

Designation: E2448 - 22

Standard Test Method for Determining the Superplastic Properties of Metallic Sheet Materials¹

This standard is issued under the fixed designation E2448; the number immediately following the designation indicates the year of original adoption or, in the case of revision, the year of last revision. A number in parentheses indicates the year of last reapproval. A superscript epsilon (ε) indicates an editorial change since the last revision or reapproval.

1. Scope

1.1 This test method describes the procedure for determining the superplastic forming properties (SPF) of a metallic sheet material. It includes tests both for the basic SPF properties and also for derived SPF properties. The test for basic properties encompasses effects due to strain hardening or softening.

1.2 This test method covers sheet materials with thicknesses of at least 0.5 mm but not greater than 6 mm. It characterizes the material under a uni-axial tensile stress condition.

Note 1—Most industrial applications of superplastic forming involve a multi-axial stress condition in a sheet; however it is more convenient to characterize a material under a uni-axial tensile stress condition. Tests should be performed in different orientations to the rolling direction of the sheet to ascertain initial anisotropy.

1.3 This method has been used successfully between strain rates of 10^{-5} mm/mm/s to 10^{-1} mm/mm/s second.

1.4 This method has been used successfully on Aluminum and Titanium alloys. The use of the method with other metals should be verified.

1.5 The values stated in SI units are to be regarded as standard. No other units of measurement are included in this standard.

1.6 This standard does not purport to address all of the safety concerns, if any, associated with its use. It is the responsibility of the user of this standard to establish appropriate safety, health, and environmental practices and determine the applicability of regulatory limitations prior to use.

1.7 This international standard was developed in accordance with internationally recognized principles on standardization established in the Decision on Principles for the Development of International Standards, Guides and Recommendations issued by the World Trade Organization Technical Barriers to Trade (TBT) Committee.

2. Referenced Documents

- 2.1 ASTM Standards:²
- E4 Practices for Force Calibration and Verification of Testing Machines
- E6 Terminology Relating to Methods of Mechanical Testing
- E21 Test Methods for Elevated Temperature Tension Tests of Metallic Materials
- E177 Practice for Use of the Terms Precision and Bias in ASTM Test Methods
- E646 Test Method for Tensile Strain-Hardening Exponents (*n* -Values) of Metallic Sheet Materials
- E691 Practice for Conducting an Interlaboratory Study to Determine the Precision of a Test Method

3. Terminology

3.1 Definitions:

3.1.1 engineering strain, e, n—a dimensionless value that is the change in length (ΔL) per unit length of original linear dimension (L_0) along the loading axis of the specimen; that is, $e = (\Delta L)/L_0$.

3.1.2 engineering stress, S [FL⁻²], *n*—the normal stress, expressed in units of applied force, *F*, per unit of original cross-sectional area, A_0 ; that is, $S = F/A_0$.

3.1.3 *true strain*, ε , *n*—the natural logarithm of the ratio of instantaneous gauge length, *L*, to the original gauge length, *L*₀; that is, $\varepsilon = \ln (L/L_0)$ or $\varepsilon = \ln (1+e)$.

3.1.4 *true stress*, σ [FL⁻²], *n*—the instantaneous normal stress, calculated on the basis of the instantaneous cross-sectional area, *A*; that is, $\sigma = F/A$; if no necking has occurred, $\sigma = S(1+e)$.

3.1.5 Refer to Terminology E6 for the definitions of the terms extensometer system, indicated temperature, necking, specified temperature, strain hardening, and stress-strain diagram.

3.2 Definitions of Terms Specific to This Standard:

3.2.1 gauge length, L, [L], n—the instantaneous distance between the shoulders of the test specimen during the test

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² For referenced ASTM standards, visit the ASTM website, www.astm.org, or contact ASTM Customer Service at service@astm.org. For *Annual Book of ASTM Standards* volume information, refer to the standard's Document Summary page on the ASTM website.

3.2.2 *testing machine crosshead velocity, V,* [L/T], *n*—the velocity of the traveling member of the testing machine to which one of the test specimen clamps is attached.

3.2.3 *strain rate*, ε , [1/T], *n*—the time rate of change of the true strain in the test specimen, measured as: $\frac{V}{L_0(1+e)}$

where:

 L_0 = original gauge length

V = machine crosshead velocity

3.2.3.1 *Discussion*—This is an operational definition of strain rate.

3.2.4 strain rate sensitivity, m, n—(ln $\Delta \sigma$)/ (ln $\Delta \dot{\epsilon}$) $\dot{\epsilon}$.

3.2.4.1 *Discussion*—In practical terms, $m = \frac{\ln(\sigma_2 \neq \sigma_1)}{\ln(\epsilon_2 \neq \epsilon_1)}$ under stated test conditions, see 7.2.1.

4. Significance and Use

4.1 The determination of the superplastic properties of a metallic sheet material is important for the observation, development and comparison of superplastic materials. It is also necessary to predict the correct forming parameters during an SPF process. SPF tensile testing has peculiar characteristics compared to conventional mechanical testing, which distort the true values of stress, strain, strain hardening, and strain rate at the very large elongations encountered in an SPF pull test, consequently conventional mechanical test methods cannot be used. This test method addresses those characteristics by optimizing the shape of the test specimen and specifying a new test procedure.

4.2 The evaluation of a superplastic material can be divided into two parts. Firstly, the basic superplastic-forming (SPF) properties of the material are measured using the four parameters of stress, temperature, strain, and strain rate. These are obtained using conversions from the raw data of a tensile test. Secondly, derived properties useful to define an SPF material are obtained from the basic properties using specific equations.

4.3 The test specimen undergoes an essentially uniform and constant necking along its length, and *S* and *e* are assumed in this standard to be valid. However at the junction to the clamp sections of the test specimen the cross section reduces from the original value to the final value, over a length of approximately 4 % at each end. Also, there are local small instabilities of cross section over the gauge length. These contribute to an error in the calculated values of ε and σ . In the absence of currently available extensioneters that could operate in the elevated-temperature environment of an SPF test, ε and σ are to be inferred from crosshead extension and force.

4.4 The derived term m is widely used to describe the SPF properties of a material. It should be used with caution, as it is dependent on strain, strain rate and temperature. Many references in the literature do not identify the strain condition at which the readings were taken, or allow multiple strains to be used in the determination of m.

4.5 Many superplastic alloys exhibit strain hardening. However, the conventional strain hardening exponent n as defined in Test Method E646 is not valid for superplastic materials as strain hardening in the latter is usually a coefficient of strain, rather than an exponent. The mechanism of strain hardening in superplastic flow is essentially due to grain growth, and although the stress/strain relationship is often linear, it is not universal for all superplastic materials. Consequently, there is no simple definition of a strain hardening coefficient and this standard does not define one. Consideration of strain hardening in superplastic deformation is discussed in Ghosh and Hamilton's, "Influences of Material Parameters and Microstructure on Superplastic Forming."³

4.6 It is assumed no local necking takes place and the cross section of the test specimen is constant over the entire gauge length.

NOTE 2—For some materials, cavitation inside the material increases the volume of the gauge section as the test progresses and changes the true cross-sectional area. For other materials, the test specimen develops a ribbed or other local texture, and changes the minimum cross section.

4.7 It is assumed that the increasingly non uniform cross section that develops at each end of the test specimen where the gauge section transitions to the original width at the clamp section is small and can be ignored.

5. Apparatus

5.1 The force-measuring system of the testing machine shall conform to the requirements of Practices E4.

5.2 The apparatus shall be calibrated according to appropriate standards or manufacturer instructions.

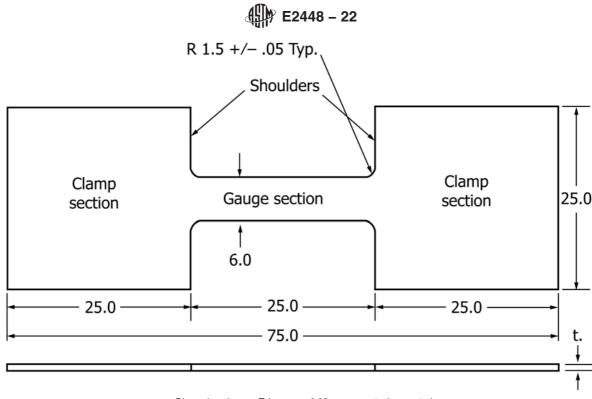
5.3 No extensioneter system is used in this test method, and the extension of the test specimen is measured at the testing machine crosshead. The accuracy of the recorded crosshead position shall be better than 0.25 mm. The testing machine compliance shall be determined before testing, and the amount of compliance subtracted from the crosshead position if it exceeds 1 % of the original gauge length of the test specimen.

Note 3—One method of determining compliance is to mount a 6 mm thick test specimen in the clamps without heating, then load the testing machine to the estimated maximum force of the test and measure the movement of the crosshead. Due to the low forces of these tests (typically 100 N maximum) compliance is likely to be small.

5.4 The tensile testing machine shall be computer controlled and capable of varying the testing machine crosshead velocity in order to maintain a near constant strain rate. The testing machine crosshead velocity may be increased in steps. The instantaneous strain rate may vary up to 1 % from nominal strain rate.

5.5 The tensile testing machine shall be provided with clamps that hold the test specimen at and under the shoulders adjacent to the gauge section. The test specimen shall not be compressed by the clamps, as this will induce superplastic flow out of the clamp area during the test. Clamp design should follow that shown in Fig. 2.

³ Ghosh, A. K., and Hamilton, C. H., "Influences of Material Parameters and Microstructure on Superplastic Forming," *Met Trans A*, Vol 13A, May 1982, pp. 733-742.



Dimensions in mm. Tolerance ± 0.25 mm except where noted. FIG. 1 Dimensions of Test Specimen

5.6 The apparatus shall be provided with a furnace that shall maintain the test specimen at a constant temperature throughout the test. Test equipment shall meet the requirements of Test Methods E21 for temperature measuring, calibration, and standardization.

6. Procedure

6.1 Test specimens shall be made to the dimensions shown in Fig. 1. The test specimen width and gauge thickness, *t*, shall be measured and recorded at a minimum of four places in the gauge section, to a tolerance of 1 % of reading, or 12 μ m, whichever is greater.

6.2 If material oxidation affects the superplastic behavior of the material, the furnace may be flooded with argon or other inert gas to reduce the effects of oxidation.

6.3 Before starting the test, bring the furnace up to the specified temperature and stabilize the temperature. Install the test specimen into the clamps. During the heat up of the test specimen, minimize external force from the testing machine to the test specimen.

Note 4—Many testing machines incorporate a "protect specimen" or "load control" option during the heating phase to accommodate the thermal expansion of the test specimen/grip assembly inside the furnace and to prevent buckling of the test specimen. This control option ensures "almost" zero force on the test specimen during heating through the movement of the cross-head beam.

6.4 Load the test specimen as soon as the indicated temperature is within the allowable range specified in 6.5.

NOTE 5—In normal tensile testing, the test specimen is not loaded until test specimen and load train have reached thermal equilibrium. A common method for determining thermal equilibrium is to observe that the crosshead has ceased to move under the "protect specimen" control,

indicating that no more thermal expansion is taking place. However, this time can be long enough to allow grain growth in the test specimen, which distorts the superplastic properties being evaluated.

6.4.1 The start of loading may be delayed until thermal equilibrium is achieved if it is known that grain growth during thermal equilibration is small enough so that SPF properties will not be affected.

6.5 For the duration of the test, defined as the time from start of loading until the termination of test or fracture, the difference between indicated temperature and specified temperature shall not exceed the following limits:

Specified temperatures less than or equal to 700 °C: ± 3 °C Specified temperatures greater than 700 °C: ± 6 °C.

6.6 During the test, increase the testing machine crosshead velocity, V, according to the equation

$$V = \dot{\varepsilon}_{\rm nom} \left(\frac{L_0}{1+e} \right) \tag{1}$$

to maintain the true strain rate, $\dot{\epsilon}$, to within ± 1 % of the nominal strain rate, $\dot{\epsilon}_{nom}$, until a predetermined strain value is reached or until fracture:

$$\left| \frac{\dot{\varepsilon} - \varepsilon_{\text{nom}}}{\dot{\varepsilon}_{\text{nom}}} \right| \le 0.01 \tag{2}$$

6.7 Record the force and crosshead extension at least twice per second to an accuracy of ± 1 % of the recorded value.

6.7.1 As the clamp extension rod is pulled out of the furnace, it cools and contracts, thereby altering the distance between crosshead and clamp. This error in reading is small compared to the test specimen gauge length, *L*, and may be ignored.