



Designation: D 5756 – 95

Standard Test Method for Microvacuum Sampling and Indirect Analysis of Dust by Transmission Electron Microscopy for Asbestos Mass Concentration¹

This standard is issued under the fixed designation D 5756; 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 covers a procedure to (a) identify asbestos in dust and (b) provide an estimate of the concentration of asbestos in the sampled dust, reported as either the mass of asbestos per unit area of sampled surface or as the mass of asbestos per mass of sampled dust.

1.1.1 If an estimate of asbestos structure counts is to be determined, the user is referred to Test Method D 5755.

1.2 This test method describes the equipment and procedures necessary for sampling, by a microvacuum technique, non-airborne dust for levels of asbestos. The non-airborne sample is collected inside a standard filter membrane cassette from the sampling of a surface area for dust which may contain asbestos.

1.2.1 This procedure uses a microvacuuming sampling technique. The collection efficiency of this technique is unknown. Variability of collection efficiency for any particular substrate and across different types of substrates is also unknown. The effects of sampling efficiency differences and variability on the interpretation of dust sampling measurements have not been determined.

1.3 Asbestos identified by transmission electron microscopy (TEM) is based on morphology, selected area electron diffraction (SAED), and energy dispersive X-ray analysis (EDXA). Some information about structure size is also determined.

1.4 This test method is generally applicable for an estimate of the concentration of asbestos starting from approximately 0.24 pg of asbestos per square centimeter (assuming a minimum fiber dimension of 0.5 μm by 0.025 μm , see 17.8), but will vary with the analytical parameters noted in 17.8.

1.4.1 The procedure outlined in this test method employs an indirect sample preparation technique. It is intended to disaggregate and disperse asbestos into fibrils and fiber bundles that can be more accurately identified, counted, and sized by transmission electron microscopy. However, as with all indirect sample preparation techniques, the asbestos observed for quantitation may not represent the physical form of the

asbestos as sampled. More specifically, the procedure described neither creates nor destroys asbestos, but it may alter the physical form of the mineral fibers.

1.5 The values stated in SI units are to be regarded as the standard. The values given in parentheses are for information only.

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

2. Referenced Documents

2.1 ASTM Standards:

- D 1193 Specification for Reagent Water²
- D 1739 Test Methods for Collection and Measurement of Dustfall (Settleable Particulate Matter)³
- D 3195 Practice for Rotameter Calibration³
- D 5755 Test Method for Microvacuum Sampling and Indirect Analysis of Dust by Transmission Electron Microscopy for Asbestos Structure Number Concentrations³
- E 832 Specification for Laboratory Filter Papers⁴

2.2 ISO Standards:

- ISO/10312 Ambient Air: Determination of Asbestos Fibers; Direct Transfer Transmission Electron Microscopy Procedure⁵
- ISO/CD13794 Ambient Air: Determination of Asbestos Fibers; Indirect-Transfer Transmission Electron Microscopy Procedure⁵

3. Terminology

3.1 Definitions:

3.1.1 *asbestiform*—a special type of fibrous habit in which the fibers are separable into thinner fibers and ultimately into fibrils. This habit accounts for greater flexibility and higher tensile strength than other habits of the same mineral. For more

¹ This test method is under the jurisdiction of ASTM Committee D-22 on Sampling and Analysis of Atmospheres and is the direct responsibility of Subcommittee D22.07 on Sampling and Analysis of Asbestos.

Current edition approved Aug. 15, 1995. Published October 1995.

² Annual Book of ASTM Standards, Vol 11.01.

³ Annual Book of ASTM Standards, Vol 11.03.

⁴ Annual Book of ASTM Standards, Vol 14.02.

⁵ Available from American National Standards Institute, 11 W. 42nd St., 13th Floor, New York, NY 10036.

information on asbestiform mineralogy, see references (1), (2) and (3).⁶

3.1.2 *asbestos*—a collective term that describes a group of naturally occurring, inorganic, highly fibrous silicate minerals, which are easily separated into long, thin, flexible fibers when crushed or processed.

3.1.2.1 *Discussion*—Included in the definition are the asbestiform varieties of: serpentine (chrysotile); riebeckite (crocidolite); grunerite (amosite); anthophyllite (anthophyllite asbestos); tremolite (tremolite asbestos); and actinolite (actinolite asbestos). The amphibole mineral compositions are defined according to the nomenclature of the International Mineralogical Association (3).

Asbestos	Chemical Abstract Service No. ⁷
Chrysotile	12001-29-5
Crocidolite	12001-28-4
Grunerite Asbestos (Amosite)	12172-73-5
Anthophyllite Asbestos	77536-67-5
Tremolite Asbestos	77536-68-6
Actinolite Asbestos	77536-66-4

3.1.3 *fibril*—a single fiber that cannot be separated into smaller components without losing its fibrous properties or appearance.

3.2 Definitions of Terms Specific to This Standard:

3.2.1 *aspect ratio*—the ratio of the length of a fibrous particle to its average width.

3.2.2 *bundle*—a structure composed of three or more fibers in a parallel arrangement with the fibers closer than one fiber diameter to each other.

3.2.3 *cluster*—an aggregate of two or more randomly oriented fibers, with or without bundles. Clusters occur as two varieties—disperse clusters and compact clusters.

3.2.3.1 *compact cluster*—a complex and tightly bound network in which one or both ends of each individual fiber or bundle are obscured, such that the dimensions of individual fibers or bundles cannot be unambiguously measured.

3.2.3.2 *disperse cluster*—a disperse and open network in which both ends of one of the individual fibers or bundles can be separately located and its dimensions measured.

3.2.4 *debris*—materials that are of an amount and size (particles greater than 1 mm in diameter as defined by a 1.0 by 1.0 mm screen) that can be visually identified (by color, texture, etc.) as to their source.

3.2.5 *dust*—any material composed of particles in a size range of ≤ 1 mm and large enough to settle by virtue of their weight from the ambient air. See Test Method D 1739.

3.2.6 *fiber*—a structure having a minimum length of 0.5 μ m with an aspect ratio of 5 to 1 or greater and substantially parallel sides (4). Fibers are assumed to have a cylindrical shape (5).

3.2.7 *fibrous mineral*—a mineral that is composed of parallel, radiating, or interlaced aggregates of fibers, from which the fibers are sometimes separable.

3.2.7.1 *Discussion*—The crystalline aggregate may be re-

ferred to as fibrous even if it is not composed of separable fibers, but has that distinct appearance. The term fibrous is used in a general mineralogical way to describe aggregates of grains that crystallize in a needle-like habit and appear to be composed of fibers. Fibrous has a much more general meaning than asbestos. While it is correct that all asbestos minerals are fibrous, not all minerals having fibrous habits are asbestos.

3.2.8 *indirect preparation*—a method in which a sample passes through one or more intermediate steps prior to final filtration.

3.2.9 *matrix*—a structure in which one or more fibers, or fiber bundles, touch, are attached to, or partially concealed by a single particle or connected group of non-fibrous particles. The exposed fiber must meet the fiber definition (see section 3.2.6). Matrices occur as two varieties—disperse matrices and compact matrices.

3.2.9.1 *compact matrix*—a structure consisting of a particle or linked group of particles, in which fibers or bundles can be seen either within the structure or projecting from it, such that the dimensions of individual fibers and bundles cannot be unambiguously determined.

3.2.9.2 *disperse matrix*—a structure consisting of a particle or linked group of particles, with overlapping or attached fibers or bundles in which at least one of the individual fibers or bundles can be separately identified and its dimensions measured.

3.2.10 *structures*—a term that is used to categorize all the types of asbestos particles which are recorded during the analysis (such as fibers, bundles, clusters, and matrices).

4. Summary of Test Method

4.1 The sample is collected by vacuuming a known surface area with a standard 25 or 37 mm air sampling cassette using a plastic tube that is attached to the inlet orifice which acts as a nozzle. The sample is transferred from inside the cassette to a 50/50 alcohol/water solution and screened through a 1.0 by 1.0 mm screen. The fine dust is filtered onto a membrane filter and ashed in a muffle furnace. The ash is mixed with distilled water to a known volume. Aliquots of the suspension are then filtered through a membrane. A section of the membrane is prepared and transferred to a TEM grid using the direct transfer method. The asbestiform structures are identified, sized, and counted by TEM, using SAED and EDXA at a magnification dependent on the size range of asbestos structures present.

5. Significance and Use

5.1 This microvacuum sampling and indirect analysis method is used for the general testing of non-airborne dust samples for asbestos. It is used to assist in the evaluation of dust that may be found on surfaces in buildings, such as ceiling tiles, shelving, electrical components, duct work, carpet, etc. This test method provides an estimate of the mass concentration of asbestos in the dust reported as either the mass of asbestos per unit area or as the mass of asbestos per mass of sampled dust as derived from a quantitative TEM analysis.

5.1.1 This test method does not describe procedures or techniques required to evaluate the safety or habitability of buildings with asbestos-containing materials, or compliance with federal, state, or local regulations or statutes. It is the

⁶ The boldface numbers refer to the list of references at the end of the test method.

⁷ The non-asbestiform variations of the minerals indicated in 3.1.2 have different Chemical Abstract Service (CAS) numbers.

user's responsibility to make these determinations.

5.1.2 At present, a single direct relationship between asbestos-containing dust and potential human exposure does not exist. Accordingly, the user should consider these data in relationship to other available information in their evaluation.

5.2 This test method uses the definition *settleable particulate matter* found in Test Method D 1739 as the definition of dust. This definition accepts all particles small enough to pass through a 1 mm screen. Thus, a single, large asbestos-containing particle(s) (from the large end of the particle size distribution) disassembled during sample preparation may result in anomalously large asbestos concentration results in the TEM analyses of that sample. Conversely, failure to disaggregate large particles may result in anomalously low asbestos mass concentrations. It is, therefore, recommended that multiple independent samples be secured from the same area, and that a minimum of three samples be analyzed by the entire procedure.

6. Interferences

6.1 The following minerals have properties (that is, chemical or crystalline structure) which are very similar to asbestos minerals and may interfere with the analysis by causing false positives to be recorded during the test. Therefore, literature references for these materials must be maintained in the laboratory for comparison to asbestos minerals so that they are not misidentified as asbestos minerals.

6.1.1 *Antigorite*.

6.1.2 *Palygorskite (Attapulgitite)*.

6.1.3 *Halloysite*.

6.1.4 *Pyroxenes*.

6.1.5 *Sepiolite*.

6.1.6 *Vermiculite scrolls*.

6.1.7 *Fibrous talc*.

6.1.8 *Hornblende* and other amphiboles not listed in 5.1.3.

6.2 Collection of any dust particles greater than 1 mm in size in this test method may cause an interference and, therefore, should be avoided.

7. Apparatus

7.1 *Transmission Electron Microscope (TEM)*, an 80 to 120 kV TEM, capable of performing electron diffraction, with a fluorescent screen inscribed with calibrated gradations, is required. The TEM must be equipped with energy dispersive X-ray spectroscopy (EDXA) and it must have a scanning transmission electron microscopy (STEM) attachment or be capable of producing a spot size of less than 250 nm in diameter at crossover.

7.2 *Energy Dispersive X-ray System (EDXA)*.

7.3 *High Vacuum Carbon Evaporator*, with rotating stage.

7.4 *High Efficiency Particulate Air (HEPA)*, filtered negative flow hood.

7.5 *Exhaust or Fume Hood*.

7.6 *Particle-Free Water* (ASTM Type II, see Specification D 1193).

7.7 *Glass Beakers*, 50 mL.

7.8 *Glass Sample Containers*, with wide mouth screw cap (200 mL), or equivalent sealable container (height of the glass

sample container should be approximately 13 cm high by 6 cm wide).

7.9 *Waterproof Markers*.

7.10 *Forceps (tweezers)*.

7.11 *Ultrasonic Bath*, table top model (100 W, approximate, see 22.5).

7.12 *Graduated Pipettes*, 1, 5, and 10 mL sizes, glass or plastic.

7.13 *Filter Funnel*, 25 mm or 47 mm (either glass or disposable). Filter funnel assemblies, either glass or disposable plastic, and using either a 25 mm or 47 mm diameter filter.

7.14 *Side Arm Filter Flask*, 1000 mL.

7.15 *Mixed Cellulose Ester (MCE) Membrane Filters*, 25 or 47 mm diameter, $\leq 0.22 \mu\text{m}$ and 5 μm pore size.

7.16 *Polycarbonate (PC) Filters*, 25 or 47 mm diameter, $\leq 0.2 \mu\text{m}$ pore size.

7.17 *Storage Containers*, for the 25 or 47 mm filters (for archiving).

7.18 *Glass Slides*.

7.19 *Scalpel Blades*.

7.20 *Cabinet-type Desiccator*, or low temperature drying oven.

8. Reagents and Materials

8.1 *Purity of Reagents*—Reagent grade chemicals shall be used in all tests. Unless otherwise indicated, it is intended that all reagents conform to the specifications of the Committee on Analytical Reagents of the American Chemical Society where such specifications are available.⁸ Other grades may be used, provided it is first ascertained that the reagent is of sufficiently high purity to permit its use without lessening the accuracy of the determination.

8.2 *Acetone*.

8.3 *Dimethylformamide (DMF)*.

8.4 *Chloroform*.

8.5 *1-methyl-2-pyrrolidone*.

8.6 *Glacial Acetic Acid*.

8.7 *Low Temperature Plasma Asher*.

8.8 *pH Paper*.

8.9 *Air Sampling Pump* (low volume personal-type pump).

8.10 *Rotameter*.

8.11 *Air Sampling Cassettes* (25 mm or 37 mm), containing 0.8 μm or smaller pore size MCE or PC filters.

8.12 *Cork Borer*, 7 mm.

8.13 *Non-Asbestos Mineral References*, as outlined in 6.1.

8.14 *Asbestos Standards*, as outlined in 3.1.2.

8.15 *Tygon*⁹ *Tubing*, or equivalent.

8.16 *Vacuum Pump* (small), that can maintain a vacuum of approximately 92 kPa.

8.17 *Petri Dishes*, large, glass (approximately 90 mm in diameter).

⁸ *Reagent Chemicals, American Chemical Society Specifications*, American Chemical Society, Washington, DC. For suggestions on the testing of reagents not listed by the American Chemical Society, see *Analar Standards for Laboratory Chemicals*, BDH Ltd., Poole, Dorset, U.K., and the *United States Pharmacopeia and National Formulary*, U.S. Pharmaceutical Convention, Inc. (USPC), Rockville, MD.

⁹ Tygon is a registered trademark of the DuPont Co.

- 8.18 *Stainless Steel or Aluminum Mesh Screen*, 30 to 40 mesh for Jaffe washer.
- 8.19 *Copper TEM Finder Grids*, 200 mesh.
- 8.20 *Carbon Evaporator Rods*.
- 8.21 *Lens Tissue*.
- 8.22 *Ashless Filter Paper*, 90 mm diameter, ASTM Class G (see Specification E 832).
- 8.23 *Reinforcement Rings*, gummed paper.
- 8.24 *Wash Bottles*, plastic.
- 8.25 *Reagent Alcohol*, HPLC grade (Fisher A995 or equivalent).
- 8.26 *Opening Mesh Screen*, plastic, 1.0 by 1.0 mm (Spectra-Mesh No. 146410 or equivalent).
- 8.27 *Static Neutralizer*.
- 8.28 *Muffle Furnace*, 480°C.
- 8.29 *Glazed Crucibles*.
- 8.30 *Jaffe Washer (4, 5)*.
- 8.31 *Diffraction Grating Replica*, with approximately 2160 lines/mm.

8.32 The following items are necessary if performing weight % calculations.

8.32.1 *Analytical Balance*, with readability of 0.01 mg or less. Exercise special care for the proper zeroing of the balance.

8.32.2 *Weighing Room*, with temperature and humidity control to allow weighing with an analytical balance to ± 0.01 mg.

8.32.3 *Class M Weights*, for calibration of the analytical balance.

9. Sampling Procedure For Microvacuum Technique

9.1 For sampling asbestos-containing dust in either indoor or outdoor environments, commercially available cassettes must be used. Cassettes and sampling nozzles must be new and not previously used. Air monitoring cassettes containing 25 mm or 37 mm diameter mixed cellulose ester (MCE) or polycarbonate (PC) filter membranes with a pore size less than or equal to 0.8 μm are required. The number of samples collected depends upon the specific circumstances of the study.

9.2 Maintain a log of all pertinent sampling information and sampling locations.

9.3 Sampling pumps and flow indicators shall be calibrated using a certified standard apparatus or assembly (see Practice D 3195).

9.4 Record all calibration information (6).

9.5 Perform a leak check of the sampling system at each sampling site by activating the pump with the closed sampling cassette in line. Any air flow shows that a leak is present that must be eliminated before initiating the sampling operation.

9.6 Attach the sampling cassette to the sampling pump at the outlet side of the cassette with plastic tubing (see 8.15). The plastic tubing shall be long enough that sample areas can be reached without interference from the sampling pump. Attach a clean, approximately 25.4 mm long, piece of plastic tubing (6.35 mm internal diameter) directly to the inlet orifice. Use this piece of tubing as the sampling nozzle. Cut the sampling end of the tubing at a 45° angle as illustrated in Fig. 1. The exact design of the nozzle is not critical as long as some vacuum break is provided to avoid simply pushing the dust around on the surface with the nozzle rather than vacuuming it

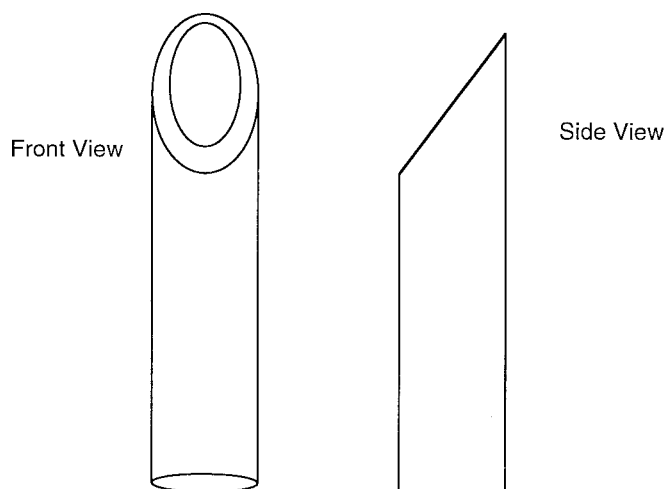


FIG. 1 Sampling End of Tubing

into the cassette. The internal diameter of the nozzle and flow rate of the pump may vary as long as the air velocity within the body of the nozzle is 100 (± 10) cm/s. This air velocity can be achieved with an internal sampling tube diameter of 6.35 mm ($\frac{1}{4}$ in.) and a flow rate of 2 L/min.

9.7 Delineate and measure the surface area of interest. A sample area of 100 cm^2 is vacuumed until there is no visible dust or particulate remaining. Perform a minimum of two orthogonal passes on the surface with a minimum of 2 min of sampling time. Avoid scraping or abrading the surface being sampled. (Do not sample any debris or dust particles greater than 1 mm in diameter (see 5.2)). Smaller or larger areas can be sampled, if needed. For example, some surfaces of interest may have a smaller area than 100 cm^2 . Less dusty surfaces may require vacuuming of larger areas. Unlike air samples, the overloading of the cassettes with dust will not be a problem. As defined in 3.2.5, only dust shall be collected for this analysis.

9.8 At the end of the sample collection, invert the cassette so that the nozzle inlet faces up before shutting off the power to the pump. The nozzle is then sealed with a cassette end plug and the cassette/nozzle taped or appropriately packaged to prevent separation of the nozzle and cassette assembly. A second option is the removal of the nozzle from the cassette, then plugging of the cassette and shipment of the nozzle (also plugged at both ends) sealed in a separate closeable plastic bag. A third option is placing the nozzle inside the cassette for shipment. The nozzle is always saved to be rinsed because a significant percentage of the dust drawn from a lightly loaded surface may adhere to the inside walls of the tubing.

9.9 Check that all samples are clearly labeled, that all dust sampling information sheets are completed, and that all pertinent information has been enclosed, in accordance with laboratory quality control practices, before transfer of the samples to the laboratory. Include an unused cassette and nozzle as a field blank.

9.10 Wipe off the exterior surface of the cassettes with disposable wet towels or “baby wipes” prior to packaging for shipment.

10. Sample Shipment

10.1 Ship dust samples to an analytical laboratory in a

sealed container, but separate from any bulk or air samples. The cassettes must be tightly sealed and packed in a material free of fibers or dust to minimize the potential for contamination. Plastic bubble pack is probably the most appropriate material for this purpose.

11. Sample Preparation

11.1 Under a negative flow HEPA hood, carefully wet-wipe the exterior of the cassettes to remove any possible contamination before taking cassettes into a clean preparation area.

11.2 Perform sample preparation in a clean facility that has a separate work area from both the bulk and air sample preparation areas.

11.3 Initial specimen preparation shall take place in a clean HEPA filtered negative pressure hood to avoid any possible contamination of the laboratory or personnel, or both, by the potentially large number of asbestos structures in an asbestos-containing dust sample. Cleanliness of the preparation area hoods is measured by the cumulative process blank concentrations (see Section 12).

11.4 If a weight % determination is reported, pre-weigh the laboratory filter in accordance with this section, otherwise, proceed with 11.5.

11.4.1 Dry the laboratory filter under vacuum in a vacuum desiccator for at least 15 min.

11.4.2 Release the vacuum, remove the desiccator cover, and equilibrate the laboratory filters in the weighing room for at least 1 h.

11.4.3 Weigh the laboratory filter and record the filter tare weight, W_1 (mg).

11.4.3.1 Zero the balance before each weighing.

11.4.3.2 Handle the filter with forceps.

11.4.3.3 Pass the filter over an antistatic radiation source. Repeat this step if the filter does not release easily from the forceps or if the filter attracts the balance pan.

11.4.4 Allow the laboratory filter to stand in the weighing room for an additional 8 to 16 h.

11.4.5 Reweigh the laboratory filter. If the second weight differs by more than 0.01 mg from the first tare weight (W_1), discard the filter.

11.5 All sample preparation steps 11.5.1-11.5.6 shall take place in the dust preparation area inside a HEPA hood.

11.5.1 Remove the upper plug from the sample cassette and container and carefully introduce approximately 10 mL of a 50/50 mixture of particle-free water and reagent alcohol into the cassette using a plastic wash bottle. If the plugged nozzle was left attached to the cassette, then remove the plug and introduce the water/alcohol into the cassette through the tubing, and then remove the tubing if it is visibly clean.

11.5.2 Replace the upper plug or the sample cap, and lightly shake the dust suspension by hand for 3 s.

11.5.3 Remove the entire cap of the cassette and pour suspension through a 1.0 by 1.0 mm opening screen into a pre-cleaned 200 mL glass specimen bottle. All visible traces of the sample contained in the cassette shall be rinsed onto the screen with a plastic wash bottle containing the 50/50 mixture of water and alcohol. Repeat this procedure two additional times for a total of three washings. Next, rinse the nozzle two or three times through the screen into the specimen bottle with

the 50/50 mixture of water and alcohol. Typically, the total amount of the 50/50 mixture used in the rinse is 50 to 75 mL. Remove the 1.0 by 1.0 mm screen and discard.

11.5.4 Use either a disposable plastic filtration unit or a glass filtering unit for filtration of the suspension. If a weight % determination is to be reported, use pre-weighed laboratory filters; see 11.4.

11.5.4.1 If a disposable plastic filtration unit is used, unwrap a new disposable plastic filter funnel unit (either 25 or 47 mm diameter) and remove the tape around the base of the funnel. Remove the funnel and discard the top filter supplied with the apparatus, while retaining the coarse polypropylene support pad in place. Assemble the unit with the adapter and a properly sized neoprene stopper, and attach the funnel to the 1000-mL side-arm vacuum flask. Place a 5.0 μm pore size MCE (backing filter) on the support pad. Wet it with a few mL of particle-free water and place an MCE or PC filter ($\leq 0.2 \mu\text{m}$ pore size) on top of the backing filter. Apply a vacuum, ensuring that the filters are centered and pulled flat without air bubbles. Any irregularities on the filter surface requires discard of that filter. After the filter has been seated properly, replace the funnel, and reseal it with the tape. Return the flask to atmospheric pressure.

11.5.4.2 If a glass filtration unit is used, place a 5 μm pore size MCE (backing filter) on the glass frit surface. Wet the filter with particle-free water, and place an MCE or PC filter ($\leq 0.2 \mu\text{m}$ pore size) on top of the backing filter. Apply a vacuum, ensuring that the filters are centered and pulled flat without air bubbles. Replace the filters if any irregularities are seen on the filter surface. Before filtration of the sample suspension, prepare a blank filter by filtration of 50 mL of particle-free water. After completion of the filtration, do not allow the filtration funnel assembly to dry because contamination is then more difficult to remove. Wash any residual suspension from the filtration assembly by holding it under a flow of water, then rub the surface with a clean paper towel soaked in a detergent solution. Repeat the cleaning operation, and then rinse several times in particle-free water.

11.5.5 Pour the suspension into the filter funnel. Rinse the beaker into the funnel at least three times, using distilled water.

11.5.6 Apply vacuum to the flask and draw the mixture through the filter.

11.5.7 Disassemble the filtering unit and carefully remove the sample filter with fine tweezers. Place the completed sample filter particle side up, into a pre-cleaned, labeled disposable plastic petri dish or other similar container.

11.6 If a weight % determination is reported, weigh the filter and solids using the method described in this section.

11.6.1 Place the filter into a low temperature drying oven (60°C) for 1 h.

11.6.2 Remove the filter and allow the filter to equilibrate in the weighing room for at least 1 h.

11.6.3 Weigh the filter and record the filter weight, W_2 (mg).

11.6.3.1 Zero the balance before each weighing.

11.6.3.2 Handle the filter with forceps.

11.6.3.3 Pass the filter over an antistatic radiation source. Repeat this step if the filter does not release easily from the forceps or if the filter attracts the balance pan.

11.6.3.4 Use the same balance for weighing the filters before and after filtration of the suspension.

11.7 Place the filter or a measured portion of the filter into a glazed crucible and cover with an appropriate lid.

11.7.1 Pre-heat the muffle furnace to 480°C. Regulate this temperature at $\pm 5^\circ\text{C}$ throughout the ashing of the sample.

11.7.2 Place the crucible in the muffle furnace. Ash the sample for 12 h, or until the weight has stabilized.

11.7.3 Remove the crucible from the muffle furnace and allow it to cool.

11.8 Rinse the ash from the crucible into a 200 mL glass specimen bottle. Rinse all visible traces of the sample contained in the crucible into the specimen bottle using a plastic wash bottle containing particle-free water. This procedure is repeated two additional times for a total of three washings.

11.8.1 Bring the volume of the suspension up to 100 mL using distilled water.

11.8.2 Adjust the pH of the suspension to 3 to 4 by using a 10.0 % solution of acetic acid. Use pH paper for testing. Filter the suspension within 24 h to avoid problems associated with bacterial and fungal growth.

11.9 Use either a disposable plastic filtration unit or a glass filtering unit for filtration of portions of the suspension. The ability of an individual filtration unit to produce a uniformly loaded filter may be tested by filtration of a colored particulate suspension such as carbon black.

11.9.1 If a disposable plastic filtration unit is used then unwrap a new disposable plastic filter funnel unit (either 25 or 47 mm diameter) and remove the tape around the base of the funnel. Remove the funnel and discard the top filter supplied with the apparatus, while retaining the coarse polypropylene support pad in place. Assemble the unit with the adapter and a properly sized neoprene stopper, and attach the funnel to the 1000 mL side arm vacuum flask. Place a 5.0 μm pore size MCE (backing filter) on the support pad. Wet it with a few mL of particle-free water and place an MCE or PC filter ($\leq 0.2 \mu\text{m}$ pore size) on top of the backing filter. Apply a vacuum, ensuring that the filters are centered and pulled flat without air bubbles. Any irregularities on the filter surface dictates discarding of that filter. After the filter has been seated properly, replace the funnel and reseal it with the tape. Return the flask to atmospheric pressure.

11.9.2 If a glass filtration unit is used, place a 5 μm pore size MCE (backing filter) on the glass frit surface. Wet the filter with particle-free water, and place an MCE or PC filter ($\leq 0.2 \mu\text{m}$ pore size) on top of the backing filter. Apply a vacuum, ensuring that the filters are centered and pulled flat without air bubbles. Replace the filters if any irregularities are seen on the filter surface. Before filtration of each set of sample portions, prepare a blank filter by filtration of 50 mL of particle-free water. If portions of the same sample are filtered in order of increasing concentration, the glass filtration unit need not be washed between filtrations. After completion of the filtration, do not allow the filtration funnel assembly to dry because contamination is then more difficult to remove. Wash any residual suspension from the filtration assembly by holding it under a flow of water, then rub the surface with a clean paper towel soaked in a detergent solution. Repeat the cleaning

operation, and then rinse several times in particle-free water.

11.9.3 With the flask at atmospheric pressure, add 20 mL of particle-free water into the funnel. Cover the filter funnel with its plastic cover if the disposable filtering unit is used.

11.10 Briefly hand shake (3 s) the capped bottle with the sample suspension, then place it in a tabletop ultrasonic bath and sonicate for 15 min. Maintain the water level in the sonicator at the same height as the solution in the sample bottle. Carry out preparation steps 11.10-11.11.5 in a HEPA hood.

11.10.1 Calibrate the ultrasonic bath in accordance with 22.5. The ultrasonic bath must be operated at equilibrium temperature.

11.11 Estimate the amount of liquid to be withdrawn to produce an adequate filter preparation. Experience has shown that a light staining of the filter surface will yield a suitable preparation for analysis. Filter at least 1.0 mL, but no more than half the total volume. Draw this portion from the suspension while it is in the sonicator. If after examination in the TEM, the smallest volume filtered (1.0 mL) yields an overloaded sample, perform additional serial dilutions of the suspension.

11.11.1 If serial dilutions are required, repeat step 11.10 before the serial dilution portion is taken. The recommended procedure for a serial dilution is to mix 10 mL of the sample solution with 90 mL of particle-free water in a clean sample bottle to obtain a 1:10 serial dilution. Follow good laboratory practices when performing serial dilutions.

11.11.2 Insert a new disposable pipette halfway into the sample suspension and withdraw a portion. Avoid pipetting any of the large floating or settled particles. Uncover the filter funnel and dispense the mixture from the pipette into the water in the funnel.

11.11.3 Apply vacuum to the flask and draw the mixture through the filter.

11.11.4 Discard the pipette.

11.11.5 Disassemble the filtering unit and carefully remove the sample filter with fine tweezers. Place the completed sample filter, particle side up, into a precleaned, labeled, disposable, plastic petri dish or other similar container.

11.11.6 In order to ensure that an optimally-loaded filter is obtained, it is recommended that filters be prepared from several different portions of the dust suspension. For this series of filters, it is recommended that the volume of each portion of the original suspension be a factor of five higher than the previous one. If the filters are prepared in order of increasing portion volume, all of the filters for one sample can be prepared using one plastic disposable filtration unit, or without cleaning of glass filtration equipment between individual filtrations.

11.11.7 There are many practical methods for drying MCE filters; the following are two examples that can be used: (1) dry MCE filters for at least 12 h (over desiccant) in an airtight cabinet-type desiccator; (2) to shorten the drying time (if desired), remove a plug of the damp filter and attach to a glass slide in accordance with 13.1.2 and 13.1.3. Place the slide with a filter plug or filter plugs (up to eight plugs can be attached to one slide) on a bed of desiccant, in the desiccator for 1 h.

11.11.8 PC filters do not require lengthy drying before

preparation, but may be placed in a desiccator for at least 30 min before preparation.

11.12 Prepare TEM specimens from small sections of each dried filter using the appropriate direct transfer preparation method.

12. Blanks

12.1 Prepare sample blanks that include both a process blank (50 mL of ultrapure water) for each set of samples analyzed and one unused filter from each new box of sample filters (MCE or PC). If glass filtering units are used, prepare and analyze a process blank each time the filtering unit is cleaned. For example, if one glass filtering unit is used for ten different individual samples, then ten process blanks are prepared and analyzed. Blanks will be considered contaminated if, after analysis, they are shown to contain more than 53 asbestos structures per square millimetre. This generally corresponds to three or four asbestos structures found in ten grid openings. The source of the contamination must be found before any further analysis can be performed. Reject samples that were processed along with the contaminated blanks and prepare new samples after the source of the contamination is found.

12.2 Prepare field blanks which are included with sample sets in the same manner as the samples, to test for contamination during the sampling, shipping, handling, and preparation steps of the method.

13. TEM Specimen Preparation of Mixed Cellulose Ester (MCE) Filters

NOTE 1—Use of either the acetone or the DMF-acetic acid method is acceptable.

13.1 Acetone Fusing Method:

13.1.1 Remove a section (plug) from any quadrant of the sample and blank filters. Sections can be removed from the filters using a 7 mm cork borer. The cork borer must be wet wiped after each time a section is removed.

13.1.2 Place the filter section (particle side up) on a clean microscope slide. Affix the filter section to the slide with a gummed page reinforcement or other suitable means. Label the slide with a glass scribing tool or permanent marker.

13.1.3 Prepare a fusing dish from a glass petri dish and a metal screen bridge. To prepare the fusing dish, make a pad from five to six ashless paper filters and place in the bottom of the petri dish. Place the screen bridge on top of the pad and saturate the filter pads with acetone. Place the slide on top of the bridge in the petri dish and cover the dish. Wait approximately 5 min for the sample filter to fuse and clear.

13.2 DMF-Acetic Acid Method:

13.2.1 Place a drop of clearing solution that consists of 35 % dimethylformamide (DMF), 15 % glacial acetic acid, and 50 % Type II water (v/v) on a clean microscope slide. Gauge the amount used so that the clearing solution just saturates the filter section.

13.2.2 Carefully lay the filter segment, sample surface upward, on top of the solution. Bring the filter and solution together at an angle of about 20° to help exclude air bubbles. Remove any excess clearing solution. Place the slide in an

oven or on a hot plate, in a fume hood, at 65 to 70°C for 10 min.

13.3 Plasma etching of the collapsed filter is required.

13.3.1 The microscope slide to which the collapsed filter pieces are attached is placed in a plasma asher. Because plasma ashers vary greatly in their performance, both from unit to unit and between different positions in the asher chamber, it is difficult to specify the exact conditions that must be used. Insufficient etching will result in a failure to expose embedded fibers, and too much etching may result in the loss of particles from the filter surface. To determine the optimum time for ashing, place an unused 25 mm diameter MCE filter in the center of a glass microscope slide. Position the slide approximately in the center of the asher chamber. Close the chamber and evacuate to a pressure of approximately 4 Pa, while admitting oxygen to the chamber at a rate of 8 to 20 cm³/min. Adjust the tuning of the system so that the intensity of the plasma is maximized. Determine the time required for complete oxidation of the filter. Adjust the system parameters to achieve complete oxidation of the filter in a period of approximately 15 min. For etching of collapsed filters, use these operating parameters for a period of 8 min. For additional information on calibration, see the AHERA (4), NISTIR (7), or ISO/10312 and ISO/CD 13794.

13.3.2 Place the glass slide containing the collapsed filters into a low-temperature plasma asher, and etch the filter.

13.4 Carbon coating of the collapsed and etched filters is required.

13.4.1 Carbon coating must be performed with a high-vacuum coating unit, capable of less than 10⁻⁴ torr (13 mPa) pressure. Units that are based on evaporation of carbon filaments in a vacuum generated only by an oil rotary pump have not been evaluated for this application and must not be used. Carbon rods used for evaporators shall be sharpened with a carbon rod sharpener to necks of about 4-mm length and 1-mm diameter. The rods are installed in the evaporator in such a manner that the points are approximately 100 to 120 mm from the surface of the microscope slide held in the rotating device.

13.4.2 Place the glass slide holding the filters on the rotation device, and evacuate the evaporator chamber to a pressure of at most 13 mPa. Perform the evaporation in very short bursts, separated by 3 to 4 s to allow the electrodes to cool. An alternate method of evaporation is accomplished by using a low continuous applied current. An experienced analyst can judge the thickness of the carbon film to be applied. Some tests should be made first on unused filters. If the carbon film is too thin, large particles will be lost from the TEM specimen, and there will be few complete and undamaged grid openings on the specimen. If the coating is too thick, it will lead to a TEM image that is lacking in contrast, and the ability to obtain electron diffraction patterns will be compromised. The carbon film should be as thin as possible and still remain intact on most of the grid openings of the TEM specimen.

13.5 Preparation of the Jaffe Washer:

13.5.1 The precise design of the Jaffe washer is not considered important, so any one of the published designs may be used (5, 8). One such washer consists of a simple stainless steel

bridge contained in a glass petri dish, as illustrated in Fig. 2.

13.5.2 Place several pieces of lens tissue on the stainless steel bridge. The pieces of lens tissue shall be large enough to completely drape over the bridge and into the solvent. In a fume hood, fill the petri dish with acetone (or DMF) until the height of the solvent is brought up to contact the underside of the metal bridge.

13.6 *Placing the Specimens into the Jaffe Washer:*

13.6.1 Place the TEM grids shiny side up on a piece of lens tissue or filter paper so that individual grids can be easily picked up with tweezers.

13.6.2 Prepare three grids from each sample.

13.6.2.1 Using a curved scalpel blade, excise at least two square (3 mm by 3 mm) pieces of the carbon-coated MCE filter from the glass slide.

13.6.2.2 Place the square filter piece carbon-side up on top of a TEM specimen grid.

13.6.2.3 Place the whole assembly (filter/grid) on the saturated lens tissue in the Jaffe washer.

13.6.2.4 Place the three TEM grid sample filter preparations on the same piece of lens tissue in the Jaffe washer.

13.6.2.5 Place the lid on the Jaffe washer and allow the system to stand for several hours.

13.7 Alternately, place the grids on a low level (petri dish filled to the $\frac{1}{8}$ mark) DMF Jaffe washer for 60 min. Add enough solution of equal parts DMF/acetone to fill the washer to the screen level. Remove the grids after 30 min if they have cleared, that is, all filter material has been removed from the carbon film, as determined by inspection in the TEM.

13.8 Carefully remove the grids from the Jaffe washer, allowing the grids to dry before placing them in a clean marked grid box.

14. TEM Specimen Preparation of Polycarbonate (PC) Filter

14.1 Cover the surface of a clean microscope slide with two strips of double-sided adhesive tape.

14.2 Cut a strip of filter paper slightly narrower than the width of the slide. Position the filter paper strip on the center of the length of the slide.

14.3 Using a clean, curved scalpel blade, cut a strip of the PC filter approximately 25 by 6 mm. Use a rocking motion of the scalpel blade to avoid tearing the filter. Place the PC strip particle side up on the slide perpendicular to the long axis of the slide. The ends of the PC strip must contact the double sided adhesive tape. Each slide can hold several PC strips. With a glass marker, label (next to each PC strip) with the individual sample number.

14.4 Carbon coat the PC filter strips as discussed in 13.4.2. PC filters do not require etching.

NOTE 2—**Caution:** Do not overheat the filter sections while carbon coating.

14.5 Prepare a Jaffe washer as described in 13.5, but fill the washer with chloroform or 1-methyl-2-pyrrolidone to the level of the screen.

14.6 Using a clean curved scalpel blade, excise three, 3 mm square filter pieces from each PC strip. Place the filter squares carbon side up on the shiny side of a TEM grid. Pick up the grid and filter section together and place them on the lens tissue in the Jaffe washer.

14.7 Place the lid on the Jaffe washer and leave the grids for at least 4 h. Best results are obtained with longer wicking times, up to 12 h.

14.8 Carefully remove the grids from the Jaffe washer. Allow the grids to dry before placing them in a clean, marked grid box.

15. Grid Opening Measurements

15.1 TEM grids must have a known grid opening area. Determine this area as follows:

15.1.1 Measure at least 20 grid openings in each of 20 random 75 μm (200-mesh) copper grids for a total of 400 grid openings for every 1000 grids used, by placing the 20 grids on a glass slide and examining them under the optical microscope. Use a calibrated graticule to measure the average length and width of the 20 openings from each of the individual grids. From the accumulated data, calculate the average grid opening area of the 400 openings.

15.2 Grid area measurements can also be made at the TEM at a calibrated screen magnification. Typically measure one grid opening for each grid examined. Measure grid openings in both the x and y directions and calculate the area.

15.3 Pre-calibrated TEM grids are also acceptable for this test method.

16. TEM Method

16.1 Calibrate the microscope routinely for magnification, camera constant, and EDXA settings (see Section 22 for more detailed discussion).

16.2 Microscope settings: 80 to 120 kV, 15 000 to 20 000 \times screen magnification for analysis as determined in 16.7.

16.3 Analyze two grids for each sample. Analyze approximately one-half of the required grid openings on one sample grid preparation and the remaining half on a second grid preparation. See 16.7 and 17.10 for the number of grid openings which must be analyzed.

16.4 *Determination of Specimen Suitability:*

16.4.1 Carefully load the TEM grid, carbon side facing up (in the TEM column) with the grid bars oriented parallel/

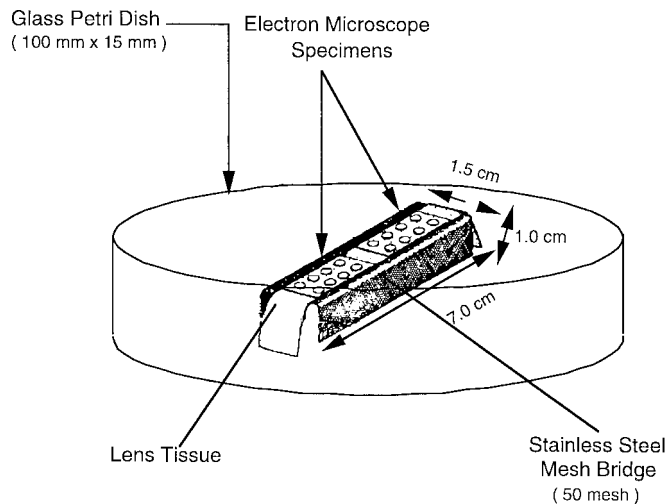


FIG. 2 Design of Solvent Washer (Jaffe Washer)

perpendicular to the length of the specimen holder. Use a hand lens or loupe, if necessary. This procedure will line up the grid with the *x* and *y* translation directions of the microscope. Insert the specimen holder into the microscope.

16.4.2 Scan the entire grid at low magnification (250× to 1000×) to determine its suitability for high magnification analysis as specified in 16.4.3.

16.4.3 Grids are acceptable for analysis if the following conditions are met:

16.4.3.1 The fraction of grid openings covered by the replica section is at least 50 %.

16.4.3.2 Relative to that section of the grid covered by the carbon replica, the fraction of intact grid openings is greater than 50 %.

16.4.3.3 The fractional area of undissolved filter is less than 10 %.

16.4.3.4 The fraction of grid openings with overlapping or folded replica film is less than 50 %.

16.4.3.5 At least 20 grid openings that have no overlapping or folded replica, are less than 5 % covered with holes and have less than 5 % opaque area due to incomplete filter dissolution.

16.5 *Determination of Grid Opening Suitability:*

16.5.1 If the grid meets acceptance criteria, choose a grid opening for analysis from various areas of the grid so that the entire grid is represented. Determine the suitability of each individual grid opening to be analyzed prior to the analysis.

16.5.2 The individual grid opening must have less than 5 % holes over its area.

16.5.3 Grid openings must be less than 25 % covered with particulate matter.

16.5.4 Grid openings must be uniformly loaded.

16.6 Observe and record the orientation of the grid, at 80 to 150×, on a grid map record sheet along with the location of the grid openings that are examined for the analysis. If indexed grids are used, a grid map is not required, but the identifying coordinates of the grid square must be recorded.

16.7 At low magnification (1000×), scan approximately 50 grid openings to find the largest width of an asbestos fiber or bundle. If an asbestos fiber or bundle is observed, adjust the microscope magnification so that 1 mm on the fluorescent screen corresponds to approximately 10 % of the width of the previously selected large structure. Examine the sample at the lower of this magnification or 20 000×. If no fibers or bundles are observed during the low magnification scan, arbitrarily choose one grid opening. Begin high magnification (15 000 to 20 000×) scans with this grid opening.

16.7.1 Record the dimensions of the structure with the largest width. Calculate the mass of this structure (see Section 17).

16.7.2 Continue scanning at the appropriate magnification until either the mass of the large structure selected in 16.7.1 is no greater than 1/10 of the total asbestos mass observed or the stopping rules (17.10) apply.

17. Data Recording Rules

17.1 Record on the count sheet (Fig. 3) any continuous grouping of particles in which an asbestos fiber is detected. Classify asbestos structures as fibers, bundles, clusters, or matrices as defined in 3.2.

17.2 Use the AHERA criteria for fiber, bundle, cluster, and matrix identification (4). Record length and width measurements.

17.2.1 For bundles and fibers, record the length and width of the structure.

17.2.2 For disperse clusters, record the overall length and width of the structure. Record, in order of decreasing length, up to five component fibers or bundles in accordance with 17.2.1. These component fibers or bundles are not counted in determining overall structure count or stopping rules.

17.2.3 For compact clusters, record the overall length and width of the structure.

17.2.4 For disperse matrices, record the overall length and width of the structure. Record, in order of decreasing length, up to five component fibers or bundles in accordance with 17.2.1. These component fibers or bundles are not counted in determining overall structure count or stopping rules.

17.2.5 For compact matrices, record the overall length and width of the structure.

17.2.6 Calculate the mass of an individual structure as shown in 20.1.

17.3 Record NSD (No Structures Detected) when no structures are detected in the grid opening.

17.4 Identify structures classified as chrysotile identified by either electron diffraction or X-ray analysis and record on a count sheet. Verify at least one out of every ten chrysotile structures by X-ray analysis.

17.5 Structures classified as amphiboles by X-ray analysis and electron diffraction are recorded on the count sheet. For more information on identification, see Yamate, et al (5) or Chatfield and Dillon (8).

17.6 Record a typical electron diffraction pattern for each type of asbestos observed for each group of samples (or a minimum of every five samples) analyzed. Record the micrograph number on the count sheet. Record at least one X-ray spectrum for each type of asbestos observed per sample. Attach the print-outs to the back of the count sheet. If the X-ray spectrum is stored, record the file and disk number on the count sheet.

17.7 *Counting Rules:*

17.7.1 At a screen magnification of between 15 000 and 20 000× evaluate the grids for the most concentrated sample loading; reject the sample if it is estimated to contain more than 50 asbestos structures per grid opening. Proceed to the next lower concentrated sample until a set of grids are obtained that have less than 30 asbestos structures per grid opening.

17.8 *Analytical Sensitivity*—An absolute analytical sensitivity specified in terms of mass cannot be defined for this test method. A nominal analytical sensitivity for an analysis can be defined on the basis of the smallest asbestos fibril reportable by this test method, 0.5 μm by 0.025 μm. Assuming the following values for the parameters defined in Section 19: EFA = 1150 mm²; GO = 10; GOA = 0.006 mm²; SPL = 100 cm²; V = 50 mL, this equates to an analytical sensitivity of 0.24 pg of asbestos per square centimetre. This sensitivity can be improved by increasing the amount of liquid filtered, increasing the number of grid openings analyzed, or decreasing the size of the final filter.

17.10.1.1 Complete the analysis of the grid square in which the stopping rules are satisfied.

17.10.2 If more than 10 % of the counted structures are either compact clusters or compact matrices, then the sample must be reprepared starting with 11.10, using either longer ultrasonic treatment or an ultrasonic probe to disperse the complex structures.

17.11 After analysis, remove the grids from the TEM, and replace them in the appropriate grid storage holder.

18. Sample Storage

18.1 The washed-out sample cassettes can be discarded after use.

18.2 Sample grids and unused filter sections must be stored for a minimum of one year.

19. Calculations

19.1 *Calculation of Results*—Use the following equation to determine the amount of asbestos in the sample:

$$\text{asbestos, g/cm}^2 = \frac{EFA \times 100 \text{ mL} \times M}{GO \times GOA \times V \times SPL \times FF} \quad (1)$$

where:

- M = total asbestos mass counted, g,
- EFA = effective filter area of the final sampling filter, mm²,
- GO = number of grid openings counted,
- GOA = average grid opening area, mm²,
- SPL = surface area sampled, cm²,
- V = volume of sample, mL, filtered in step 11.5.9, representing the actual volume taken from the original 100 mL suspension, and
- FF = fraction of filter ashed in 11.7.

19.1.1 If the mass of asbestos per mass of analyzed dust is to be reported, calculate the weight % as follows: Sum the mass of asbestos from 20.1 to obtain a total mass of asbestos. Estimate the weight % using the following equation:

$$\text{asbestos, weight percent} = \frac{EFA \times 100 \text{ mL} \times M \times 100 \%}{GO \times GOA \times V \times FF \times (W_2 - W_1)} \quad (2)$$

where:

- W_1 = pre-weight of the filter, 11.4.1, mg, and
- W_2 = weight of redeposit filter after the suspension has been filtered, as in 11.6.1, mg.

20. Reporting

20.1 Mass for each asbestos structure is calculated using the methods described in ISO/CD 13794.

20.1.1 The mass of a fiber or bundle is calculated as:

$$\text{Mass}_{\text{Fiber}}(\text{g}) = \frac{\pi}{4} \times L \times W^2 \times D \times 10^{-12} \quad (3)$$

where:

- L = length of the fiber, μm
- W = width of the fiber, μm, and
- D = density of fiber, Mg/m³ (1), (2), or (3).

20.1.1.1 The following densities are used:

Mineral	Density, Mg/m ³
Chrysotile	2.55
Crocidolite	3.37
Amosite	3.43
Anthophyllite	3.00
Tremolite	3.00
Actinolite	3.10

20.1.2 The mass of a disperse matrix or a disperse cluster is estimated by summing the mass of the component fibers or bundles. The mass of a compact cluster or compact matrix is not determined.

20.2 Report the following information for each dust sample analyzed:

- 20.2.1 Types of asbestos present,
- 20.2.2 Number of asbestos structures present,
- 20.2.3 Concentration in mass per cm² for each asbestos type present,
- 20.2.4 The analytical sensitivity,
- 20.2.5 Effective filtration area,
- 20.2.6 Average size of the TEM grid openings that were counted,
- 20.2.7 Number of grid openings examined,
- 20.2.8 Sample dilution used,
- 20.2.9 Area of the surface sampled,
- 20.2.10 Listing of size data for each structure counted, and
- 20.2.11 A copy of the TEM count sheet for each sample or a complete listing of the raw data.

21. Quality Control/Quality Assurance

21.1 In general, the laboratory's quality control checks are used to verify that a system is performing according to specifications regarding accuracy and consistency. In an analytical laboratory, spiked or known quantitative samples are normally used. However, due to the difficulties in preparing known quantitative asbestos samples, routine quality control testing focuses on re-analysis of samples (duplicate recounts).

21.1.1 Re-analyze samples at a rate of 1/10 of the sample sets (1 out of every 10 samples analyzed not including laboratory blanks). The re-analysis shall consist of a second sample preparation obtained from the final filter.

21.1.2 In addition, quality assurance programs must follow the criteria shown in the Federal Register for AHERA samples (4) and in the NISTIR document (7). These documents describe sample custody, sample preparation, blank checks for contamination, calibration, sample analysis, analyst qualifications, and technical facilities.

22. Calibrations

22.1 Perform calibrations of the instrumentation on a regular basis, and retain a calibration record for each TEM in the laboratory, in accordance with the laboratory's quality assurance program.

22.2 Record calibrations in a log book along with dates of calibration and the attached backup documentation.

22.3 A calibration list for the instrument is as follows:

22.3.1 *TEM*:

22.3.1.1 Check the alignment and the systems operation. Refer to the TEM manufacturer's operational manual for detailed instructions.

22.3.1.2 Calibrate the camera length of the TEM in electron

diffraction (ED) operating mode before ED patterns of unknown samples are observed. Camera length can be measured by using a carbon coated grid on which a thin film of gold has been sputtered or evaporated. A thin film of gold is evaporated on the specimen TEM grid to obtain zone-axis ED patterns superimposed with a ring pattern from the polycrystalline gold film. In practice, it is desirable to optimize the thickness of the gold film so that only one or two sharp rings are obtained on the superimposed ED pattern. Thicker gold films will tend to mask weaker diffraction spots from the fibrous particles. Since the unknown d-spacings of most interest in asbestos analysis are those which lie closest to the transmitted beam, multiple gold rings from thicker films are unnecessary. Alternatively, a gold standard specimen can be used to obtain an average camera constant calculated for that particular instrument that can then be used for ED patterns of unknowns taken during the corresponding period.

22.3.1.3 Perform magnification calibration at the fluorescent screen. This calibration must be performed at the magnification used for structure counting. Calibration is performed with a grating replica (for example, one containing at least 2160 lines/mm).

(a) Define a field of view on the fluorescent screen. The field of view must be measurable or previously inscribed with a scale or concentric circles (all scales should be metric).

(b) Frequency of calibration will depend on the service history of the particular microscope.

(c) Check the calibration after any maintenance of the microscope that involves adjustment of the power supply to the lens or the high-voltage system or the mechanical disassembly of the electron optical column (apart from filament exchange).

(d) The analyst must ensure that the grating replica is placed at the same distance from the objective lens as the specimen.

(e) For instruments that incorporate a eucentric tilting specimen stage, all specimens and the grating replica must be placed at the eucentric position.

22.3.1.4 The smallest spot size of the TEM must be checked.

(a) At the crossover point, photograph the spot size at a screen magnification of 15 000× to 20 000×. An exposure time of 1 s is usually adequate.

(b) The measured spot size must be less than or equal to 250 nm.

22.4 EDXA:

22.4.1 The resolution and calibration of the EDXA must be verified.

22.4.1.1 Collect a standard EDXA Cu peak from the Cu grid.

22.4.1.2 Compare the X-ray energy versus channel number for the Cu peak and be certain readings are within ±10 eV.

22.4.2 Collect a standard EDXA of crocidolite asbestos (NIST SRM 1866).

22.4.2.1 The elemental analysis of the crocidolite must resolve the Na peak.

22.4.3 Collect a standard EDXA of chrysotile asbestos.

22.4.3.1 The elemental analysis of chrysotile must resolve both Si and Mg on a single chrysotile fiber.

22.5 Perform ultrasonic bath calibration as follows:

22.5.1 Fill the bath with water to a level equal to the height of suspension in the glass sample container that is used for the dust. Operate the bath until the water reaches the equilibrium temperature.

22.5.2 Place 100-mL of water (at approximately 20°C) in the 200-mL glass sample container, and record its temperature.

22.5.3 Place the sample container into the water in the ultrasonic bath (with the power turned off). After 60 s, remove the glass container and record its temperature.

22.5.4 Place 100-mL of water (at approximately 20°C) in another 200-mL glass sample container, and record its temperature.

22.5.5 Place the second sample container in the water in the ultrasonic bath (with the power turned on). After 60 s, remove the glass container and record its temperature.

22.5.6 Calculate the rate of energy deposition into the sample container using the following formula:

$$R = 4.185 \times \sigma \times \rho \times \frac{\theta_2 - \theta_1}{t} \quad (4)$$

where:

4.185 = joules/cal,

R = energy deposition, MW/m³,

θ_1 = temperature rise with the ultrasonic bath not operating, °C,

θ_2 = temperature rise with the ultrasonic bath operating, °C,

t = time, s (to 60 s (22.5.3 and 22.5.5)),

σ = specific heat of the liquid in the glass sample container (1.0 cal/g), and

ρ = density of the liquid in the glass sample container (1.0 g/cm³).

22.5.7 Adjust the operating conditions of the bath so that the rate of energy deposition is in the range of 0.08 to 0.12 MW/m³, as defined by this procedure.

23. Precision and Bias

23.1 *Precision*—The precision of the procedure in this test method has not been determined. The failure to adequately disaggregate large structures, however, will decrease the precision of the results.

23.2 *Bias*—Bias of this test method has not been determined. The failure to adequately disaggregate large structures, however, will induce a negative bias to the results.

24. Keywords

24.1 asbestos; indirect; mass; microvacuuming; settled dust; TEM



D 5756

REFERENCES

- (1) Steel, E., and Wylie, A., "Mineralogical Characteristics of Asbestos," *Geology of Asbestos Deposits*, Riordon, P. H., Ed., SME-AIME, 1981, pp. 93–101.
- (2) Zussman, J., "The Mineralogy of Asbestos," *Asbestos: Properties, Applications and Hazards, Vol. 1*, Chissick, S. S., and Derricott, R., Eds., John Wiley and Sons, 1979, pp. 45–67.
- (3) Leake, B. E., "Nomenclature of Amphiboles," *American Mineralogist*, Vol 63, 1978, pp. 1023–1052.
- (4) "USEPA, Asbestos-Containing Materials in Schools: Final Rule and Notice," *Federal Register*, 40 CFR Part 763, Appendix A to Sub-part E., October 30, 1987.
- (5) Yamate, G., Agarwall, S. C., and Gibbons, R. D., "Methodology for the Measurement of Airborne Asbestos by Electron Microscopy," EPA Draft Report, Contract No. 68-02-3266, 1984.
- (6) *OSHA Technical Manual, OSHA Instruction CPL 2-20B*, Directorate of Technical Support, OSHA, U.S. Department of Labor, Washington, DC 20210. Feb. 5, 1990, pp. 1–8–1–11.
- (7) *NIST/NVLAP Program Handbook for Airborne Asbestos Analysis*, NISTIR, August 1989, pp. 89–4137.
- (8) Chatfield, E. J., and Dillon, M. J., "Analytical Method for the Determination of Asbestos in Water," EPA No. 600/4-83-043, 1983.

ASTM International takes no position respecting the validity of any patent rights asserted in connection with any item mentioned in this standard. Users of this standard 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 standard is subject to revision at any time by the responsible technical committee and must be reviewed every five years and if not revised, either reapproved or withdrawn. Your comments are invited either for revision of this standard or for additional standards and should be addressed to ASTM International Headquarters. 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 the ASTM Committee on Standards, at the address shown below.

This standard is copyrighted by ASTM International, 100 Barr Harbor Drive, PO Box C700, West Conshohocken, PA 19428-2959, United States. Individual reprints (single or multiple copies) of this standard may be obtained by contacting ASTM at the above address or at 610-832-9585 (phone), 610-832-9555 (fax), or service@astm.org (e-mail); or through the ASTM website (www.astm.org).