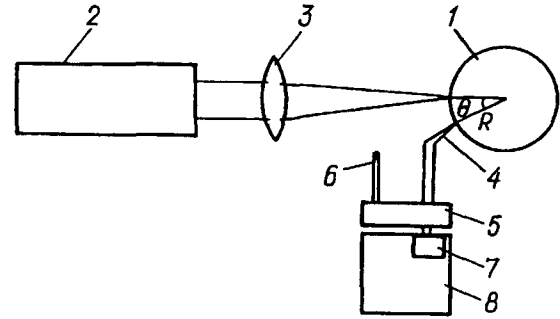


Fig. 2. Schematic of the experimental setup.

$$\times \exp \left(at \left(\frac{2R - 2r \cos \theta}{4at} - \frac{\alpha}{\lambda} \right)^2 \right) \operatorname{erfc} \left(\sqrt{at} \left(\frac{2R - 2r \cos \theta}{4at} - \frac{\alpha}{\lambda} \right) \right) \right]. \quad (3)$$

The temperature $T(r; \theta; t) = T' - T_0$ will be referred to as the excess temperature if the ambient temperature $T_0 \neq 0$ and T' is the sample temperature.

To measure thermal diffusivity, it is convenient to use dimensionless parameters. For this, expression (3) at $r = R$, which corresponds to the temperature-field distribution on the sphere surface, can be written in terms of the relative quantities $\vartheta(\theta; Fo; Bi)$ as

$$\vartheta = \frac{\exp \left(-\frac{1 - \cos \theta}{2Fo} \right)}{2\sqrt{\pi} Fo^{3/2}} \left[1 - Bi \sqrt{\pi} Fo \exp \left(Fo \left(\frac{1 - \cos \theta}{2Fo} - Bi \right)^2 \right) \operatorname{erfc} \left(\sqrt{Fo} \left(\frac{1 - \cos \theta}{2Fo} - Bi \right) \right) \right], \quad (4)$$

where $\vartheta = 2\pi R^3 \rho c_p T/Q$; $Fo = at/R^2$ has the meaning of dimensionless time; $Bi = \alpha R/\lambda$.

Figure 1 shows calculated dependences of the relative excess temperature on the Fourier number at different points of the spherical surface for $Bi = 0$ and $Bi = 0.1$.

Obtained expression (4) allows determination of the thermal diffusivity of metals and alloys on the experimental setup (Fig. 2). A thermal pulse generated by GOR-100M laser 2 with a wavelength of $0.694 \mu\text{m}$ is sent to the pole of spherical sample 1 of radius R . The condition of a point source, the size of which is 0.3 mm ($0.3 \ll d$), at the point of exposure is achieved by focusing the thermal pulse by lens 3 with a focal power of 10 diopters. To record a temperature signal in time, chromium-nickel thermocouple 4 with an electrode thickness of 0.05 mm was welded to the sphere surface at the angle θ . The thermocouple was protected against direct thermal radiation with Al_2O_3 paste. The angle θ was measured by means of a BMI-1Ts toolmaker's microscope with an accuracy of up to $1'$. Since the temperature drop reaches $\sim 2^\circ\text{C}$ on the sphere surface with the coordinates $(R; \theta; 0)$, amplifier 5 is used in the setup. To obtain a low level of noise and a high common-mode rejection factor, the input stages of the amplifier are connected as differentiators on low-noise operational amplifiers of the type K140UD17. To determine the reference point of a thermal signal, photodiode 6 is used. Amplified signals of the photodiode and the thermocouple are sent via analog-to-digital converter 7 to computer 8 of the type IBM PC [5].

The value of $Fo_{1/2}$ found from the theoretical (see Fig. 1) and computer-processed experimental (Fig. 3) curves for the given θ allows determination of the thermal diffusivity by the formula

TABLE 1. Thermal-Diffusivity Values ($a \cdot 10^6$, m^2/sec) According to Experimental Data

Metal	1	2	3
Nickel	16.5	16	22.7
Iron	17.4	18	22.0
Zirconium	13.3	—	12.7
Aluminum	92.4	94	93.2

Note: 1 denotes data of the present authors; 2, [6]; 3, [7].

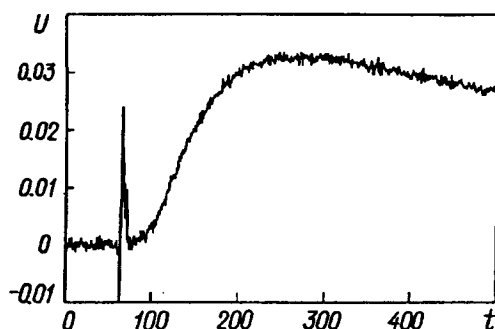


Fig. 3. Experimental curve of the thermoelectromotive force obtained by a Chromel-Alumel thermocouple for a nickel sample at $R = 6.967$ mm, $\theta = 33^\circ 51'$, U , mV; t , msec.

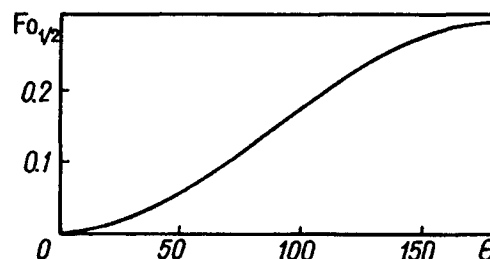


Fig. 4. Plot of $Fo_{1/2}$ vs. the angle θ .

$$a = Fo_{1/2} \frac{R^2}{\tau_{1/2}} \quad (5)$$

Figure 4 shows $Fo_{1/2}$ as a function of the angle θ .

The results of measurement of the thermal diffusivity of Ni (99.99), Zr (99.9), Al (99.99), and Armco iron carried out at a temperature of 300 K (see Table 1) agree with the values obtained by the Parker pulse method [6] and the method of temperature waves [7]. It should be noted that the method of temperature waves gives the overestimated results as compared to the pulse methods, and the discrepancy in the thermal-diffusivity values obtained by different methods reaches 30% (see the references in [7]). The error in thermal-diffusivity measurement does not exceed 5% and consists of:

- the measurement error due to the inaccuracy in the geometric dimensions of the sample in the experiment (no more than 1%);
- the standard deviation of the thermal diffusivity due to time measurement (in the course of the same experiment it does not exceed 1%);
- the error of thermal-diffusivity measurement related to inaccurate determination of the angle θ , which is related, in turn, to $Fo_{1/2}$ (within the limits of 2–2.5%).

To sum up, the developed version of the pulse method of thermal-diffusivity measurement will allow determination of thermophysical characteristics of small samples in the form of a drop at high temperatures in both the solid and liquid states without changing the sample itself.

NOTATION

R , sphere radius; θ , angle; Fo , Fourier number; ϑ , relative temperature; Bi , Biot number; $Fo_{1/2}$, Fourier number corresponding to half the maximum relative temperature; $\tau_{1/2}$, time in which the temperature signal reaches half the maximum value; Q , released heat; ρ , density; a , thermal diffusivity; c_p , heat capacity; α , heat-transfer coefficient; λ , thermal conductivity.

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