

Thermal Analysis & Rheology

THERMAL APPLICATIONS NOTE DETERMINING MINIMUM USABLE SAMPLE THICKNESS IN TMA

Thermomechanical analysis (TMA) is a technique widely used to determine transition temperatures and expansion/ contraction properties of materials, including the coefficient of thermal expansion (α). When determining α , however, there are three factors which need to be considered to obtain reproducible quantitative results. These are TMA sensitivity and baseline drift, sample thickness, and the magnitude of α .

The TA Instruments TMA 2940 has a maximum sensitivity of 0.1 μ m, a maximum resolution of 3nm, and a baseline drift (0 to 500°C) of about 0.5 μ m. Therefore, the recommended measured sample dimensional change over this range for α determinations is >5 μ m (roughly 10x the baseline drift). Knowing this, the minimum sample thickness required for testing can be determined from:

$$\alpha = \frac{\Delta L}{\Delta T} * \frac{1}{L_o} \implies L_o = \frac{\Delta L}{\Delta T} * \frac{1}{\alpha}$$
[1]

where: ΔL = measured dimension change

 $L_o = original sample thickness$

 ΔT = temperature range of measurement

For example, a material with an expected α of 50 μ m/m°C would yield a recommended minimum sample thickness for testing of:

$$L_{o} = \left(\frac{5\mu m}{500^{\circ} C}\right) * \left(\frac{1}{50\mu m / m^{\circ} C}\right) = 200\mu m$$
[2]

This minimum sample thickness will obviously change with α and temperature range. Clearly, it is better to use samples much thicker than this minimum if possible, as this will further improve signal-to-noise ratio.

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