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Standard Practice for Preparation of Contaminated Test Coupons for the Evaluation of Cleaning Agents¹

This standard is issued under the fixed designation G121; 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.

ε¹ NOTE—Editorial changes were made throughout in October 2015.

1. Scope

- 1.1 This practice describes the procedure for the preparation of single- and double-sided contaminated metallic-test coupons for the evaluation of cleaning agents. It is applicable for the evaluation of cleaning agents proposed for the cleaning of oxygen-enriched systems and components. oxygen equipment, pharmaceutical manufacturing equipment, and medical devices (see Test Method G122 and Guide G127). It also is applicable to other systems where contamination is a concern.
- 1.2 Several classes of <u>eontaminants_contaminants/residues</u> most likely to be found in <u>oxygen-enriched systems and components oxygen equipment, pharmaceutical manufacturing equipment, and <u>medical devices</u> are identified. However, if the user of this practice has identified contaminants not included in these classes, such identified contaminants may be substituted for the preparation of the test eoupons.coupons if appropriate for this test method.</u>
- 1.3 Preparation of nonmetallic substrates is not addressed, although similar methodology may be used. Solvent and cleaning agent compatibility with the nonmetallic substrates should be verified prior to the preparation of the test coupons. Typical nonmetallic materials utilized in oxygen systems are contained in Guide G63.
- 1.4 This practice may involve hazardous materials, operations, and equipment. This practice does not purport to address all of the safety concerns associated with its use. It is the responsibility of whomever uses this practice to consult and establish appropriate safety safety, health, and health environmental practices and determine the applicability of regulatory limitations prior to use.
- 1.5 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 a a/catalog/standards/sist/ed38ee50-5584-4578-881c-f132d83a3fd1/astm-g121-18
 - 2.1 ASTM Standards:²
 - D1193 Specification for Reagent Water
 - E1235 Test Method for Gravimetric Determination of Nonvolatile Residue (NVR) in Environmentally Controlled Areas for Spacecraft
 - E3106 Guide for Science-Based and Risk-Based Cleaning Process Development and Validation
 - F303 Practices for Sampling for Particles in Aerospace Fluids and Components
 - F312 Test Methods for Microscopical Sizing and Counting Particles from Aerospace Fluids on Membrane Filters
 - F324 Test Method for Nonvolatile Residue of Volatile Cleaning Solvents Using the Solvent Purity Meter (Withdrawn 1987)³
 - F331 Test Method for Nonvolatile Residue of Solvent Extract from Aerospace Components (Using Flash Evaporator)
 - F3127 Guide for Validating Cleaning Processes Used During the Manufacture of Medical Devices
 - G63 Guide for Evaluating Nonmetallic Materials for Oxygen Service
 - G94 Guide for Evaluating Metals for Oxygen Service
 - G122 Test Method for Evaluating the Effectiveness of Cleaning Agents

¹ This practice is under the jurisdiction of ASTM Committee G04 on Compatibility and Sensitivity of Materials in Oxygen Enriched Atmospheres and is the direct responsibility of Subcommittee G04.02 on Recommended Practices.

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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 The last approved version of this historical standard is referenced on www.astm.org.



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2.2 ANSI:ANSI Standard:⁴

B 46.1B46.1 Surface Texture (Surface Roughness, Waviness, and Lay)

3. Terminology

- 3.1 Definitions:
- 3.1.1 blank, n—the contamination level of the fluid when the test coupon is omitted.

3.1.1.1 Discussion—

Sometimes referred to as "background" level.

- 3.1.2 *cleaning agent, n*—an agent used to support the removal of contaminant from equipment surfaces or other critical objects (such as a medical device).
- 3.1.3 <u>contaminant, contaminant (contamination), n</u>—unwanted <u>molecular and molecular, non-volatile residue (NVR), and/or</u> particulate matter that could <u>adversely</u> affect or degrade the <u>performance operation, life, or reliability</u> of the <u>systems or components</u> upon which they reside.

3.1.3.1 Discussion—

The contaminant could be a drug substance, an intermediate, a formulated product, a lubricant, a machining oil, a solvent, a cleaning agent, etc. The contaminant may migrate to other surfaces or users may become exposed to the contaminant during the use of equipment.

- 3.1.4 *contamination*, <u>contaminate</u>, <u>n—v—a</u> process of contaminating. <u>applying contaminant (non-volatile residue (NVR) and/or particulate matter).</u>
- 3.1.3 surface roughness, R_{α} n—the arithmetic average deviation of the surface profile from the centerline, normally reported in micrometres.
- 3.1.4 nonvolatile residue (NVR), n—residual molecular and particulate matter remaining following the filtration of a solvent containing contaminants and evaporation of the solvent at a specified temperature.
- 3.1.5 particle (particulate contaminant), control coupon (witness coupon), n—a piece of matter in a solid state with observable length, width, and thickness coupon made from the same material as the test coupons, but in this test method is not coated with the contaminant.

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3.1.5.1 Discussion os. iteh.ai/catalog/standards/sist/ed38ee50-5584-4578-881c-f132d83a3fd1/astm-g121-18

The size of a particle is usually defined by its great dimension and is specified in micrometres.

- 3.1.6 dirty hold time (DHT), n—the maximum time interval between equipment use and cleaning.
- 3.1.7 *molecular contaminant (nonparticulate*(non-particulate contamination), n—the molecular contaminant contaminants that may be exist in a gaseous, liquid, or solid state.state and may be uniformly or non-uniformly distributed.

3.1.7.1 Discussion—

It may be uniformly or nonuniformly distributed, or be Molecular contaminants may be found as a solution, an emulsion, or in the form of droplets. Molecular contaminants account for most of the what constitutes NVR.

3.1.8 blank, nonvolatile residue (NVR), n—the contamination level of the fluid when the test coupon is omitted: a molecular and particulate matter remaining following the filtration and controlled evaporation of a solvent containing contaminants.

3.1.7.1 Discussion—

Sometimes referred to as "background" level.

3.1.9 *control coupon (witness coupon)*, *particle (particulate contaminant)*, *n*—a coupon made from the same material as the test eoupons, but in this test method is not coated with the contaminant.general term used to describe a finely divided solid of organic or inorganic matter with observable length, width, and thickness.

⁴ Dusts of varying particle sizes are available from the AC Spark Plug Division of General Motors, 1300-T N. Dort Hwy., Flint, MI 48556: Available from American National Standards Institute (ANSI), 25 W. 43rd St., 4th Floor, New York, NY 10036, http://www.ansi.org.

3.1.9.1 Discussion—

Particulates are usually reported as the amount of contaminant by the population of a specific micrometer size, usually defined by its greatest dimension. See methods described in Test Methods F312, Practices F303, or ARP 598 for particle size and population determination.

- 3.1.10 surface roughness, R_{α} n—the arithmetic average deviation of the surface profile from the centerline, normally reported in micrometres.
- 3.1.11 *test coupon*, *n*—representative surface that is typically a rectangular piece of a material of construction on which a known amount of a compound is deposited to simulate a process residue.

4. Summary of Practice

- 4.1 A solution of the contaminant The contaminant under study is applied to either one side or both sides of the precleaned test coupons and dried under standard conditions. in a defined area. The amount of contaminant on the test coupons is determined determined by weighing the test coupons before and after application. The amount of contaminant applied should be controlled and the variation in weight between test coupons minimized. Test coupons should be tested in a manner that simulates the actual cleaning conditions for the product, equipment, or system. Coupons may be tested immediately to simulate actual cleaning conditions or dried under specified conditions (for example, ambient overnight, 8 h at 104 °C, etc.) prior to testing (for example, for Dirty Hold Time studies). Nonmetallic material test coupons used as inserts, seats, seals, gaskets, etc. may also be prepared by this procedure and are evaluated in the as-used condition under actual cleaning conditions.
 - 4.2 Three methods of test coupon preparation are used:

Method A,A: NVR sample, single side,

Method B,B: NVR sample, double side side, and

Method C,C: NVR and particulate samplesample.

5. Significance and Use

- 5.1 This practice will be suitable to direct the preparation of test coupons with a known amount of contaminant on the surface. A standard test coupon is described and a list of contaminants that have typically been found in oxygen-enriched systems and components is provided.
- 5.2 These <u>test</u> coupons shall be used in the evaluation of cleaning agents for oxygen-enriched systems and components. This will permit direct comparison within and between test facilities.
- 5.3 Materials used in other fluid handling systems such as nitrogen, helium, hydrogen, gasoline, etc. may also be prepared for evaluation by this practice.

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6. Apparatus

6.1 Test Coupon—Metal panels Panels of the same material as the component part of construction and finish as the equipment, product, or system to be cleaned. Other alloys that may be used if the specific alloy is unknown are included in Guide G94. The coupon Test coupons should be numbered and in a manner that prevents removal during testing. A specified area of the test coupon may be designated as the test area where the test material is applied. Test coupons may be of different test area dimensions (for example, 5 cm x 5 cm, 10 cm x 10 cm) depending on the test procedure they are used in. An example test coupon configuration is shown in Fig. 1.

Note 1—The surface finish of the test coupon should be the same as the part to be cleaned.

- 6.2 Balance-Range to a minimum of 50 g with an 0.1-mg accuracy capable of weighing to ±0.1 mg.
- 6.3 Oven-Convection, Convection Oven—eapableCapable of maintaining 50°C ± 5°C.50 °C ± 5 °C.
- 6.4 Spray Applicator—Capable of applying an even coat of contaminant; that is, an artist's airbrush, perfume atomizer, or a spray device such as that used with window or tile cleaners has been found to apply an even coating of the contaminant in a controlled manner.
- 6.5 Other standard equipment such as a vacuum filtration system, solvent resistant filters, gloves, laboratory glassware, syringes, pipettes, desiccator, laboratory tongs, tweezers, and wire.

7. Reagents

7.1 <u>Contaminant Materials—Contaminant materials-general classes of materials that have typically been found in oxygen-enriched systems and components as a result of the manufacturing, assembly, fabrication, and construction processes include:</u>

silicone oils and greases,

fluorinated aerospace fluids and greases,