$01/2008{:}20938$

2.9.38. PARTICLE-SIZE DISTRIBUTION ESTIMATION BY ANALYTICAL SIEVING

Sieving is one of the oldest methods of classifying powders and granules by particle-size distribution. When using a woven sieve cloth, the sieving will essentially sort the particles by their intermediate size dimension (i.e., breadth or width). Mechanical sieving is most suitable where the majority of the particles are larger than about 75 µm. For smaller particles, their light weight provides insufficient force during sieving to overcome the surface forces of cohesion and adhesion that cause the particles to stick to each other and to the sieve, and thus cause particles that would be expected to pass through the sieve to be retained. For such materials other means of agitation such as air-jet sieving or sonic sifting may be more appropriate. Nevertheless, sieving can sometimes be used for some powders or granules having median particle sizes smaller than 75 μ m where the method can be validated. In pharmaceutical terms, sieving is usually the method of choice for classification of the coarser grades of single powders or granules. It is a particularly attractive method in that powders and granules are classified only on the basis of particle size, and in most cases the analysis can be carried out in the dry state.

Among the limitations of the sieving method are the need for an appreciable amount of sample (normally at least 25 g, depending on the density of the powder or granule, and the diameter of the test sieves) and difficulty in sieving oily or other cohesive powders or granules that tend to clog the sieve openings. The method is essentially a two-dimensional estimate of size because passage through the sieve aperture is frequently more dependent on maximum width and thickness than on length.

This method is intended for estimation of the total particle-size distribution of a single material. It is not intended for determination of the proportion of particles passing or retained on 1 or 2 sieves.

Estimate the particle-size distribution as described under Dry sieving method, unless otherwise specified in the individual monograph. Where difficulty is experienced in reaching the endpoint (i.e., material does not readily pass through the sieves) or when it is necessary to use the finer end of the sieving range (below 75 μ m), serious consideration must be given to the use of an alternative particle-sizing method.

Sieving is carried out under conditions that do not cause the test sample to gain or lose moisture. The relative humidity of the environment in which the sieving is carried out must be controlled to prevent moisture uptake or loss by the sample. In the absence of evidence to the contrary, analytical test sieving is normally carried out at ambient humidity. Any special conditions that apply to a particular material must be detailed in the individual monograph.

Principles of analytical sieving. Analytical test sieves are constructed from a woven-wire mesh, which is of simple weave that is assumed to give nearly square apertures and is sealed into the base of an open cylindrical container. The basic analytical method involves stacking the sieves on top of one another in ascending degrees of coarseness, and then placing the test powder on the top sieve. The nest of sieves is subjected to a standardised period of agitation, and then

This sieving process for estimating the particle-size distribution of a single pharmaceutical powder is generally intended for use where at least 80 per cent of the particles are larger than 75 μm . The size parameter involved in determining particle-size distribution by analytical sieving is the length of the side of the minimum square aperture through which the particle will pass.

TEST SIEVES

Test sieves suitable for pharmacopoeial tests conform to the most current edition of *ISO 3310-1: Test sieves – Technical requirements and testing – Part 1: Test sieves of metal wire cloth* (see Table 2.9.38.-1). Unless otherwise specified in the monograph, use those ISO sieves listed as principal sizes in Table 2.9.38.-1 that are recommended in the particular region.

Table 2.9.38.-1.

ISO N	ominal Ap	erture	US	Recom-	European	
Princi- pal sizes	Supplementary sizes		Sieve No.	mended USP Sieves (mesh)	Sieve No.	Sieve No.
R 20/3	R 20	R 40/3		(,		
11.20 mm	11.20 mm	11.20 mm			11 200	
	10.00 mm					
		9.50 mm				
	9.00 mm					
8.00 mm	8.00 mm	8.00 mm				
	7.10 mm					
		6.70 mm				
	6.30 mm					
5.60 mm	5.60 mm	5.60 mm			5600	3.5
	5.00 mm					
		4.75 mm				4
	4.50 mm					
4.00 mm	4.00 mm	4.00 mm	5	4000	4000	4.7
	3.55 mm					
		3.35 mm	6			5.5
	3.15 mm					
2.80 mm	2.80 mm	2.80 mm	7	2800	2800	6.5
	2.50 mm					
		2.36 mm	8			7.5
	2.24 mm					
2.00 mm	2.00 mm	2.00 mm	10	2000	2000	8.6
	1.80 mm					
		1.70 mm	12			10
	1.60 mm					
1.40 mm	1.40 mm	1.40 mm	14	1400	1400	12
	1.25 mm					
		1.18 mm	16			14
	1.12 mm					
1.00 mm	1.00 mm	1.00 mm	18	1000	1000	16
	900 µm					

o	2.
fana	Me
alysi	thoc
S	ds

Principal sizes Supplementation sizes Sive mended (mesh) Sive m	ISO Nominal Aperture			US Sieve No.		European	Japanese
800 µm 800 µm 18 800 µm 710 µm 710 µm 710 µm 25 710 710 22 630 µm 25 710 710 22 26 630 µm 30 26 26 26 560 µm 500 µm 500 µm 35 500 500 30 500 µm 500 µm 355 µm 40 36 30 36 505 µm 425 µm 40 36 355 355 42 355 µm 355 µm 45 355 355 42 35 355 µm 355 µm 60 250 50 60 250 60 212 µm 300 µm 60 250 250 60 60 250 250 60 224 µm 120 121 120 120 120 100 <td< th=""><th colspan="2"></th><th>Sieve No.</th><th>Sieve No.</th></td<>			Sieve No.			Sieve No.	
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110 m 10 m 10 m 20 10 10 22 100 m 100 m 30 10 10 100 m 100 m 30 10 10 100 m 100 m 35 500 m 30 30 100 m 100 m 10 10 10 100 m 10 10 10 10 1			850 µm	20			18
$630 \mum$ $600 \mum$ 30 26 $500 \mum$ $500 \mum$ 35 500 500 30 $500 \mum$ $500 \mum$ $500 \mum$ 35 500 500 30 $500 \mum$ $425 \mum$ 40 36 36 36 36 $450 \mum$ $400 \mum$ $450 \mum$ 355 355 42 356 42 $315 \mum$ $355 \mum$ 45 355 355 42 356 42 $400 \mum$ $450 \mum$ $360 \mum$ $500 \mum$ $700 \mum$ $710 \mum$ <		800 µm					
SolumiSolum <th< td=""><td>710 µm</td><td>710 µm</td><td>710 µm</td><td>25</td><td>710</td><td>710</td><td>22</td></th<>	710 µm	710 µm	710 µm	25	710	710	22
560 µm 500 µm 500 µm 500 500 500 30 500 µm 600 µm 40 36 300 36 400 µm 400 µm 40 36 355 355 42 355 µm 555 µm 50 50 50 42 355 µm 550 µm 50 50 50 42 300 µm 50 50 50 50 50 280 µm 50 µm 50 50 50 60 280 µm 50 µm 60 250 250 60 210 µm 50 µm 60 250 250 60 210 µm 50 µm 60 250 100 70 101 µm 101 µm 100 100 100 102 µm 101 µm 101 100 100 102 µm 101 µm 100 100 100 101 µm 101 µm 100 100 100 102 µm 101 µm 100 100 100 101 µm 101 µm 100 100 100 102 µm 101 µm 100 100 100 101 µm 101 µm 100		630 µm					
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450 µm 425 µm 40 36 400 µm 355 µm 455 µm 355 355 µm 42 355 µm 355 µm 45 355 355 42 355 µm 30 µm 40 50 50 50 300 µm 50 250 50 60 50 280 µm 200 µm 60 250 250 60 201 µm 212 µm 70 70 70 200 µm 180 µm 80 180 83 83 160 µm 180 µm 80 180 83 83 151 µm 151 µm 125 110 100 100 152 µm 125 µm 125 125 110 100 152 µm 125 µm 125 125 126 126 160 µm 170 90 90 166 100 170 µm 170 170 90 90 166 100 170 µm 170 µm 170 170 170 170 170 <td< td=""><td></td><td>560 µm</td><td></td><td></td><td></td><td></td><td></td></td<>		560 µm					
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$400 \mum$ $355 \mum$ 45 355 355 42 $315 \mum$ $300 \mum$ 50 50 50 $280 \mum$ $200 \mum$ 60 250 250 60 $224 \mum$ $212 \mum$ 70 70 70 $200 \mum$ $212 \mum$ 70 70 70 $180 \mum$ $800 \mum$ 80 800 800 830 $180 \mum$ $150 \mum$ 80 180 830 830 $160 \mum$ $150 \mum$ 120 125 125 110 $125 \mum$ $125 \mum$ 120 125 125 110 $120 \mum$ $100 \mum$ $100 \mum$ $100 \mum$ $100 \mum$ $100 \mum$ $90 \mum$ $90 \mum$ $75 \mum$ $90 \mum$ $90 \mum$ $90 \mum$ $100 \mum$ $63 \mum$ $51 \mum$ $200 \mum$ $63 \mum$ $63 \mum$ $63 \mum$ $200 \mum$ $100 \mum$ $610 \mum$ $51 \mum$ $51 \mum$ $51 \mum$ $51 $		450 µm					
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315 µm 300 µm 50 50 280 µm 250 µm 250 µm 60 250 250 60 250 µm 250 µm 60 250 250 60 60 224 µm 70 70 70 70 200 µm 212 µm 70 70 70 180 µm 80 µm 80 180 83 160 µm 150 µm 160 100 100 125 µm 150 µm 100 125 119 125 µm 125 µm 120 125 125 119 121 µm 100 µm 140 140 140 140 140 125 µm 106 µm 140		400 µm					
300 µm5050280 µm250 µm6025025060250 µm250 µm6025025060240 µm70707070200 µm70707070180 µm8018018083160 µm8018018083160 µm100100100155 µm125 µm120125125125 µm126 µm140100121 µm100 µm140100121 µm120125125126125 µm126 µm160 µm100100126 µm127 µm120125125127 µm128 µm120125125128 µm120120125125129 µm12012512	355 µm	355 µm	355 µm	45	355	355	42
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50 μm 45 μm 45 μm 325 45 45 330		56 µm					
45 μm 45 μm 45 μm 325 45 45 330			53 µm	270			282
		50 µm					
40 µm	45 µm	45 µm	45 µm	325	45	45	330
		40 µm					
38 µm 38 391			38 µm			38	391

Sieves are selected to cover the entire range of particle sizes present in the test sample. A nest of sieves having a $\sqrt{2}$ progression of the area of the sieve openings is recommended. The nest of sieves is assembled with the coarsest screen at the top and the finest at the bottom. Use micrometres or millimetres in denoting test sieve openings (Note: mesh numbers are provided in the table for conversion purposes only).

Test sieves are made from stainless steel or, less preferably, from brass or other suitable non-reactive wire.

Calibration and recalibration of test sieves is in accordance with the most current edition of ISO 3310-1. Sieves are carefully examined for gross distortions and fractures, especially at their screen frame joints, before use. Sieves may be calibrated optically to estimate the average opening size, and opening variability, of the sieve mesh. Alternatively, for the evaluation of the effective opening of test sieves in the size range of 212-850 μ m, standard glass spheres are available. Unless otherwise specified in the individual monograph, perform the sieve analysis at controlled room temperature and at ambient relative humidity.

Cleaning test sieves. Ideally, test sieves are cleaned using only a low-pressure air jet or a liquid stream. If some apertures remain blocked by test particles, careful gentle brushing may be used as a last resort.

Test sample. If the test sample mass is not given in the monograph for a particular material, use a test sample having a mass between 25-100 g, depending on the bulk density of the material, for test sieves having a 200 mm diameter. For 76 mm sieves, the amount of material that can be accommodated is approximately 1/7 that which can be accommodated by a 200 mm sieve. Determine the most appropriate mass for a given material by test sieving accurately weighed samples of different masses, such as 25 g, 50 g, and 100 g, for the same time period on a mechanical shaker (note: if the test results are similar for the 25 g and 50 g samples, but the 100 g sample shows a lower percentage through the finest sieve, the 100 g sample size is too large). Where only a sample of 10-25 g is available, smaller diameter test sieves conforming to the same mesh specifications may be substituted, but the endpoint must be redetermined. The use of tests samples having a smaller mass (e.g. down to 5 g) may be needed. For materials with low apparent particle density, or for materials mainly comprising particles with a highly iso-diametrical shape, sample masses below 5 g for a 200 mm screen may be necessary to avoid excessive blocking of the sieve. During validation of a particular sieve analysis method, it is expected that the problem of sieve blocking will have been addressed.

If the test material is prone to absorbing or losing significant amounts of water with varying humidity, the test must be carried out in an appropriately controlled environment. Similarly, if the test material is known to develop an electrostatic charge, careful observation must be made to ensure that such charging does not influence the analysis. An antistatic agent, such as colloidal silicon dioxide and/or aluminum oxide, may be added at a 0.5 per cent (m/m) level to minimise this effect. If both of the above effects cannot be eliminated, an alternative particle-sizing technique must be selected.

Agitation methods. Several different sieve and powder-agitation devices are commercially available, all of which may be used to perform sieve analyses. However, the different methods of agitation may give different results for sieve analyses and endpoint determinations because of the different types and magnitudes of the forces acting on the individual particles under test. Methods using mechanical agitation or electromagnetic agitation, and that can induce either a vertical oscillation or a horizontal circular motion. or tapping or a combination of both tapping and horizontal circular motion are available. Entrainment of the particles in an air stream may also be used. The results must indicate which agitation method was used and the agitation parameters used (if they can be varied), since changes in the agitation conditions will give different results for the sieve analysis and endpoint determination, and may be sufficiently different to give a failing result under some circumstances.

Endpoint determination. The test sieving analysis is complete when the mass on any of the test sieves does not change by more than 5 per cent or 0.1 g (10 per cent in the case of 76 mm sieves) of the previous mass on that sieve. If less than 5 per cent of the total sample mass is present on a given sieve, the endpoint for that sieve is increased to a mass change of not more than 20 per cent of the previous mass on that sieve.

If more than 50 per cent of the total sample mass is found on any one sieve, unless this is indicated in the monograph, the test is repeated, but with the addition to the sieve nest of a more coarse sieve intermediate between that carrying the excessive mass and the next coarsest sieve in the original nest i.e., addition of the ISO series sieve omitted from the nest of sieves.

SIEVING METHODS

Mechanical agitation (Dry sieving method). Tare each test sieve to the nearest 0.1 g. Place an accurately weighed quantity of test sample on the top (coarsest) sieve, and replace the lid. Agitate the nest of sieves for 5 min, then carefully remove each sieve from the nest without loss of material. Reweigh each sieve, and determine the mass of material on each one. Determine the mass of material in the collecting pan in a similar manner. Re-assemble the nest of sieves, and agitate for 5 min. Remove and weigh each sieve as previously described. Repeat these steps until the endpoint criteria are met (see Endpoint determination under Test sieves). Upon completion of the analysis, reconcile the masses of material. Total losses must not exceed 5 per cent of the mass of the original test sample.

Repeat the analysis with a fresh sample, but using a single sieving time equal to that of the combined times used above. Confirm that this sieving time conforms to the requirements for endpoint determination. When this endpoint has been validated for a specific material, then a single fixed time of sieving may be used for future analyses, providing the particle size distribution falls within normal variation.

If there is evidence that the particles retained on any sieve are aggregates rather than single particles, the use of mechanical dry sieving is unlikely to give good reproducibility, and a different particle size analysis method must be used.

Air-entrainment methods (Air-jet and sonic-sifter sieving). Different types of commercial equipment that use a moving air current are available for sieving. A system that uses a single sieve at a time is referred to as, air-jet, sieving. It uses the same general sieving methodology as that described under Dry sieving method, but with a standardised air jet replacing the normal agitation mechanism. It requires sequential analyses on individual sieves starting with the finest sieve to obtain a particle size distribution. Air jet sieving often includes the use of finer test sieves than used in ordinary dry sieving. This technique is more suitable where only oversize or undersize fractions are needed.

In the sonic sifting method, a nest of sieves is used, and the test sample is carried in a vertically oscillating column of air that lifts the sample and then carries it back against the mesh openings at a given number of pulses per minute. It may be necessary to lower the sample amount to 5 g, when sonic sifting is employed.

The air-jet sieving and sonic sieving methods may be useful for powders or granules when the mechanical sieving techniques are incapable of giving a meaningful analysis. These methods are highly dependent upon proper dispersion of the powder in the air current. This requirement may be hard to achieve if the method is used at the lower end of the sieving range (i.e., below 75 μ m), when the particles tend to be more cohesive, and especially if there is any tendency

for the material to develop an electrostatic charge. For the above reasons endpoint determination is particularly critical, and it is very important to confirm that the oversize material comprises single particles and is not composed of aggregates.

INTERPRETATION

The raw data must include the mass of test sample, the total sieving time, the precise sieving methodology, and the set values for any variable parameters, in addition to the masses retained on the individual sieves and in the pan.

It may be convenient to convert the raw data into a cumulative mass distribution, and if it is desired to express the distribution in terms of a cumulative mass undersize, the range of sieves used must include a sieve through which all the material passes. If there is evidence on any of the test sieves that the material remaining on it is composed of aggregates formed during the sieving process, the analysis is invalid.

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2.9.40. UNIFORMITY OF DOSAGE UNITS

To ensure the consistency of dosage units, each unit in a batch should have an active substance content within a narrow range around the label claim. Dosage units are defined as dosage forms containing a single dose or a part of a dose of an active substance in each dosage unit. The uniformity of dosage units specification is not intended to apply to suspensions, emulsions, or gels in single-dose containers intended for cutaneous administration.

The term "Uniformity of dosage unit" is defined as the degree of uniformity in the amount of the active substance among dosage units. Therefore, the requirements of this chapter apply to each active substance being comprised in dosage units containing one or more active substances, unless otherwise specified elsewere in this Pharmacopoeia. The uniformity of dosage units can be demonstrated by

either of 2 methods: content uniformity or mass variation (see Table 2.9.40.-1).

The test for content uniformity of preparations presented in dosage units is based on the assay of the individual contents of active substance(s) of a number of dosage units to determine whether the individual contents are within the limits set. The content uniformity method may be applied in all cases.

The test for mass variation is applicable for the following dosage forms:

(1) solutions enclosed in single-dose containers and in soft capsules;

(2) solids (including powders, granules and sterile solids) that are packaged in single-dose containers and contain no active or inactive added substances;

(3) solids (including sterile solids) that are packaged in single-dose containers, with or without active or inactive added substances, that have been prepared from true solutions and freeze-dried in the final containers and are labelled to indicate this method of preparation;

(4) hard capsules, uncoated tablets, or film-coated tablets, containing 25 mg or more of an active substance comprising 25 per cent or more, by mass, of the dosage unit or, in the case of hard capsules, the capsule contents, except that uniformity of other active substances present in lesser proportions is demonstrated by meeting content uniformity requirements.