

Designation: D 38 – 94 (Reapproved 2000)^{€1}

Standard Test Methods for Sampling Wood Preservatives Prior to Testing¹

This standard is issued under the fixed designation D 38; the number immediately following the designation indicates the year of original adoption or, in the case of revision, the year of last revision. A number in parentheses indicates the year of last reapproval. A superscript epsilon (ϵ) indicates an editorial change since the last revision or reapproval.

 ϵ^1 Note—A keyword was added editorially in March 2000.

1. Scope

1.1 These test methods cover sampling of creosote, creosote-coal tar solutions, creosote-petroleum solutions, and oilborne and waterborne preservatives.

NOTE 1—See Specification D 390, for Coal-Tar Creosote for the Preservative Treatment of Piles, Poles, and Timbers for Marine, Land, and Fresh Water Use,² Specification D 391 for Creosote-Coal Tar Solution,² and Specification D 1858, for Creosote-Petroleum Solution.²

1.1.1 Creosote and creosote solutions are homogeneous liquids, except for their moisture content. At atmospheric temperatures, however, part of the creosote is usually in crystalline form, and shall be heated to a temperature at which it is entirely liquid before sampling.

1.1.1.1 When sampling creosote and creosote solutions, the object is to obtain a sample that will be representative of the water content, and the methods of sampling described are intended to accomplish that purpose.

1.1.2 Recommended test methods are given for sampling a preservative in a liquid or solid state and in the several types of containers in which they are likely to be encountered. Creosote, creosote-coal tar, and creosote-petroleum solutions may be in the form of a liquid or semiliquid. Oilborne and waterborne preservatives may be in the solid state, in concentrated solutions, or in ready-to-use solutions.

1.2 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 95 Test Method for Water in Petroleum Products and Bituminous Materials by Distillation²

- D 168 Test Method for Coke Residue of Creosote³
- D 246 Test Method for Distillation of Creosote and Creosote-Coal Tar Solutions³
- D 346 Practice for Collection and Preparation of Coke Samples for Laboratory Analysis⁴
- D 367 Test Method for Xylene-Insoluble Matter in Creosote³
- D 368 Test Method for Specific Gravity of Creosote and Oil-Type Preservatives³
- D 369 Test Method for Specific Gravity of Creosote Fractions and Residues³
- D 370 Test Method for Dehydration of Oil-Type Preservatives³
- D 1033 Methods of Chemical Analysis of Chromated Zinc Chloride³
- D 1035 Test Methods for Chemical Analysis of Fluor-Chrome-Arsenate-Phenol³
- D 1274 Test Methods for Chemical Analysis of Pentachlorophenol³
- D 1326 Methods for Chemical Analysis of Ammoniacal Copper Arsenate³
- D 1627 Methods for Chemical Analysis of Acid Copper Chromate³
- D 1628 Test Methods for Chemical Analysis of Chromated Copper Arsenate³
- D 3176 Practice for Ultimate Analysis of Coal and Coke⁴
- D 3180 Practice for Calculating Coal and Coke Analyses from As-Determined to Different Bases⁴

3. Significance and Use

3.1 The test methods described are for use in obtaining representative samples of wood preservatives under most phases of shipment, use, or storage. Since these materials may have to be sampled under varying conditions, it is not possible to provide a definite standard for each type of wood preservative under every sampling condition. Individual cases may have to be worked out by the sampler on the basis of judgement, skill, and experience. The basic principle of each

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² Annual Book of ASTM Standards, Vol 05.01.

³ Annual Book of ASTM Standards, Vol 04.10.

⁴ Annual Book of ASTM Standards, Vol 05.05.

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method, however, is to obtain a sample or a composite of several samples in such manner and from such locations in the container that the sample or composite will be truly representative of the material being sampled.

4. Apparatus

4.1 Continuous-Drip Sampling:

4.1.1 Line Connection constructed substantially as shown in Fig. 1 shall be inserted in the pipeline through which the preservative is being transferred, preferably on the discharge side of the pump. The sampling pipe shall be provided with a valve.

4.1.2 Receiver—Use a clean, dry container of convenient size to receive the sample. The container shall be covered with a lid to protect the sample from foreign contamination. Creosote, creosote-coal tar solutions, and creosote-petroleum solutions may become semisolid as they cool in the sample container during the course of the sampling. These samples should be heated to complete liquidity before taking an aliquot for testing.

4.2 Zone Sampling:

4.2.1 Sampling Bottle (see Figs. 2 and 3) with a 19-mm (3/4-in.) diameter opening.

4.3 Thief Sampling:

4.3.1 For sampling liquids or semiliquids in tank cars, tank trucks, and storage tanks, use a sampling devise designed so that a sample can be obtained within 13 mm (1/2in.) of the bottom of the vessel used. A sampling thief for this purpose is illustrated in Fig. 4.

4.3.2 For sampling granular material loose carried in bags, drums, or boxes, use a solids thief sampler 38 mm $(1\frac{1}{2} \text{ in.})$ in diameter and a least 457 mm (18 in.) long similar to that shown in Fig. 5.

4.4 Grab Sampling:

4.4.1 Shovel or Scoop, with a width at least twice as great as the maximum dimension of the largest pieces sampled and a sample container suitable for the product being sampled.

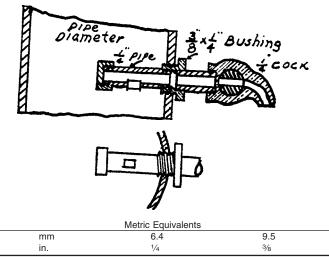


FIG. 1 Continuous Sampling Connection

4.5 *Tube Sampling*:

4.5.1 Glass or Metal Tube, designed so that it will reach to within 3 mm (1/8 in.) of the bottom of the container. A metal tube for sampling 208-L (55-gal) drums is shown in Fig. 6.

4.6 Tap Sampling:

4.6.1 Standard Pipe, with a suitable valve is satisfactory. The valve should be equipped with a delivery tube which will not contaminate or affect the product being sampled.

5. Procedure

5.1 Recommended Sampling Methods:

5.1.1 See recommended methods for sampling creosote and creosote solutions in Table 1.

5.1.2 See recommended methods for sampling oilborne and waterborne preservatives in Table 2.

5.2 Continuous Drip Sampling-Use this method of sampling wherever a liquid is being loaded or discharged from a

Shipping, Treating or Storage to be Sampled	Time of Sampling	Preferred Sampling Method	Type of Sample	Size of Sample to be Taken from Each Container	Additional Instructions
Ship and barge tanks	immediately after loading or prior to unloading	bottle or beaker sampling	all-levels sample	approximately 3 gal	Gage free water separately if creosote is not agitated thoroughly.
Tank cars and tank trucks	immediately after loading or prior to unloading	bottle or beaker sampling	composite of upper middle and lower samples	1 gal	Gage free water separately if creosote is not agitated thoroughly. Use Table 1 to determine proportions of subsamples in composite sample.
Vertical storage and working tanks	whenever required for inventory or other purposes	bottle or beaker sampling	all-levels sample	1–8 gal depending on size of tank	Gage free water separately if creosote is not agitated thoroughly.
Horizontal storage and working tanks	whenever required for inventory or other purposes	bottle or beaker sampling	composite of upper, middle and lower samples	1–3 gal depending on size of tank	Gage free water separately if creosote is not agitated thoroughly. Use Table 1 to determine proportions of subsamples in composite sample.
Pressure treating cylinders	immediately after cylinder is filled	tap sampling	tap sample	1 gal	
Non-pressure treating vats	immediately after vat is filled for cold bath	bottle or beaker sampling	all-levels sample	1 gal	

TABLE 1 Sampling of Creosote and Creosote Solutions^A

^A When the sample is taken the temperature of the material should be at least 150°F and the material should be completely liquefied and thoroughly agitated.

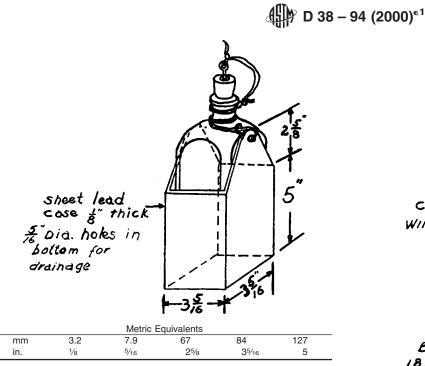


FIG. 2 Weighted Bottle

vessel by means of pumping. Obtain the sample from a pipe line conveying the product in such manner as to give a representative average of the stream throughout the period of transit.

5.2.1 Insert a ¹/₄-in. sampling pipe in the line through which the liquid preservative is being pumped, on the discharge side of the pump and preferably in a rising section of the pipe line. Extend this sampling pipe halfway to the center of the main pipe, with the inner open end of the sampling pipe turned at an angle of 1.57 radius (90°) and facing the flow of the liquid. Provide the sampling pipe with a valve and discharge it into a receiver of 4 to 19-L (1 to 5-gal) capacity. Clean the sampling pipe of foreign material by quickly collecting and discarding the first 500 mL of discharge.

5.2.2 Adjust the valve so that with a steady continuous drip of the preservative the receiver will be filled in the time required to pump the entire quantity of material being transferred.

5.2.3 Immediately upon completion of the pumping, thoroughly mix the contents of the receiver by agitation (heat to complete liquidity if necessary), rolling, or shaking, and immediately take duplicate 1-L (1-qt) samples for the purpose of testing. Do not collect less than 4 L (1 gal) of drop samples except in the case of large volumes, such as boat shipments, where a maximum of 19 L (5-gal) is sufficient.

5.3 *Zone Sampling*—Use this method of sampling to sample liquid-wood preservatives in tank cars, storage tanks, tank trucks and trailers, work tanks, ship tanks, and barge tanks. Semiliquid materials, which can be heated until sufficiently fluid, may be sampled by this procedure. Thoroughly agitate creosote and solutions of creosote with coal tar or petroleum and heat to 65 to 93°C (150 to 200°F) to effect solution. Obtain the sample by submerging a closed sampler to a point as near

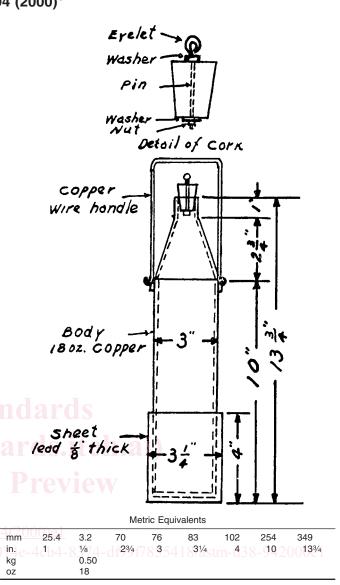


FIG. 3 Weighted Beaker

as possible to the selected draw-off level, then open the sampler, allowing the sample container to fill at that point, and slowly withdraw it.

5.3.1 Obtain three samples. Take each in a similar manner, with the sampling apparatus lowered to the point selected as the following zones (Table 3):

Upper zone	obtained from the middle of the upper third of the vessel
	contents.
Middle zone	obtained from the middle of the vessel contents.
Lower zone	obtained from the middle of the lower third of the vessel
	contents.

5.3.2 The individual samples may be analyzed separately or composited for testing. If composited, combine the three samples (Table 3) and mix at once while the material is thoroughly liquid. Take no sample for analysis from material that contains crystals.

5.4 *Thief Sampling*—Use this method of sampling to obtain bottom samples of liquids or semiliquids from tank cars, and storage tanks, by means of the liquid thief apparatus, and to obtain samples of granular material in bags, drums, or boxes, by means of the solids thief apparatus.