

We thank Dr J. Sikorski of the Textile Department, University of Leeds, for access to the Stereoscan microscope and for his support, and Mr T. Buckley of the same department for his practical assistance.

I. J. SMALLEY  
J. G. CABRERA

Department of Civil Engineering,  
University of Leeds,  
Leeds.

Received February 24, 1969.

<sup>1</sup> Lambe, T. W., *Proc. Amer. Soc. Civ. Eng.*, **79**, No. 315 (1953).

<sup>2</sup> Rosenqvist, I. T., *J. Soil Mech. and Found. Div., Amer. Soc. Civ. Eng.*, **85**, No. SM2, 31 (1959).

<sup>3</sup> Sloane, R. L., and Kell, T. R., *Clays and Clay Min., Proc. Fourteenth Nat. Conf.*, 289 (1966).

<sup>4</sup> Kirkham, D., De Boodt, M. F., and De Leenheer, L., *Soil Sci.*, **87**, 141 (1959).

<sup>5</sup> Smart, P., *Clays and Clay Min., Proc. Fifteenth Nat. Conf.*, 241 (1967).

<sup>6</sup> Schofield, R. K., and Samson, H. R., *Disc. Farad. Soc.*, **18**, 135 (1954).

## Determination of the Firing Temperature of Ancient Ceramics by Measurement of Thermal Expansion

THE thermal expansion method for determining firing temperatures<sup>1-3</sup> is based on the assumption that, when clays are fired, shrinkage occurs as a result of various sintering processes such as vitrification. Consequently when a clay ceramic is heated up from room temperature (see Fig. 1) it typically exhibits a reversible expansion, until temperatures comparable with the original firing temperature are reached. With continued increase in temperature the ceramic begins to contract because superimposed on the reversible expansion there is an irreversible shrinkage associated with resumption of sintering (that is, the firing of the ceramic is being continued beyond the point reached during the original firing). The temperature ( $T_a$ ) at which a net shrinkage is first observed should therefore provide an indication of the original firing temperature ( $T_e$ ) of the ceramic.

Firing temperature determinations were undertaken on a selection of ceramics with a wide range of provenance and age (Table 1). Specimens (2.5 × 1.0 × 1.0 cm) were cut from the pottery sherds and the shrinkage temperature,  $T_a$ , was measured using extension rod dilatometers<sup>3,4</sup>. The specimen was then refired for 1 h at temperature,

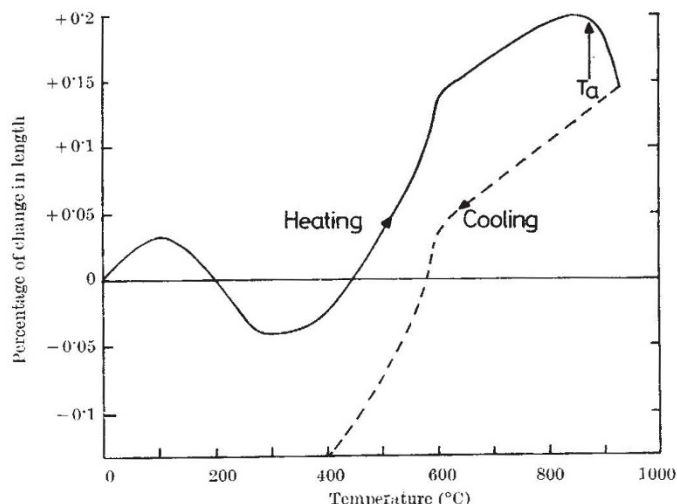


Fig. 1. Thermal expansion curves during heating and cooling for a typical ancient ceramic specimen (Pottersbury ware).

Table 1. FIRING TEMPERATURE OF ANCIENT CERAMICS

Provenance	Archaeological data Period/age Type	Firing temperature ( $T_e$ °C)
Turkey	c. 5000 BC	750–820
Iraq	c. 4500 BC	970–1,050†
Iraq	c. 3500 BC	1,140–1,180
Cyprus	c. 3000 BC	500–700
Cyprus	c. 1300 BC	940–1,000†
Cyprus	c. 1300 BC	1,030–1,070†
Turkey	c. 500 BC	1,000–1,100
Nigeria	—	< 500
China	c. 1100 AD	1,070–1,140*
China	c. 1100 AD	1,070–1,190*
China	c. 1700 AD	960–1,050*
England	Iron Age	Calcite gritted ware
England	Roman	Black burnished ware
England	Roman	Grey ware
England	Roman	Colour coated ware
England	Roman	Mortarium
England	Roman	Samian ware
England	Saxon	—
England	Saxon	Theftord ware
England	Mediaeval	Laverstock ware
England	Mediaeval	Cowick ware
England	c. 1650 AD	Pottersbury ware

\* Presence of low viscosity liquid phase— $T_e$  possibly too low.

† Bloating— $T_e$  possibly too high.

$T_e'$ , and the new shrinkage temperature,  $T_a'$ , was measured.

The values obtained for  $T_a$  ranged from 620°–1,230° C. When  $T_a$  was less than 700° C, the thermal expansion data provided no precise information on the firing temperature and it was only possible to suggest that  $T_e$  was either less than 500° C or in the 500–700° C range, depending on whether or not hydrated clay minerals were present in the ceramics<sup>5</sup>. Similarly when the ceramic contained calcite,  $T_e$  was probably less than 800° C; the value for  $T_a$  being invalid because this mineral produced a complex pattern of expansion and contraction. For the remainder of the ceramics,  $T_e$  was calculated using the relationship

$$(T_e - T_a) = (T_e' - T_a')$$

Measurements on clay specimens, prefired at known temperatures, indicated that this relationship was valid for a wide range of firing temperatures provided vitrification occurred during the original firing and  $T_e'$  was selected such that  $(T_e' - T_e)$  equalled approximately 20–30° C. Even in these circumstances, however, the value obtained for  $T_e$  may be too high as a result of bloating caused by the expansion of gases trapped in the liquid phase of the clay body. Alternatively  $T_e$  may be too low because shrinkage, resulting from the pressure exerted on the specimen by the dilatometer components, can occur when the viscosity of the liquid phase decreases at high temperatures.

The values estimated for  $T_e$  (Table 1) show that firing temperatures ranging from 500°–1,200° C were used in the manufacture of ancient ceramics and that low or high firing temperatures were not confined to particular periods or parts of the world. The data provide valuable information on the performance of ancient pottery kilns and on the technological capability of the potters. A knowledge of the firing temperature can also be valuable in other scientific investigations of ancient ceramics and kilns such as thermoluminescent and magnetic dating.

I thank Professor J. P. Roberts for helpful discussions on this problem. This work was supported by the Science Research Council and an ICI research fellowship.

M. S. TITE\*

Department of Ceramics,  
Houldsworth School of Applied Science,  
University of Leeds.

Received January 9, 1969.

\* Present address: Department of Physics, University of Essex, Colchester.

<sup>1</sup> Kiefer, Ch., *Bull. Soc. Franc. Ceram.*, **30**, 3 (1956); **31**, 17 (1956).

<sup>2</sup> Terrisse, J. R., *Rei Cretariae Romanae Faكتورum*, **2**, 63 (1959).

<sup>3</sup> Roberts, J. P., *Archaeometry*, **6**, 21 (1963).

<sup>4</sup> Baudran, A., *Bull. Soc. Franc. Ceram.*, **27**, 13 (1955).

<sup>5</sup> Perinet, G., *Trans. Seventh Intern. Ceram. Cong., London*, 371 (1960).