



Designation: ~~D5145-03~~ Designation: D 5145 - 09

Standard Test Methods for Nonvolatile and Pigment Content of Electrocoat Baths¹

This standard is issued under the fixed designation D 5145; 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 These test methods cover the characterization of electrocoat baths through the determination of nonvolatile content of inorganic pigment content.

1.2

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

1.3 *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 and health practices and determine the applicability of regulatory limitations prior to use.*

2. Referenced Documents

2.1 *ASTM Standards:*²

D 1193 [Specification for Reagent Water](#)

D 2832 [Guide for Determining Volatile and Nonvolatile Content of Paint and Related Coatings](#)

E 180 [Practice for Determining the Precision of ASTM Methods for Analysis and Testing of Industrial Chemicals and Specialty Chemicals](#)

3. Summary of Test Method

3.1 Two specimens are accurately weighed into aluminum weighing dishes. The dishes are placed in an oven at 110°C for 1 h, reweighed to obtain the nonvolatile matter content and, if required, placed in a muffle furnace at 500°C for 2 h and weighed a third time to obtain the inorganic pigment content.

4. Significance and Use

4.1 The nonvolatile content and pigment content are measures of total solids and inorganic pigment solids, respectively, in electrocoat paints. In addition to production quality control, these properties are important in maintaining electrocoat baths in the optimum range.

4.2 Other test methods for determining nonvolatile content of paint and paint related materials are described in Guide D 2832.

5. Apparatus

5.1 *Analytical Balance* with a sensitivity of 0.1 mg.

5.2 *Aluminum Weighing Dishes*, 57 mm in diameter and 17 mm deep. These commercial dishes may contain a lubricant used during their manufacture. This should be removed by heating the aluminum dishes on a hot plate at 300°C until vapors are no longer visible. Store the dishes in a desiccator until needed.

5.3 *Syringes*, 5-mL, disposable variety.

5.4 *Oven* circulating, maintained at $110 \pm 2^\circ\text{C}$.

5.5 *Muffle Furnace*, maintained at $500 \pm 15^\circ\text{C}$.

6. Reagents

6.1 *Purity of Water*—References to water shall be understood to mean water conforming to Type II of Specification D 1193.

¹ These test methods are under the jurisdiction of ASTM Committee D01 on Paint and Related Coatings, Materials, and Applications and are the direct responsibility of Subcommittee D01.21 on Chemical Analysis of Paints and Paint Materials.

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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.

*A Summary of Changes section appears at the end of this standard.

7. Sampling and Sample Preparation

7.1 Obtain the sample while the electrocoat bath is under proper circulation so a uniform sample is obtained. In the case of a ultrafiltrate sample, the material should be thoroughly mixed or stirred prior to drawing the sample, thereby ensuring uniformity.

7.2 After sampling, prior to removing the test specimen, it is mandatory the sample be shaken or stirred until it is homogeneous and free of any settled material. This is particularly important if there is a delay between sampling the bath and performing this test procedure. The absence of settled material should be ascertained visually or by inserting a spatula and scraping the bottom of the container. Continue to shake or stir the sample until specimens are taken for measurement. *This Point is Very Important.*

NONVOLATILE CONTENT

8. Procedure

8.1 Weigh two aluminum dishes separately, each to 0.1 mg and record as W_1 .

8.2 Using a syringe, withdraw 1.0 to 1.5 mL of the well mixed sample, then quickly weigh the syringe to 0.1 mg, recording this weight as W_2 . Transfer the entire contents of the syringe into the aluminum dish. Reweigh the empty syringe to 0.1 mg and record as W_3 . In the case of ultrafiltrate clear liquids or of low solids paints, increase the specimen size to 5 mL and preheat at 60°C for 2 h. Duplicate this step with the second aluminum dish (8.1).

8.3 Add a few millilitres of water to the specimen in the aluminum dishes prior to placing them in the oven. This facilitates uniform spreading of the material. Place the dishes in the 110°C oven for 1 h.

8.4 Remove the dishes from the oven and allow to cool to room temperature in a desiccator. Reweigh them to 0.1 mg and record the weights as W_4 .

8.5 Retain the dishes for measurement of inorganic pigment content as detailed in a following section of these test methods.

9. Calculation

9.1 Calculate the percent nonvolatile content as follows:

$$\% NV = \frac{W_4 - W_1}{W_2 - W_3} \times 100 \quad (1)$$

where:

W_1 = weight of empty aluminum dish, g,

W_2 = weight of syringe filled with sample, g,

W_3 = weight of empty syringe, g, and

W_4 = weight of dish and contents after 1 h at 110°C, g.

10. Precision and Bias

10.1 Precision is based on an interlaboratory study in which the operators in each of ten laboratories analyzed in duplicate on 2 days, four different electrocoat-bath samples with nonvolatile contents ranging from 0.30 to 25.2 %. The results were analyzed statistically in accordance with Practice E 180. The interlaboratory coefficient of variance was 1.6 % at 3040 df and the interlaboratory coefficient of variation was 2.3 % at 2736 df. Based on these coefficients, the following criteria should be used for judging the acceptability of results at the 95 % confidence level: level.

10.1.1 *Repeatability*: Two results, each the mean of duplicate determinations obtained by the same operator on different days, should be considered suspect if they differ by more than 4.6 % relative.

10.1.2 *Reproducibility*: Two results, each the mean of duplicate determinations obtained by operators in different laboratories, should be considered suspect if they differ by more than 6.7 % relative.

10.2 Bias cannot be determined because there are no accepted standards for nonvolatile content of electrocoat baths.

INORGANIC PIGMENT CONTENT

11. Procedure

11.1 Place the weighing dishes used for the nonvolatile content determination in the muffle furnace at 500°C for 2 h.

11.2 After 2 h, remove the dishes from the muffle furnace, transfer them to a desiccator and allow to cool to room temperature. After cooling, remove dishes and weigh to 0.1 mg, recording these weights at W_5 .

NOTE 1—If an organic char is still present in the aluminum dish after 2 h in the muffle furnace, continue heating in the furnace until no char is present.

12. Calculation

12.1 Calculate the percent inorganic pigment as follows:

$$\% \text{ Inorganic Pigment (EC Bath)} = \frac{W_5 - W_1}{W_2 - W_3} \times 100 \quad (2)$$