



Association of Textile, Apparel & Materials Professionals

AATCC takes no position respecting the validity of any patent rights asserted in connection with any item mentioned in this test method. Users of this test method are expressly advised that determination of the validity of any such patent rights, and the risk of infringement of such rights, are entirely their own responsibility.

This test method is subject to revision at any time by the responsible technical committee and must be reviewed every five years and if not revised, either reaffirmed or withdrawn. Your comments are invited either for revision of this test method or for additional procedures and should be addressed to the AATCC Technical Center. Your comments will receive careful consideration at a meeting of the responsible technical committee, which you may attend. If you feel that your comments have not received a fair hearing, you should make your views known to Diana A. Wyman, Technical Director, at the address shown below.

This test method is copyrighted by AATCC, PO Box 12215, Research Triangle Park, NC 27709 USA. Individual reprints (single or multiple copies) of this method may be obtained by contacting AATCC at the above address or tel: 919-549-3526; fax: 919-549-8933, or e-mail: orders@aatcc.org.

AATCC License Agreement: This test method is copyrighted by the American Association of Textile Chemists and Colorists (AATCC), PO Box 12215, 1 Davis Drive, Research Triangle Park, NC USA. All rights reserved.

AATCC grants you a license as follows: The right to download one electronic file of this AATCC test method for temporary storage on one computer for purposes of viewing, and/or printing one copy of the AATCC test method for individual use. Neither the electronic file nor the hard copy print may be reproduced in any way. In addition, the electronic file may not be distributed elsewhere over computer networks or otherwise. The hard copy print may only be distributed to other employees for their internal use within your organization.

This test method is not for resale.

Thermal Fixation Properties of Disperse Dyes

Developed in 1978 by AATCC Committee RA87; reaffirmed 1981, 2006, 2011; editorially revised and reaffirmed 1986, 1991, 1996, 2001; editorially revised 2008, 2010.

1. Purpose and Scope

1.1 This test method is used to determine the fixation properties of disperse dyes on polyester/cellulose blends as a function of fixation conditions. The variable described herein is temperature; however, variations in time and/or concentration of dyes and/or auxiliaries may be studied by this method.

2. Principle

2.1 The dye is padded onto the blend fabric at a specified concentration; the fabric is dried and the dye is fixed by exposure to controlled conditions of time and temperature. The cotton portion of the blend is dissolved in concentrated sulfuric acid followed by neutralization and thorough rinsing.

2.2 The reflectance values of the dyed fabrics, exposed to the several fixation conditions, are determined by spectrophotometry and the concentrations of dye relative to the heaviest dyeing (as 100%) are evaluated by applying the Kubelka-Munk function (K/S). These results express the relative dye fixation for a particular set of dyeing conditions. Alternatively, the dye on the fabric can be extracted with a suitable solvent and the concentrations of dye determined by transmittance spectrophotometry. When the dye concentrations on the fixed samples are compared to the corresponding concentration on a padded but *unfixed* sample, a measure of the true dye fixation, as a function of applied dye, may be obtained.

3. Terminology

3.1 **disperse dye**, *n.*—an essentially water insoluble dye having affinity, when properly dispersed, for polyester, polyamide and some other manufactured polymeric fibers.

3.2 **thermal fixation**, *n.*—the use of dry heat to achieve a degree of permanence when applying colorants to textile materials.

4. Safety Precautions

NOTE: These safety precautions are for information purposes only. The pre-

cautions are ancillary to the testing procedures and are not intended to be all inclusive. It is the user's responsibility to use safe and proper techniques in handling materials in this test method. Manufacturers MUST be consulted for specific details such as material safety data sheets and other manufacturer's recommendations. All OSHA standards and rules must also be consulted and followed.

4.1 Good laboratory practices should be followed. Wear safety glasses in all laboratory areas.

4.2 All chemicals should be handled with care.

4.3 Perform leaching procedure (see 7.3) using sulfuric acid, 70%, in a hood. In preparing, dispensing and handling sulfuric acid, use chemical goggles or face shield, impervious gloves and an impervious apron. Concentrated acids should be handled only in an adequately ventilated hood. CAUTION: Always add acid to water.

4.4 In preparing, dispensing and handling ammonium hydroxide, use chemical goggles or face shield, impervious gloves and an impervious apron. Dispense, mix and handle ammonium hydroxide only in an adequately ventilated hood.

4.5 Perform the extraction procedure (see 8.2) using chlorobenzene inside an adequately ventilated hood. CAUTION: Chlorobenzene vapors are toxic and flammable.

4.6 Chlorobenzene is a flammable liquid and should be stored in the laboratory only in small containers away from heat, open flame and sparks.

4.7 An eyewash/safety shower should be located nearby and a self-contained breathing apparatus should be readily available for emergency use.

4.8 Exposure to chemicals used in this procedure must be controlled at or below levels set by governmental authorities (e.g., Occupational Safety and Health Administration's [OSHA] permissible exposure limits [PEL] as found in 29 CFR 1910.1000; see web site: www.osha.gov for latest version). In addition, the American Conference of Governmental Industrial Hygienists (ACGIH) Threshold Limit Values (TLVs) comprised of time weighted averages (TLV-TWA), short term exposure limits (TLV-STEL) and ceiling limits (TLV-C) are recommended as a general guide for air contaminant exposure which should be met (see 11.1).

5. Uses and Limitations

5.1 Evaluations by reflectance measurements are limited to relatively low dye concentrations because of the non-linearity of the K/S function at high dye concentrations (see 11.2).

5.2 Fixation values obtained by reflectance measurements can differ from those determined by extraction techniques because of differences in dye distribution within the fiber (see 11.2).

5.3 Every effort should be made during padding, drying and fixation to avoid two-sided dyeings. Should this occur to any significant extent, the preferred method of measuring fixation is extraction.

6. Apparatus and Materials (see 11.3)

- 6.1 Laboratory padder.
- 6.2 Forced air thermofixation unit (see 11.4).
- 6.3 Analytical balance.
- 6.4 Spectrophotometer (see 11.5).
- 6.5 Volumetric pipettes and flasks.
- 6.6 65/35 polyester/cotton fabric; adequately pre-scoured and ready for dyeing.
- 6.7 Sulfuric acid, 70%.
- 6.8 Chlorobenzene.
- 6.9 Ammonium hydroxide, 5%.
- 6.10 Acetic acid, 56%.

7. Procedure

7.1 Pad polyester/cotton blend cloth at 50-60% wet pickup with:

- 10 g/L—Disperse dye
- 20 g/L—Antimigrant (see 11.6)
- Acetic acid (56%) to pH 5.5-6.0

Use the same pad liquor for all the dyeings in the set.

7.2 Dry each sample thoroughly under minimum migration conditions (see 11.5), then thermofix for 90 s at 196°C (385°F), 205°C (400°F), 213°C (415°F) and 221°C (430°F), respectively, in a properly calibrated oven (see 11.4).

7.3 Leach out the cotton portion of the blend with 70% sulfuric acid for 3-4 min at 55°C (130°F). Rinse the sample in cold running water and neutralize with 5% ammonium hydroxide solution for 1 min followed by another cold water rinse.

7.4 In cases where a change in shade of a disperse dye is suspected due to leaching, the following procedure may be employed: Attach a 100% polyester rider to the blended cloth to be dyed by the method in 7.1. Leach a portion of the rider separately by the method described in 7.3 and evaluate for any changes that