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Standard Test Method for Measurement of Internal Stress of Plated Metallic Coatings with the Spiral Contractometer¹

This standard is issued under the fixed designation B 636; 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.

 ϵ^1 Note—Section 12 was added, the title changed to "Test Method," and other editorial changes made in January 1992.

1. Scope

1.1 This test method covers the use of the spiral contractometer for measuring the internal stress of metallic coatings as produced from plating solutions on a helical cathode. The test method can be used with electrolytic and autocatalytic deposits.

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

2. Terminology

2.1 Definitions:

2.1.1 *compressive stress* (-)—stress that tends to cause a deposit to expand.

2.1.2 *internal stress*—the net stress that remains in a deposit when it is free from external forces. The internal stress tends to compress or stretch the deposits.

2.1.3 *tensile stress* (+)—stress that tends to cause a deposit to contract.

3. Summary of Test Method

3.1 The test method of measuring stress with the spiral contractometer is based on plating on the outside of a helix. The helix is formed by winding a strip around a cylinder, followed by annealing. In operation, one end of the helix is fixed and the other is allowed to move as stresses develop. The free end is attached to an indicating needle through gears that magnify the movement of the helix. As the helix is plated, the stress in the deposit causes the helix to wind more tightly or to unwind, depending on whether the stress is compressive (-) or tensile (+). From the amount of needle deflection and other data, the internal stress is calculated.

3.2 With instrument modifications, the movement of the helix can be measured electronically instead of mechanically as described in 3.1.

4. Significance and Use

4.1 The spiral contractometer, properly used, will give reproducible results (see 8.5) over a wide range of stress values. Internal stress limits with this method can be specified for use by both the purchaser and the producer of plated or electroformed parts.

4.2 Plating with large tensile stresses will reduce the fatigue strength of a product made from high-strength steel. Maximum stress limits can be specified to minimize this. Other properties affected by stress include corrosion resistance, dimensional stability, cracking, and peeling.

4.3 In control of electroforming solutions, the effects of stress are more widely recognized, and the control of stress is usually necessary to obtain a usable electroform. Internal stress limits can be determined and specified for production control.

4.4 Internal stress values obtained by the spiral contractometer do not necessarily reflect the internal stress values found on a part plated in the same solution. Internal stress varies with many factors, such as coating thickness, preparation of substrate, current density, and temperature, as well as the solution composition. Closer correlation is achieved when the test

b4740a50 conditions match those used to coat the part. b636-841992e1

5. Apparatus

5.1 The spiral contractometer is described by A. Brenner and S. Senderoff.²

NOTE 1—Spiral contractometers are available from many of the suppliers of nickel sulfamate.

5.2 Helices shall be stopped-off on the inside to prevent plating. Helices are available with or without a permanent inert coating on the insides (see Appendix X1).

5.3 The clamps holding the helix to the contractometer shall be coated with an inert nonconductive coating to prevent their plating and acting as thieves.

5.4 For testing electroplating solutions, anodes are placed equidistant from the helix and symmetrically positioned to produce even plate distribution. A minimum of four anodes is required. A concentric anode arrangement is preferred.

¹ This test method is under the jurisdiction of ASTM Committee B-8 on Metallic and Inorganic Coatingsand is the direct responsibility of Subcommittee B08.10on General Test Methods.

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² Brenner, A., and Senderoff, S., *Proceedings of the American Electroplaters Society*, Vol 35, 1948, p. 53.

5.5 Laboratory tests on electroplating solutions shall utilize at least 3.7 L of solution. A 4-L beaker with an annular anode arrangement is convenient. Use of this volume or larger will minimize solution changes due to electrolysis during the test.

5.6 Laboratory tests on autocatalytic plating solutions are done in a 1-L, tall-form beaker. Obviously, no anodes are used.

6. Factors Affecting Accuracy

6.1 Variations in the preparation of the helix may cause substantial variations in results.

6.1.1 Stop-off material shall be applied properly to the interior of the helix. The stop-off material shall be thin and flexible to permit the helix to move freely during the test. A coating weight of less than 50 mg/dm² is satisfactory.

NOTE 2—The inside must be stopped-off with some inert, flexible coating. One acceptable stop-off material is "Microstop."³ One part of "Microstop" is diluted with two parts of acetone before use. Any nickel deposited on the inside of the helix will exhibit an opposing effect.

6.1.2 Helices that have been permanently coated on the inside with TFE-fluorocarbon may give variable results when testing near-zero stresses.

6.1.3 Cleaning variations and surface preparation of the helix before the test can produce varying results. For example, electrocleaning of the helix shall always be cathodic and controlled with respect to current, time, and temperature. Anodic cleaning at this stage can give wide variations. Abrasive cleaning of the helix and the use of etchants shall be avoided.

6.1.4 Very thin deposits of less than about 3 μ m are influenced more by the surface conditions and preparation of the helix than are thicker deposits.

6.2 Internal stress varies with current density used in electroplating. The variation is not predictable and depends on the metal being deposited, impurities or additives, and the current density range under consideration. It is important that the current be measured and controlled closely throughout the stress test. Variations in currents shall be held to less than 2 %.

6.3 Because the temperature of the plating solution may affect the internal stress, it shall be maintained within 2°C during the test. The initial rest point of the indicator and the final rest point are both taken at the operating temperature of the plating solution to eliminate thermal stresses.

6.4 The solution composition shall not vary during the test. Usually, if the repeatability tests in 8.5 are met, the solution can be assumed to be unchanged during the test runs. Conversely, when the repeatability tests are not met, the plating solution shall be analyzed to determine if any changes in solution composition have occurred during the test.

6.4.1 Tests run on electroplating solutions using insoluble or inefficient anodes could result in significant solution changes during the test.

6.4.2 When testing autocatalytic plating solutions, the constituents of the plating solution may be significantly depleted during the test, unless replenished.

6.5 A relationship between the surface area to be plated and the volume of autocatalytic plating solutions exists that may affect the character of the deposit. In testing autocatalytic plating solutions, the ratio of plated surface area to the volume of solution that is normally used in the plating tank shall be maintained. When using proprietary solutions, the supplier's recommendation shall be followed.

7. Calibration

7.1 Calibrate the instrument as directed in the manufacturer's instructions.

7.2 The frequency of calibration will vary with use and extent of attack on the helices from the chemical stripping. When visible attack is noted, discard the helix.

7.3 The calibration procedure consists essentially of determining the force required per degree of dial deflection. A known mass is suspended over a small pulley on a lever arm with the helix mounted in place. The degree of deflection is read from the dial. The data required for the calibration calculations as expressed in metric units are as follows:



8. Procedure

6-8.1 The procedure will vary with the solution being tested. Follow the instructions given by the supplier carefully. Variations in the procedure can produce variations in results. Give appropriate attention to the factors in Section 6. A detailed procedure for nickel plating solutions appears in Appendix X1.

8.2 Position the spiral contractometer in electroplating solutions so that it is equidistant from the anodes. Position the anodes on at least four sides when they are used in a production tank or use a concentric anode arrangement. Do not place the spiral contractometer between the tank anodes and the work being plated in a production tank. A separate ammeter and current control are required. If the test is run on a sample of the solution, use a 4-L beaker.

8.3 When testing autocatalytic plating solutions, maintain the proper surface area-to-solution volume ratio (see 6.5).

8.4 The conditions of the test are usually chosen to match those of the production parts as closely as practicable.

8.4.1 The thickness of the deposit plated on the helix may not be required to be the same as the thickness plated on production parts. This is especially true when production part thickness requirements are greater than about 15 μ m.

8.4.2 The current density used for the spiral contractometer is sometimes lower than that of production parts. Electroplating at the lower current density will often indicate the cumulative effects of solution impurities before these have much effect on the work being plated at the higher currents.

³ Manufactured by Michigan Chrome and Chemical Co., 8615 Grinell Ave., Detroit, MI 48174.