



Designation: D6323 – 98 (Reapproved 2003)

Standard Guide for Laboratory Subsampling of Media Related to Waste Management Activities¹

This standard is issued under the fixed designation D6323; 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 guide covers common techniques for obtaining representative subsamples from a sample received at a laboratory for analysis. These samples may include solids, sludges, liquids, or multilayered liquids (with or without solids).

1.2 The procedures and techniques discussed in this guide depend upon the sample matrix, the type of sample preparation and analysis performed, the characteristic(s) of interest, and the project specific instructions or data quality objectives.

1.3 This guide includes several sample homogenization techniques, including mixing and grinding, as well as information on how to obtain a specimen or split laboratory samples.

1.4 This guide does not apply to air or gas sampling.

1.5 The values stated in SI units are to be regarded as the standard.

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:*²

C702 Practice for Reducing Samples of Aggregate to Testing Size

C859 Terminology Relating to Nuclear Materials

D346 Practice for Collection and Preparation of Coke Samples for Laboratory Analysis

D2234/D2234M Practice for Collection of a Gross Sample of Coal

D4547 Guide for Sampling Waste and Soils for Volatile Organic Compounds

¹ This guide is under the jurisdiction of ASTM Committee D34 on Waste Management and is the direct responsibility of Subcommittee D34.01.01 on Planning for Sampling.

Current edition approved Aug. 10, 2003. Published December 1998. DOI: 10.1520/D6323-98R03.

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.

D4823 Guide for Core Sampling Submerged, Unconsolidated Sediments

D5743 Practice for Sampling Single or Multilayered Liquids, With or Without Solids, in Drums or Similar Containers

D5792 Practice for Generation of Environmental Data Related to Waste Management Activities: Development of Data Quality Objectives

D5956 Guide for Sampling Strategies for Heterogeneous Wastes

D6051 Guide for Composite Sampling and Field Subsampling for Environmental Waste Management Activities

3. Terminology

3.1 *contaminant unit, n*—the largest particle size that contains the contaminant of interest

3.1.1 *Discussion*—The contaminant of concern, as defined by the project objectives, may be associated with all the particle sizes or associated with only a certain particle size or sizes. At the time of waste generation, discharge or spill, the particle size of this contaminant of concern may be on the atomic or molecular scale, such as solvent spill into sand, or a macro scale, such as lead acid batteries at a dump site. The contaminant unit may also be in-between these scales, such as lead particles encapsulated in coal. In practice, the contaminant unit may change if the contaminant unit becomes absorbed or adsorbed to particles larger than the contaminant unit. It is the size of the contaminant unit at the time of subsampling, not at the time of generation, that is referred to as the contaminant unit.

3.2 *maximum allowable particle size, n*—the largest lineal dimension of a sample's individual particles accepted for a given sample mass.

3.2.1 *Discussion*—The maximum allowable particle size is sometimes referred to as the allowable particle size. A simple method of measurement is a sieve.

3.3 *multilayered sample, n*—a sample consisting of two or more clearly differentiated components.

3.3.1 *Discussion*—Multilayered samples are those with two or more distinct visual layers of material. These layers may be the result of differences in density, such as liquid/liquid layers (for example, chlorinated solvents and water, water and oil),

liquid/solid layers (for example, sludge), solid/solid layers (for example, small rocks and large rocks), or combinations of these layers (for example water, oil, and soil). These layers may also be the result of depositional layering, such as green clay and silty sand from a coring sample.

3.4 *particle size, n*—the controlling lineal dimension of individual particles (see Terminology C859).

3.5 *representative subsample, n*—a subsample collected in such a manner that it reflects one or more characteristics of interest (as defined by the project objectives) of the laboratory sample from which it was collected.

3.5.1 *Discussion*—A representative subsample can apply to a single sample, or a composite sample.

3.6 *sample, n*—a portion of material taken from a larger quantity for the purpose of estimating properties or composition of the larger quantity, (see Guide D6051).

3.7 *sludge, n*—Any mixture of solids that settles out of solution. Sludges contain liquids that are not apparent as free liquids, (see Practice D5743).

3.8 *subsample, n*—a portion of a sample taken for the purpose of estimating properties or composition of the whole sample, (see Guide D6051).

3.8.1 *Discussion*—A subsample is also commonly referred to by such terms as specimen, aliquot, subsample, or analytical subsample. To avoid confusion, the term specimen is used to refer to the final portion of the sample used during analysis. The material received at the laboratory will be referred to as the laboratory sample. All portions subsampled in-between receipt at the laboratory and the specimen are referred to as subsamples. The subsample may be the specimen.

4. Significance and Use

4.1 This guide discusses options for taking a subsample from a sample submitted to a laboratory. If followed, it will minimize the bias and variance of the characteristic of interest of the laboratory sample prior to analysis.

4.2 The guide will describe appropriate instructions to be submitted to the laboratory with the field sample.

4.3 This guide is intended for use in the laboratory to take a representative subsample or specimen of the whole field sample for direct analysis or sample preparation for analysis. It is intended for field personnel, data users, laboratory sample reception personnel, analysts, and managers.

4.4 To obtain a representative subsample, layer analysis, grinding, mixing, and changing the physical state such as digesting, drying, melting or freezing may be required. This guide considers cone and quartering, riffle splitting, and particle size reduction.

5. General Considerations

5.1 Successful implementation of this standard depends on effective communication between the data user and the laboratory staff. The selection of optimal subsampling procedures, techniques, and strategy by the laboratory depends on the intended use of the data. The data user should submit appropriate instructions with all samples and, when necessary, the laboratory staff should contact the data user for confirmation or further clarification of these instructions.

5.1.1 The appropriate instructions must be reviewed by both the laboratory receiving personnel and the analyst(s) or supervisor. If there are no instructions, the appropriate laboratory personnel should contact the data user. Options should be discussed and clarified prior to initiating any subsampling procedure. These instructions may include such options as those found in Table 1. The limitations and advantages of these methods are also found in this table. The data user should be informed about the limitations and advantages of all subsampling procedures prior to deciding which one to use.

5.1.2 If the data user still provides no instructions upon being contacted, laboratory personnel should explain to the data user that the laboratory's standard operating procedures, which reflect the concerns and issues discussed in this standard, will be used. The sample should be treated as if the scale of the contamination is on the micro level, and no artifacts can be removed. Since sample matrices and types and mechanisms of contamination are infinitely variable and require judgments to be made, it is advisable that experienced analysts decide which subsampling techniques be employed. These procedures must be discussed and clarified with the data user prior to initiating any subsampling procedure.

5.2 If the sample integrity or composition is not as anticipated, the data user must be contacted to confirm or clarify the instructions. An example of when this would be necessary would be a case where a coring sleeve was received at the laboratory. On opening the container, the analyst notices clay in one end of the sleeve, and sand at the other end. Before the analyst can proceed, the appropriate instructions from the data user must be obtained.

5.2.1 Field samples should be collected in appropriate containers for the analyses requested. If the submitted sample is improperly collected, the data user should be contacted by laboratory personnel. If the data user authorizes the laboratory to continue with the analysis, a note should be made in the receipt documentation, and also in the case narrative in the final report.

5.3 Documentation during the subsampling process is critical. Since subsampling techniques may bias the results, the subsampling method used must be noted in the analytical logbook.

5.3.1 Anytime the analytical result will be biased, it must be documented, and the data user should be notified prior to beginning any subsampling technique. For example, if headspace exists in a container arriving at the laboratory, some volatile components will have partitioned into that headspace. However, if the data user decides to proceed with the analysis, the analytical logbook and the case narrative on the final report should indicate this condition.

5.4 Particle size is the physical dimension of an object's pieces or parts. The maximum particle size contained within a laboratory sample is the largest of these pieces. The contaminant of concern, as defined by the project objectives, may be associated with all particle sizes or associated with only a certain particle size or sizes. The largest of these particle sizes, that contain the contaminant of interest, would be the contaminant unit. The contaminant unit, at the time of waste generation, discharge or spill, may be on the atomic or molecular

TABLE 1 Limitations and Advantages of Sample Preparation Options

Instruction	Limitations	Advantages
Remove artifacts, such as rocks and twigs, from the sample prior to subsampling	(1) May bias analytical results by altering contaminant concentration, (2) May bias sample if results are not properly weight averaged.	(1) May be easier to subsample, (2) May be easier to analyze, (3) Appropriate if the target population is material minus artifacts.
Digest or extract the contaminant from the outside of the large particles	May bias sample if contaminant is within the large particles. ^A	(1) May be easier to analyze, (2) May prevent need for weight average calculation.
Digest or extract particle sizes separately	(1) Separation of particle sizes may be difficult, ^A (2) May bias sample if results are not properly weight averaged, (3) Higher cost.	(1) Allows some particle size consistency during analysis, (2) May be easier to subsample within portions after separation.
Form an emulsion layer so that the material may be treated as homogeneous liquid	May bias the sample if a complete emulsion is not achieved. ^A	(1) May be easier to subsample as a homogeneous liquid,
Separate liquid layers	(1) Separation of layers may be difficult, especially at the interface, (2) May bias sample if results are not properly weight averaged.	(1) May be easier to analyze, (2) May be easier to subsample within portions after separation, (3) Allows different preparation methods within each layer.
Dry sample	May alter chemistry or change stability of some compounds. ^A	(1) Allows for consistency of subsampling for liquid/solid mixtures, (2) Analytes reported unbiased by moisture content.
Change the physical state, such as freeze the material so that it may be treated as a solid, or melt the material so that it may be treated as a liquid	May be difficult to achieve complete freezing or melting and maintain it long enough to get a subsample. ^A	Allows for consistency of subsampling.
Analyze only one layer of multilayered samples, such as analyze only the oil portion of an oil/water mixture for PCBs	May bias the sample if there is cross-contamination between layers	(1) Possible cost savings to customer, (2) May be easier to subsample from a single layer, (3) May be easier to analyze.
Composite portions of the sample for volatile analysis directly in a purge unit vs. individual analysis of these portions	(1) May overload gas chromatography columns if sample has high amounts of solvents in each portion, (2) Separation of portions may be difficult.	(1) May prevent losses of volatile because the sample is handled only once, (2) Possible cost savings to customer, (3) May prevent need for weight average calculation.
Reduce particle size	(1) Increasing surface area may effect data in some procedure with particle digestion or extraction, such as TCLP, ^A (2) May be difficult, depending on matrix.	Allows for consistency of subsampling.
Use standard methods for solids (for example, cone and quarter, grind, riffle, sieve)	Dependent on method. ^A See 7.1.5 for more information.	Dependent on method. See 7.1.5 for more information.

^AMay be unsuitable for samples to be analyzed for volatile constituents.

scale, such as a solvent spill into sand, or a macro scale, such as lead acid batteries at a dump site. The contaminant unit may also be in-between these scales, such as lead particles encapsulated in coal. In practice, the contaminant unit may change if the contaminant becomes absorbed or adsorbed to particles larger than the contaminant unit.

5.4.1 Knowledge of the contaminant unit may be used to determine the preliminary steps to subsampling. For example, if the contaminant unit is on a molecular scale and was adsorbed to soil particles and rocks, removal of large rocks with their relatively small surface area may not affect the data as long as the results are weight averaged. If one is unsure of the mechanism of contamination, that determines how the contamination is dispersed within the sample matrix, one may not be able to discard any particles during subsampling.

5.5 Sampling theory requires that subsample *mass* should increase as the size of the largest particle in the sample increases. If the subsample mass recommended by sampling theory is larger than that normally used in the sample preparation method, the subsample mass may be increased and the extraction/digestion procedure scaled accordingly. The standard volume of digestate/extract is then submitted for analysis. However, if the subsample mass is too large to be accommo-

dated by the sample preparation procedure, multiple subsamples (of equal mass) can be extracted/digested, and the extracts/digests combined and mixed prior to removal of the standard volume specimen needed for analysis. Another alternative is to reduce the particle size of the entire sample or subsample as specified by sampling theory. If the particle size is reduced enough, a subsample of the mass recommended by sampling theory and the extraction/digestion method can be obtained.

5.6 All subsampling should be performed in an area which is free from contamination, easily decontaminated, and vented to control dust and remove fumes.

5.7 Prior to subsampling liquids, the analyst must consider the property or characteristic requested, and the container size received. If the analyte has the ability to adsorb onto the container, the field sample should arrive at the laboratory in an appropriately sized bottle, such that the whole sample will be used. The sides of the container should then be rinsed properly to assure that all the contaminants are transferred into the analytical vessel.

5.8 Subsampling techniques are different when analyzing for volatile compounds than non-volatile compounds. The differences are discussed for each sample matrix.

TABLE 2 Recommended Minimum Subsample Mass for Particulate Materials to Achieve a Fundamental Error of 15 %

Recommended Minimum Subsample Mass, g	Maximum Allowable Particle Size ^A , cm	Size Class ^B	U.S. Standard Sieve Mesh Size
0.1	0.05	Clay to coarse sand	35
1	0.1	Coarse to very coarse sand	18
2	0.13	Very coarse sand	13
5	0.17	Very coarse sand	12
10	0.21	Granule gravel	10
30	0.31	Granule gravel	7
50	0.37	Granule gravel	6
100	0.46	Pebble gravel	5

^AThe maximum allowable particle size allowed can be approximated for other sample masses by using the equation, allowable particle size (cm) = the cube root of $0.001 \times$ sample mass in grams

^BWentworth Size Class (See (3) and (4)) These soil descriptions are added for those readers who equate particle size with the Wentworth descriptions and are not meant to indicate that the application of this table is limited to soil particles. The table can also be used for non-soil particles such as waste.

6. Matrix Specific Subsampling

6.1 Solids:

6.1.1 **Table 2** gives the relationship between the maximum particle size and sample mass to achieve a fundamental error of 15 %. For information about fundamental error, see Ref (1)³. If the maximum allowable particle size is greater than that listed in the table, then sampling theory would suggest particle size reduction, such as grinding, or use of a larger sample mass. For example, if granule gravel has an allowable particle size of 0.21 cm, according to this table, a subsample mass of 10 g is needed. Many digestions/extractions commonly use 1 g. The 10 g subsample would be ground to a size of 0.1 cm, and the specimen would be taken from the ground material. Approximations were employed to construct this table. The actual fundamental error could be much larger. More information on calculating the maximum allowable particle size for various fundamental errors, and particles shapes is included in **Annex A1**.

6.1.1.1 Particle size reduction can be avoided by modifying the sample digestion/extraction methods to use large specimen masses. Following digestion/extraction of the large specimen, the homogeneous digestate/extract is analyzed. For example, if the contaminant of concern was a chromium solution spilled into a mixture of gravel and soil, knowing that the chromium coats the medium allows a choice of sample preparation. The analyst may totally dissolve the specimen. He/she may digest the chromium from the outside of the particles, analyzing the digestate. Or, he/she may extract the material, and analyze the extractant. If the analyst had no information about the mechanism of contamination, he/she would have to assume that the contamination is throughout all particles of the laboratory sample.

6.1.1.2 If information is available regarding the type and mechanism of contamination, then it may be possible to avoid particle size reduction and the use of large specimen masses. For example, if contamination occurred on a molecular level

³ The boldface numbers given in parentheses refer to a list of references at the end of the text.

(for example, a solvent is discharged to soil), an argument can be made to preclude large particles (for example, rocks), since insignificant amounts of contamination will be adsorbed to the small surface of the large particles. If the large particles are excluded, the appropriate mass can be based on the smaller particle sizes, which have a much larger surface area. If the decision is made to remove the large particles, the mass of both the large and small particles would be used to calculate the final concentration of the contaminant of interest.

6.1.2 If larger sample masses are required, two ASTM Standards may be helpful. Test Method **D2234/D2234M** discusses test methods and procedures for the collection of a sample under various conditions of sampling. It describes general and special purpose sampling procedures for coal by size, condition, and characteristics. The document also includes in the annex, a test method for determining the variance of components of a coal and a test method for estimating the overall variance for increments of one fixed weight of a given coal.

6.1.2.1 A second method which is often referenced by other ASTM standards dealing with coals is Practice **D346**. The practice is designed to provide a representative sample of the coke from which it is collected. It considers the variability of coke and the wide variety of sampling equipment.

6.1.3 If the desired subsample mass is equal to or less than the maximum allowable particle size, subsampling can proceed without any need for particle size reduction. If the particle size is greater than the allowable particle size in **Table 2**, then the sample matrix, the type and mechanism of contamination, the scale of the contaminant unit, as well as particle size reduction should be considered. These allowable particle sizes are based on the assumption that the contaminant exists as a particle or is adsorbed to particles. In those cases where the contaminant unit is on the atomic or molecular scale or is a particle much smaller than the allowable particle size of the sample, then alternative approaches can be employed with acceptable bias.

6.1.4 Solid Samples for Volatile Analysis:

6.1.4.1 Due to the volatility of this class of compounds, laboratory subsampling procedures addressing the sample's allowable particle size could result in large losses of the parameter of interest, rendering the results unrepresentative. The possible exception is when cryogenic freezing conditions are maintained throughout the handling process.

NOTE 1—If cryogenic techniques (working with frozen samples) or other special techniques are used, the procedure should be validated on laboratory control samples (LCS) which have been spiked with known concentrates of the parameter of interest. The nature of the test matrix for the LCS and the spiking solution should be chosen to allow uniform distribution of the spike in the matrix prior to subsampling. Regardless of the particle size reduction techniques chosen, it is advisable to carry an LCS through each preparation batch as a quality control sample. A subsample of the field sample without particle size reduction through the validation procedure will demonstrate the amount of volatile loss (from the LCS results) and increased volatiles recovery (from a comparison of the unreduced field sample with the size reduced sample) using particle size reduction.)

6.1.4.2 When analyzing samples for low levels of volatile constituents (0.5 to 200 $\mu\text{g}/\text{kg}$ range), discrete samples should be placed into a container that, once sealed, is analyzed without

opening (see Guide [D4547](#)). For this reason, and because sample mass is often limited by the analysis system, low level samples typically are not amenable to either particle size reduction procedures or the analysis of larger sample mass. When contaminant levels exceed 200 µg/kg, discrete samples can be transferred to vessels containing an appropriate solvent, such as methanol. Because the solvent retains the volatiles in solution, and only a small portion of the solvent is removed for analysis, particle and contaminant unit size with respect to subsample mass requirements can be considered.

NOTE 2—Current analytical guidance recommends solvent extraction only for volatile organic concentrations at greater than 200 µg/kg. However, solvent extraction may be used for solvents of lower concentration levels if the solvent/solid ratio is reduced, or by using an analysis system with lower detection limits.

6.1.4.3 The following subsections discuss different subsampling procedures that can be used as particle size and contaminant unit vary in relationship to the allowable particle sizes listed in [6.1.1](#).

6.1.5 *Laboratory Sample Particle Size and Contaminant Unit Less Than the Allowable Particle Size*—Prepare sample vessel with the appropriate solvent and solvent volume as specified in the method. Add the appropriate mass of sample to the prepared vessel with a pre-cleaned coring tool (see Guide [D4547](#)) or spatula as quickly as possible. The coring tool must be large enough to accommodate the largest sample matrix particle. After volatile extraction, remove the appropriate volume of the solvent for analysis from the liquid portion of the specimen.

6.1.6 *Laboratory Sample Particle Size Greater Than the Allowable Particle Size and the Contaminant Unit Less Than the Allowable Particle Size*—Particle size reduction cannot generally be used, since it may result in a loss of volatile contaminants. There are cases where the chemistry of the contaminant and the support will not allow the volatile to be lost. For these types of contaminants, see the methods in [6.1.5](#). Alternate approaches to subsampling may be viable if knowledge exists regarding the sample matrix, type and mechanism of contamination, and the contaminant unit.

6.1.6.1 If the contaminant unit is on a molecular scale, then the distribution of the contaminant will be a function of the absorptive properties, and the larger surface area of the smaller particles results in more contaminants being adsorbed to the smaller particles. If the data user is willing to accept a biased-high concentration for the contaminant of interest, a subsample of only the smaller particles may be made.

6.1.6.2 Another method is to use a solvent appropriate for analysis, such as methanol, to extract the material. Use a 4 g specimen of the original laboratory sample in 10 mL of solvent. Increase the volumes proportionally to accommodate the specified sample mass. An appropriate portion of the solvent is analyzed once the contaminant is dissolved in the solvent.

6.1.7 *Laboratory Sample Particle Size and Contaminant Unit Larger than the Allowable Particle Size*—When sample masses become prohibitive with regard to logistical constraints, sample extracts can be composited. An example of this is the case where soil of particle size 0.46 cm is contaminated with volatile organic compounds. The suggested

subsample mass in this case from [Table 2](#) is 100 g, therefore requiring a vessel with 250 mL of solvent. As an alternate to a single large sample, five 20 g subsamples could be extracted. Microliter amounts of each of the five resulting extracts could be composited into the same purge chamber. Once the contaminants are dissolved in the solvent, the concern of contaminant heterogeneity is decreased and the use of small subsamples of the extracts can be justified.

6.1.8 *Solid Sample for Non-Volatile Analysis*—The method of subsampling is dependent on the size of the allowable particle size and the size of the contaminant unit. The following subsections discuss different subsampling procedures that can be used as laboratory sample particle size and contaminant unit vary in relationship to the allowable particle sizes listed in [Table 2](#). If the laboratory sample requires mixing prior to subsampling, the methods in section [6.1.6](#) should be used.

6.1.9 *Laboratory Sample Particle Size and Contaminant Unit Less Than the Allowable Particle Size*:

6.1.9.1 *Transversal Subsampling (Rectangular Scoop)* (see [Fig. 1](#))—One method of subsampling the materials is to use transversal subsampling (see Guide [D6051](#)). The entire laboratory sample is emptied onto a non-contaminating smooth surface. The sample is shaped into an elongated pile with a flattened top surface ([1](#)). Complete top-to-bottom transversal cuts are made across the pile and the extracted material is transferred into a tared container. The transversal cuts are repeated until the appropriate mass is obtained. Cuts are made with a rectangular scoop. The width of the transversal section should be wide enough to accommodate the largest particle size, but narrow enough to require multiple transversal sections be used to obtain the appropriate mass. Two thin sheets of Teflon, metal, or polyethylene may be used to facilitate the removal of material within the transversal cut. After the removal of a few transversal sections, the pile may be reshaped to facilitate further cuts. The entire unused portion of the laboratory sample should be returned to the original sample container.

6.1.9.2 *Cone and Quartering* (see [Fig. 2](#))—Another method of subsampling materials involves cone and quartering ([2](#)). This method may introduce large biases from the grouping and segregation error. It is discussed here due to its widespread use, but the user should take note of its limitations (see [Table 4](#)). The entire laboratory sample is emptied out onto a non-contaminating smooth surface. Material is piled into a cone with a flattened top surface. Two top-to-bottom cuts are made through the cone at perpendicular angles to form four equal portions, or quarters. Two opposite quarters are compiled into a new cone, and the process is repeated until the proper sample mass is achieved. See [C702](#) for more information.

6.1.9.3 *Riffle Splitter* (see [Fig. 3](#))—A riffle splitter is a mechanical device with an equal number of narrow sloping chutes ([2](#)). Alternate chutes discharge the sample into opposite directions into two collection pans. Subsampling is achieved by pouring the entire laboratory sample into the riffle splitter. One portion is removed, and the collection pan is replaced with a clean pan. This portion is poured back through the riffle splitter so that half of the subsample will combine with the original material, and the other half will drop into the clean

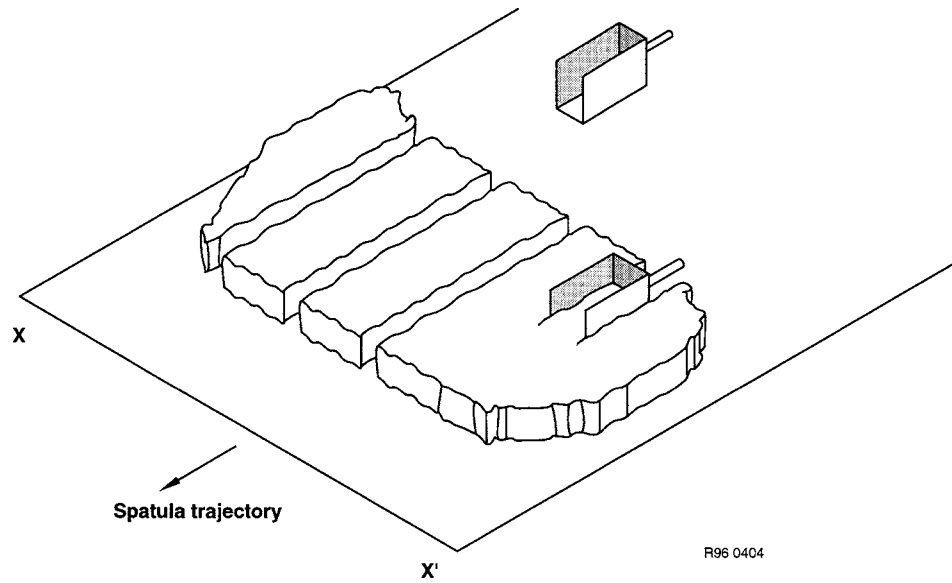


FIG. 1 Transversal Subsampling

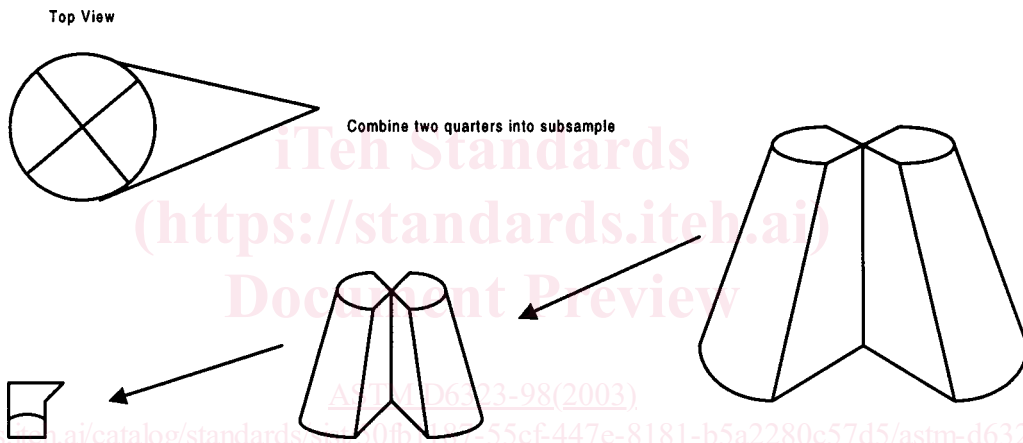


FIG. 2 Cone and Quartering

collection pan. This process is repeated until the proper sample mass is achieved. There are many types of riffle splitters available. The analyst must select the one that meets the data user's needs. The riffle splitter opening should be about three times the largest particle diameter, and have an even number of chutes.

6.1.9.4 *Alternate Scoop*—This method is discussed in Guide 6051. Scoops of the mixed laboratory sample are either placed in the analytical vessel, or discarded. In the example in Fig. 4, three scoops are discarded for every scoop saved. Care should be taken that each scoop of material is of the same size and is taken in a consistent manner to minimize bias.

6.1.9.5 *Sectorial Splitter (Sample Divider)*—(See Fig. 5)—This is a spinning riffler-type sample divider. The sample is placed in the opening, and the material is vibrated through shoots to the spinning bottles. The specimen may be taken directly from the bottle, or it may be further subsampled by discarding every other bottle, placing the remaining sample back in the opening, and starting the process over again.

6.1.10 *Laboratory Sample Particle Size Greater Than the Allowable Particle Size and the Contaminant Unit Less Than the Allowable Particle Size:*

6.1.10.1 If the data user can accommodate worst-case, such as biased-high concentration of the parameter of interest, then subsampling can be performed without the need for large sample masses or particle size reduction. The procedure described in 6.1.11.1 will work, except that the particles larger than the allowable particle size are removed from the transversal section prior to addition to the subsample.

6.1.10.2 If the data user cannot accommodate worst-case, refer to 6.1.11.

6.1.11 *Laboratory Sample Particle Size and Contaminant Unit Larger than the Allowable Particle Size:*

6.1.11.1 Since the contaminant is associated with the large particles in this type of laboratory sample, the option of selectively removing large particles is not viable. Optimum solutions would be to increase the sample mass, or reduce the particle size. If these methods are not practical, a multiple