AMRL#:		
AWINL #.		

AGGREGATE WORKSHEET INDEX REPORT #: _____

Test Method	AASHTO/ASTM	Page	Technician	√ or NP
SAMPLING AGGREGATES	T2 / D75	2		
MINUS NO. 200 WASH	T11 / C117	3		
UNIT WEIGHT	T19 / C29	4		
ORGANIC IMPURITIES	T21 / C40	6		
SIEVE ANALYSIS OF AGGREGATE	T27 / C136	7		
SIEVE ANALYSIS OF MINERAL FILLER	T37 / D546	9		
SPECIFIC GRAVITY OF FINE AGGREGATE	T84 / C128	10		
SPECIFIC GRAVITY OF COARSE AGGREGATE	T85 / C127	12		
LOS ANGELES ABRASION	T96 / C131	13		
SULFATE SOUNDNESS	T104 / C88	15		
CLAY LUMPS & FRIABLE PARTICLES	T112 / C142	18		
LIGHTWEIGHT PIECES	T113 / C123	19		
SAND EQUIVALENT	T176 / D2419	21		
AGGREGATE DURABILITY INDEX	T210 / D3744	26		
REDUCING FIELD SAMPLES	T248 / C702	31		
TOTAL MOISTURE CONTENT	T255 / C566	32		
UNCOMPACTED VOID CONTENT	T304 / <i>C1252</i>	33		
RESISTANCE TO ABRASION BY MICRO- DEVAL (COARSE AGGREGATE)	T327 / D6928	36		
FRACTURED PARTICLES	T335 / D5821	39		
LOS ANGELES ABRASION FOR COARSE AGGREGATE	/C535	41		
FLAT & ELONGATED PARTICLES	/D4791	43		
SPECIFIC GRAVITY USING INFRARED	/D7172	45		
DENSITY/ABSORPTION OF AGG. USING VACUUM SATURATION / SUBMERSION	/D7370	47		
RESISTANCE TO ABRASION BY MICRO- DEVAL (FINE AGGREGATE)	/D7428	53		

^{**} NP for Not Presented

^{★ -} Indicates the line has been modified since the version of the worksheets dated 2013-09-13.

SAMPLING AGGREGATES

(T2)	
(D75)	

		Date:
_		hod (complete applicable sections):
l.	<u>Flowi</u>	ng Aggregate Stream (Bins or Belt Discharge)
	(a)	Each increment taken from entire cross-section of the material as it is discharged?
		Note: It is usually necessary to have a special device constructed for use at each particular plant,
		consisting of a pan of sufficient size to intercept the entire cross section of the discharge stream and
		hold the required amount of material without overflowing.
	Conve	eyor Belt Sampling
	(a)	Conveyor belt stopped while the sample increments are being obtained?
	(b)	Two templates placed on the belt so that the material between them will yield one increment?
	(c)	All material between templates carefully scooped into container including any fines?
		Note – automatic belt samplers may be used if properly maintained and if regular inspection ensures all
		material is being removed from the belt. ★
3.	Stock	<u>piles</u>
	(a)	Every effort made to sample from stockpiles using appropriate power equipment, using it to create a
		separate small sampling pile following the instructions and diagrams in the test method?*
	(b)	Where power equipment is not available, samples from stockpiles made of at least three increments of
	(-)	material from the top third, mid-point, and bottom third of the stockpile?
	(c)	If sampling fine material without power equipment, outside layer of pile removed before sampling?★
	(d)	Bias in test results due to segregation of materials avoided?
	` '	portation Units (trucks, rail cars, barges, etc.), typically avoided
•	(a)	If sampling of these transportation units cannot be avoided, sampling plan for specific case created?
	(b)	Sampling plan defines number of samples necessary to represent lots of specific sizes?
	` /	
		ways (Bases and Sub-bases)
	(a)	All increments taken from roadway for full depth of material?
	(b)	Specific area of increment to be taken clearly marked (such as by a metal template)?
All M	lethods:	
	Samp	les obtained from finished product (when practicable)?
2.	Samp	le for abrasion testing not subjected to further crushing or reduction in particle size (unless the
	sampl	e is of such a size that it requires further reduction for testing purposes)?
		ial to be sampled visually inspected for variations and if any variations are noted, corrective action
		establish homogeneity in the material prior to sampling?
	tuito ti	residential frontogenerty in the indicental prior to sampling.
4.	At lea	st three approximately equal increments selected at random from the material and combined to form
••		sample (see table below for recommended size)?
	Note t	o assessors: For stockpiles and transportation units, number of increments should be based on the sampling plan.
	11016	s assessors. For stockplies and transportation unus, number of increments should be bused on the sampling plan.

	Recommended Min.	Recommended Min.	Recommended Min.	Recommended Min.
Aggregate size	Mass (kg)	Mass (lb)	Volume (L)	Volume (Gal)
\leq 9.5 mm (3/8 in.)	10	22	8	2
12.5 mm (1/2 in.)	15	35	12	3
19.0 mm (3/4 in.)	25	55	20	5
25.0 mm (1 in.)	50	110	40	10
37.5 mm (1 ½ in.)	75	165	60	15
50 mm (2 in.)	100	220	80	21
63 mm (2 ½ in.)	125	275	100	26
75 mm (3 in.)	150	330	120	32
90 mm (3 ½ in.)	175	385	140	37

Note: For combinations of coarse and fine aggregate, recommended size is coarse aggregate mass plus 10 kg (22 lb).

5.	Field samples	reduced to test	ing size in ac	cordance w	ith standard p	rocedures	s (T248/C	C702)?	·····	
_			_							

COMMENTS (T2 / D75): (T2 / D75)

MATERIALS FINER THAN 75- μm (No. 200) SIEVE IN MINERAL AGGREGATES BY WASHING

(T11)	
(C117)	

			<u>APPARATUS</u>		Date:	
1.			0.1% of sample mass?			
2.	Sieves (Nest of two):	(a) (b)	75-μm (No. 200)?	8) to 1.1	'8 mm (No. 16)?	
3. 4.			5? 9°F)?			
5.	Wetting agent (Method	d B only)?				
6.	Mechanical washing a (a) Results are co (b) Degradation	onsistent w	optional): ith those obtained using manual methods? ble is avoided?			
			<u>PROCEDURE</u>			
1. 2.	Test sample mass conf	forms to fo e nominal	•	ted is n	ot listed below,	
	100		Nominal Maximum Size	Mi	nimum Mass, g	
			No.4 or finer		300	
		9.5 mm	(3/8 in.) ASTM: Greater than No. 4 to 3/8 in		1000	
		19.0 m	m (3/4 in.) ASTM: Greater than 3/8 to 3/4 in		2500	
		37.5 mm	(1½ in.) or larger ASTM: Greater than 3/4 in.		5000	
Note:	If same sample is to be to	ested as in T	27 (C136), minimum mass should conform to rec	uirement	ts of that method.	
3. 4. 5. 6. 7. 8. 9. 10. 11. 12. 13. 14.	Test sample mass dete Placed in container an Optional: Wetting ag Contents of container Complete separation of Wash water poured the Wash water free of continued underial on sieves returned Excess water decanted Washed aggregate dried Washed aggregate mass	rmined to ordered ent added? vigorously of coarse arrough siever arse particle ntil wash warned to was from was ed to constant of the constant	ss at 110±5°C (230±9°F)?) sieve?		
			Original dry mass			
COMN	MENTS (T11 / C117):					(T11 / C117)

BULK DENSITY ("UNIT WEIGHT") AND VOIDS IN AGGREGATE

(T19)	
(C29)	

Date: __

APPARATUS

l.	Unit we	ight measure:
	(a)	Top rim is smooth, watertight, and plane to 0.25 mm (0.01 in) when measure with feeler gauge?
	(b)	Interior wall of measure is a smooth and continuous surface (no open seams, large welds, etc)?
	(c)	Height is 80 to 150% of diameter?

Measure recalibrated at least annually or whenever the accuracy is called into question?......

Note to Assessors: The standardization requirements are included here because they are listed in the test method. If the laboratory is seeking accreditation, these issues will be covered in the R18 evaluation and the notes should be written under the quality system section. Only if they are not seeking R18 accreditation would you write a note here.

(e) Capacity and design of measure conforms to requirements in table below?

Capacity of measure	Nominal Max. size of	Min. thickness	Min. thickness top 38 mm	Min. thickness
	Aggregate	bottom	(1.5 inches) of wall	remainder of wall
$2.8 L (1/10 ft^3)$	12.5 mm (1/2 in.)	5.0 mm	2.5 mm	2.5 mm
9.3 L (1/3 ft ³)	25.0 mm (1 in.)	5.0 mm	2.5 mm	2.5 mm
$14 L (1/2 ft^3)$	37.5 mm (1.5 in.)	5.0 mm	5.0 mm	3.0 mm
$28 L (1 ft^3)$	75 mm (3 in.)	5.0 mm	5.0 mm	3.0 mm

2.	Tamping rod?
	(a) Round, straight steel rod approximately 600 mm (24 in.) [AMRL: ± 4 in] long.
	(b) 16 mm (5/8 in.) in diameter with hemispherical tip.
3.	<u>Shovel or scoop</u> ?
4.	Piece of plate glass (larger than the measure's diameter)?
5.	Grease, such as chassis or water pump grease suitable for forming a water-tight seal?
6.	Balance, graduated to at least 0.05 kg (0.1 lb) increments:
	(a) AASHTO: Readable to 0.1% of sample mass?
	ASTM: Accurate to 0.1% of test load?
7.	Thermometer, with a range of at least 10 to 32°C (50 to 90°F) and that is readable to at least 0.5°C (1°F)?
Sample	e Preparation
1.	Sample obtained by (T248 / C702), approx. 125 to 200% of quantity needed to fill the measure?
2.	Sample dried to essentially constant mass or at 110±5°C (230±9°F)?

COMMENTS (T19 / C29):

 $(T19\,/\,C29)$

BULK DENSITY ("UNIT WEIGHT") AND VOIDS IN AGGREGATE

(T19)	
(C29)	

	<u>PROCEDURE</u>	Date:
Observe	ve one of the following:	
	ing Procedure (up to 37.5-mm [1 ½-in.] particles)	
1.	Measure filled 1/3 full and leveled with fingers?	
2.	Aggregate rodded with 25 evenly distributed tamping strokes?	
3.	Tamping rod does not forcibly strike the bottom of the measure?	
4.	Tamping strokes limited to layer being tamped?	
5.	Measure filled with two more similar layers and third layer filled to overflowing (before	
6.	Surface leveled with the fingers or the straightedge (tamping rod)?	
7.	Average level surface obtained (aggregate projections above the rim balance the voids be	
8.	Net mass determined to the nearest 0.05 kg (0.1 lb)? (G)	
9.	Bulk density reported to the nearest 10 kg/m^3 (1 lb/ft ³)? {Bulk density = $(G - T) / V$ or	
10.	Void content (if determined) reported to the nearest 1 percent?	······ <u> </u>
	g procedure (37.5 to 150-mm [1 ½ to 5-in.] particles):	
1.	Measure filled 1/3 full and leveled with fingers?	
2.	Layer compacted by raising alternate sides about 50 mm (2 in.) and dropping on floor 25	times
	on each side (a total of 50)?	
3.	Measure filled with two more similar layers and third layer filled to overflowing (before	compaction)?
4.	Surface leveled with the fingers or the straightedge (tamping rod)?	
5.	Average level surface obtained (aggregate projections above the rim balance the voids be	
6.	Net mass determined to the nearest 0.05 kg (1 lb)?	
7.	Net mass of aggregate multiplied by calibration factor or divided by volume of the meas	ure?
8.	Bulk density reported to the nearest 10 kg/m ³ (1 lb/ft ³)?	······
9.	Bulk density reported to the nearest 10 kg/m³ (1 lb/ft³)?	<u>-</u>
Shovelin	eling procedure (up to 150-mm [6-in.] particles): Note: This method only used when specified.	
1.	Measure filled to overflowing with scoop or shovel?	
2.	Aggregate discharged from height not exceeding 50 mm (2 in.) above top of measure?	
3.	Care taken to prevent segregation of the particle sizes?	
4.	Surface leveled with the fingers or the straightedge (tamping rod)?	
5.	Average level surface obtained (aggregate projections above the rim balance the voids be	elow the rim)?
6.	Net mass determined to the nearest 0.05 kg (0.1 lb)?	
7.	Net mass of aggregate multiplied by calibration factor or divided by volume of the meas	
8.	Bulk density reported to the nearest 10 kg/m ³ (1 lb/ft ³)?	
9.	Void content (if determined) reported to the nearest 1 percent?	······· <u> </u>
COMM	MENTS (T19 / C29):	(T19 / C29)

ORGANIC IMPURITIES IN FINE AGGREGATES FOR CONCRETE

(121)	
(C40)	

			APPARATUS Date:
1.		Glass	bottles:
1.		(a)	Clear (colorless) glass with outside dimension [ASTM: thickness] between 40 and 60 mm [ASTM: 38.1 to 63.5 mm] (1.5 to 2.5 in.)?
		(b)	Approximately 240 to 470-mL (8 to 16 oz) nominal capacity?
		(c)	Graduation lines in milliliters or ounces?
			Note: If bottle is unmarked, lines may be scribed onto the bottle and are required only at the 75, 130 &
		(1)	200-mL (2 ½, 4 ½, & 7-oz) levels. (Lines at 2 ½ oz only necessary when using the standard color solution.)
		(d)	Stoppers or caps which are not soluble in specified reagents?
2.		Reage	ent, 3 parts NaOH (sodium hydroxide) to 97 parts water by mass [ASTM: weight]?
3.		Refere	ence color standards (One of the following):
		(a)	Standard solution:
			Note to assessor: The standard color solution is a hazardous material and should not be handled without the proper safety equipment and training. Consult the SDS document for details.
			(1) Reagent grade Potassium Dichromate dissolved in concentrated sulfuric acid?
			(2) Equal to Organic Color No. 3?
			(3) Solution is freshly made (less than 2 hours old)?
,	or	(b)	Glass color plate with Organic Color Nos. 1-5 (Gardener Color Nos. 5, 8, 11, 14, & 16)?
			<u>PROCEDURE</u>
1.		Samp	le obtained by Method (T248 / C702)?
2.			ITO only: If sample is dried prior to testing, is it dried only by air drying?
3.		Sampl	le mass about 450 g (1 lb)?
4.		Sand a	added to the 130-mL (approximately 4 ½-ozs) level in the bottle?
5.		NaOH	I solution added until volume of fine aggregate and liquid, after shaking, is 200 mL (approx. 7 oz) level?
6.			stoppered and shaken vigorously?
7.			ed to stand for 24 hours?
8.			comparison made against color standards?
9.		soluti	or is darker than organic plate No. 3 (Gardner Color Standard No. 11) or darker than the standard color ion, is the material considered to possibly contain injurious organic compounds and further tests acted before the aggregate is used in concrete?
		Conaa	
COM	ИM	ENTS	(T21 / C40): (T21 / C40

SIEVE ANALYSIS OF FINE AND COARSE AGGREGATES

(T27)	
(C136)	

		<u>APPARA</u>	<u>rus</u>	Date:	
1.	ASTM: Fine agg: Bat	able to 0.1% of sample mass? lance, readable to 0.1 g, accur lance, readable & accurate to	ate to 0.1 g or 0.1% of	test load (greater)?	
2.		C (230±9°F)?			
3.	(b) Shaker runs for control Note to assessors: Check that either to	aker, Manufacturer / ID #: ients the particles on the sieving correct amount of time (determine). The time set on the dial may not be the set time and the actual time run- ring assessment:	ned during annual star te the same amount of time are the same, or that the	dardization)?e that the shaker will operate laboratory is aware of any o	 e. effset.
	(c) Sieving accuracy	y met in a reasonable time periodess of approx. 10 minutes may resu	d?		
4.	Sieves: circle which type((s) were used: 8-in. si	eves 12-in. sieve	s Other (such as sq	uare sieves)
Note to	Aggoggover Please evaluate n	PROCEI procedure(s) and sieving efficiency f		ione chakara if applicable	
		Mixtures of Coarse and Fine	-		
Cuarso				_	
1. 2. 3. 4. 5.	Sample obtained by (T24: Minimum sample mass: 2 in 20 kg; 2 ½ in 35 Sample dried to constant AASHTO only: Mass dete	8 / C702) or whole field sample $3/8$ in 1 kg; $1/2$ in 2 kg; $3/8$ kg; 3 in 60 kg; $3\frac{1}{2}$ in 10 mass at 110 ± 5 °C (230 ± 9 °F) ermined to nearest 0.1% (unless not forced to pass through open	e used?	kg; 1 ½ in 15 kg; coarse agg. only)?	······
6.	during one minute of cont	ued until not more than 0.5% by tinuous hand sieving (check by d until not more than one mas	hand with 8-in. diamet	er sieve)?	
		ute of continuous hand sievin			
	AASHTO: Sieve Size	Initial specimen mass	Mass passing	sieve % Passing	
	ASTM: Sieve Size	Mass on sieve before hand-ch	eck Mass passing	sieve % Passing	
7. 8. 9.	Fine sieves: mass of resid sieving surface (200 g for Coarse sieves: mass of residue). This is note to Assessors: This is note to Assessors:	ighed to 0.1% of original dry nue on each sieve [finer than 4.7 8-in. diameter sieve; 469 g for sidue on each sieve [for 4.75-m) * (effective sieving area [which to identical to (T30/D5444), they	5-mm (No. 4) sieves] 12-in. diameter sieve) m (No. 4) sieves and la ch is <u>smaller</u> than its nature calculated differently.	does not exceed 7 kg/m ² o? arger] does not exceed aminal diameter], m ²)?	f
	Sieve < #4	Opening (mm) < 4.75	Mass (g) – 8 in. dia.	Mass (g) – 12 in. dia.	
	#4	4.75	338	796	
	1/4 in. 3/8 in.	6.3 9.5	449 677	1055 1592	
	1/2 in. 3/4 in.	12.5 19.0	891 1354	2094 3183	
10.	Total mass of material after	er sieving agrees with mass bef	ore sieving to within 0	.3% (If not, do not	
11.	Percentages calculated to	the nearest 0.1% and reported	to the nearest whole nu	mber (except	
12. COMN		than 10%, percentage -200 reporting original dry sample mass, incl		m fraction from (T11 / C1	

COMMENTS (T27 / C136):

SIEVE ANALYSIS OF FINE AND COARSE AGGREGATES

(127)	
(C136)	

	Initial Mass:			Final Mass:		
	e obtained by (T24	48 / C702) or whole field sam	nple used.	minimum samı	ple mass 30	00 g?
Janpi		mass at $110 \pm 5^{\circ}$ C (230 ± 9°				
$\Lambda \Lambda CH$	TO only: Mass dat	ermined to nearest 0.1% (uni	loss alroad	lv datarminad i	n (T11 / C1	117))2
111511	10 omy. mass act	crimined to nearest 0.170 (inte	iess airead	y acterminea i	n (1117 C1	.17))
4ASH	TO: Sieving contin	ued until not more than 0.5%	6 by mass	of the total spe	cimen pass	es a given sieve
luring	one minute of con	ntinuous hand sieving (check	by hand w	rith 8-in. diame	eter sieve)?	
		ed until not more than one n				
		inute of continuous hand sie				
AAS	HTO: Sieve Size	Initial specimen mas	SS	Mass passin	g sieve	% Passing
AS	TM· Sieve Size	Mass on sieve hefore hand	d-check	Mass passin	o sieve	% Passino
Residu	not overloaded:	Mass on sieve before hand eighed to 0.1% of original dr	ry mass?			
Residu	ne on each sieve wo not overloaded: Mass of residue sieving surface Mass of residue 2.5 * (sieve ope	eighed to 0.1% of original dr on each sieve [finer than 4.7 (200 g for 8-in. diameter sieve on each sieve [for 4.75-mm ening, mm) * (effective sievir	y mass? 75-mm (No ve; 469 g f (No. 4) sio ng area (wh	o. 4) sieves] do or 12-in. diame eves and larger hich is <u>smaller</u>	es not exce eter sieve)?] does not of than its no.	eed 7 kg/m ² of
Residu Sieves (a)	ne on each sieve wo not overloaded: Mass of residue sieving surface Mass of residue 2.5 * (sieve ope	eighed to 0.1% of original dr on each sieve [finer than 4.7 (200 g for 8-in. diameter sieve on each sieve [for 4.75-mm ening, mm) * (effective sievir s: This is not identical to (T30/I	ry mass? 75-mm (No. ve; 469 g f (No. 4) sion area (wh. D5444), the	o. 4) sieves] do or 12-in. diame eves and larger hich is <u>smaller</u> y are calculated	es not exce eter sieve)?] does not e than its no. differently.	eed 7 kg/m ² of exceed minal diameter
Residu Sieves (a)	ne on each sieve wo not overloaded: Mass of residue sieving surface Mass of residue 2.5 * (sieve ope	eighed to 0.1% of original dr e on each sieve [finer than 4.7 (200 g for 8-in. diameter sieve on each sieve [for 4.75-mm ening, mm) * (effective sievir servires: This is not identical to (T30/10) Opening (mm)	ry mass? 75-mm (No. ve; 469 g f (No. 4) sion area (wh. D5444), the	o. 4) sieves] do or 12-in. diame eves and larger nich is <u>smaller</u> y are calculated (g) – 8 in. dia.	es not exce eter sieve)?] does not e than its no differently.	eed 7 kg/m ² of exceed minal diameter
Residu Sieves (a)	ne on each sieve we not overloaded: Mass of residue sieving surface Mass of residue 2.5 * (sieve ope	eighed to 0.1% of original dr on each sieve [finer than 4.7] (200 g for 8-in. diameter sieve on each sieve [for 4.75-mm] ening, mm) * (effective sieving) S: This is not identical to (T30/1) Opening (mm) < 4.75	ry mass? 75-mm (No. ve; 469 g f (No. 4) sion area (wh. D5444), the	o. 4) sieves] do or 12-in. diame eves and larger nich is smaller y are calculated (g) – 8 in. dia.	es not exce eter sieve)?] does not e than its no differently.	eed 7 kg/m ² of exceed minal diameter
Residu Sieves (a)	ne on each sieve we not overloaded: Mass of residue sieving surface Mass of residue 2.5 * (sieve ope Note to Assessor Sieve < #4 #4	eighed to 0.1% of original dr on each sieve [finer than 4.7 (200 g for 8-in. diameter sieve on each sieve [for 4.75-mm ening, mm) * (effective sievir s: This is not identical to (T30/1 Opening (mm) < 4.75 4.75	ry mass? 75-mm (No. ve; 469 g f (No. 4) sion area (wh. D5444), the	o. 4) sieves] do or 12-in. diameters and larger nich is smaller y are calculated (g) – 8 in. dia.	es not exce eter sieve)?] does not e than its not differently. Mass (g)-	eed 7 kg/m ² of exceed minal diameter
Residu Sieves (a)	ne on each sieve we not overloaded: Mass of residue sieving surface Mass of residue 2.5 * (sieve ope	eighed to 0.1% of original dr on each sieve [finer than 4.7] (200 g for 8-in. diameter sieve on each sieve [for 4.75-mm] ening, mm) * (effective sieving) S: This is not identical to (T30/1) Opening (mm) < 4.75	ry mass? 75-mm (No. ve; 469 g f (No. 4) sion area (wh. D5444), the	o. 4) sieves] do or 12-in. diame eves and larger nich is smaller y are calculated (g) – 8 in. dia.	es not exce eter sieve)?] does not e than its not differently. Mass (g)-	eed 7 kg/m ² of exceed minal diameter
Residu Sieves a)	ne on each sieve we not overloaded: Mass of residue sieving surface Mass of residue 2.5 * (sieve ope Note to Assessor Sieve < #4 #4 #4 #4 #4 #4 #4 #4 #4 #4	eighed to 0.1% of original dr on each sieve [finer than 4.7 (200 g for 8-in. diameter sieve on each sieve [for 4.75-mm ening, mm) * (effective sievir S: This is not identical to (T30/1 Opening (mm) < 4.75 4.75 6.3	ry mass? 75-mm (No. ve; 469 g f (No. 4) sion area (wh. D5444), the	o. 4) sieves] do or 12-in. diame eves and larger nich is smaller y are calculated (g) – 8 in. dia.	es not exce eter sieve)?] does not e than its no. differently. Mass (g)- 4 7	eed 7 kg/m ² of exceed minal diameter 12 in. dia. 169 196

(T27 / C136)

SIEVE ANALYSIS OF MINERAL FILLER FOR ROAD AND PAVING MATERIALS

(T37)
(D546)

	APPARATUS Date:
	AFFARATUS Date.
1.	Sieves: 600 μm (No. 30), 300 μm (No. 50), and 75 μm (No. 200) and [ASTM Only: 1.18 mm (No. 16)]?
2.	Satisfactory water spray or rubber hose?
3.	Oven, maintains 110±5°C (230±9°F)?
4.	Balance: AASHTO: Class G2?
	ASTM: Class GP-1, readable to 0.01 g, capacity at least 200 g?
	<u>PROCEDURE</u>
1.	Sample obtained by (T248 / C702)?
2.	Sample mass at least 100 g?
3.	AASHTO: Sample dried to constant mass at 110±5°C (230±9°F) and mass determined to nearest 0.1 g?
	ASTM: Sample dried to constant mass at 110±5 °C (230±9 °F) and mass determined to nearest 0.01 g?
4.	Sample placed on specified nest of sieves: [ASTM Only: No. 16], No. 30, No. 50, No. 200?
5.	Material washed with stream of water until water coming through sieves is clear?
6.	Velocity of water not sufficient to splash sample over side?
7.	Care taken to avoid clogging of 75-µm sieve?
8.	Residue on each sieve dried to constant mass at 110±5°C (230±9°F) and mass determined?
9.	AASHTO only: Excess water decanted (if necessary) from washed samples only through the
	75-µm sieve (prior to drying)?
10.	Masses of material retained on each sieve calculated as a percentage of the original sample mass?
11.	Results reported as total percent passing each sieve to nearest 0.5%?
~~.	
COMM	IENTS (T37 / D546): (T37 / D546)

SPECIFIC GRAVITY AND ABSORPTION OF FINE AGGREGATE

(T84)	
(C128)	

					<u>APPARAT</u>	<u>'US</u>	Date:	
1	D	4						
1.	Pycnon				1	2		
	(a)						1. 1	
	(b)					> space required	d to accommodate test sample	le?
	(c) One of the following types of containers: (1) Volumetric flask, capacity 500 mL (or more)?							
		(1) (2)						
	or or	(3)						
	OI	(3)	(a)				etween highest graduation ma	
			(a)					
			(b)				o 24 mL?	
			(c)				dentification markings?	
			(d)					
			(e)				tion line (mL)?	
2.	Conical	mold	(0)	ome of capaci	ily marked doore	ingnest grada	tion fine (in2):	
2.	(a)		of metal	0.8 mm minimu	m thickness wit	h a height of 75	±3 mm?	
	(b)						00±3 mm?	
3.							ss of 340±15g?	
4.								
5.	Balance							
J.	Burance	<u> </u>					mple mass, sensitive to 0.1 g	
6.	Ontion	ıl AASH						
	2			9				
		7/			PROCED	<u>URE</u>		
		1/						
	Preparat							
1.							to 1200 g] in size?	
2.								
	Note: O	ven dryin	g not nec	essary if naturally	moist condition is	desired.		
3.	Allowed to cool to comfortable handling temperature [ASTM: approximately 50 °C]?							
4.	Covered with water or at least 6% moisture added?							
5.	Allowed to stand 15-19 hours [ASTM 20-28 hours], or naturally moist?							
6.	Excess	water de	ecanted (i	if necessary) with	nout loss of fines	i?		
7.	Sample spread on flat, nonabsorbent surface, and uniformly dried by current of warm air?							
8.	Mold placed on flat, nonabsorbent surface and filled to overflowing?							
9.	Tamper allowed to fall freely under gravitational attraction, 25 times with a 5 mm drop?							
10.								
10.	Loose sand removed from around base and mold lifted vertically?							
12.	Sample fails to slump on first test?							
13.							ole slumps slightly?	
13.	Drymg	Commu	za ana si	amp test repeated	a at mequent mite	i vais anui samp	no orampo ongmy:	
COMM	ENTS (1	84 / C1	28):					(T84 / C128)

SPECIFIC GRAVITY AND ABSORPTION OF FINE AGGREGATE

PROCEDURE (Continued)

(184)	
(C128)	

Date: _____

Proce	dure:
1.	Pycnometer partially filled with water, 500±10 g sample added, and SSD sample mass recorded? (S)
2.	Pycnometer filled to 90% of total capacity and agitated to eliminate air bubbles?
3.	Mechanical agitation permitted if performed in a manner that will not degrade the sample and comparison to
	manual agitation on the same material performed every 6 m, and the two results fall within the Table 1 range?
4.	Temperature of contents adjusted to 23.0±1.7°C (73.4±3°F) [ASTM: 23.0±2.0 \mathcal{C}]?
5.	Water level adjusted to calibrated capacity and mass of pycnometer and contents determined? (C)
	Note: Paper towel or isopropyl alcohol may be used to disperse foam on the water surface.
6.	Sample removed and dried to constant mass at 110±5°C (230±9°F)?
	Note , AASHTO only: Second sample taken at the same time, within 0.2 g of the sample placed in the pycnometer, may be used to determine the oven-dry mass.
7.	Sample cooled in air at room temperature for 1.0±0.5 hour and dry specimen mass determined? (A)
8.	Empty pycnometer filled to its calibration capacity with water at 23.0±1.7°C (73.4±3°F)
	[ASTM: 23.0±2.0 °C] and mass determined (pycnometer may be previously calibrated)? (B)
9.	All masses determined to nearest 0.1 g?
10.	Bulk specific gravity calculated as follows {Bulk sp gr = $A / (B + S - C)$ } and reported to nearest 0.001 (or
	reported to nearest 0.01 for fine aggregate meeting M6 requirements)?
11.	If sample tested in a naturally moist condition, source of the sample and the procedures used to prevent
	drying prior to testing reported?
Burett	e Method (AASHTO only)
	nate method to determine weight of pycnometer, specimen, & water)
1.	Mass of saturated surface-dry specimen determined? (S)
2.	Mass of empty pycnometer determined? (W)
3.	Specimen added to pycnometer as in Step 1 of Procedure?
4.	Water at 23.0 \pm 1.7 °C (73.4 \pm 3 °F) added to pycnometer from burette, quantity of water read from burette? (V_a) .
5.	Total mass of pycnometer, specimen, and water (C) calculated from equation $\{C = 0.9975 \ V_a + S + W\}$?
6.	Specific gravity reported to nearest 0.001 (or reported to nearest 0.01 for fine agg. meeting M6 reqs.)?
	atelier Method
(Alter	nate procedure)
1.	Le Chatelier flask filled with water to point on stem between 0 and 1 mL marks and initial volume recorded?
2.	Temperature of flask and contents at 23.0±1.7°C (73.4±3°F) [ASTM: 23.0±2.0 °C]?
3.	Approximately 55±5 g (other masses acceptable) of saturated surface dry fine aggregate added to flask?
4.	Separate 500±10 g sample of saturated surface-dry material taken for absorption determination?
	Note: This sample must be obtained at the same time as the sample is introduced into the Le Chatelier flask.
	Note , AASHTO only: Second sample taken at the same time, within 0.2 g of the sample placed in the pycnometer,
_	may be used to determine the oven-dry mass. Stopper placed in flesh, and flesh and contents egitated to remove entrepped sin?
5.	Stopper placed in flask, and flask and contents agitated to remove entrapped air?
6. 7	Flask and contents check to be within 1°C (1.8°F) of temperature in Step 3, water level read and recorded?
7.	ASHTO: aggregate removed from flask and dried to constant mass at 110 ± 5 °C $(230\pm9$ °F)?
0	ASTM: separate 500 g sample used to determine absorption?
8.	Lab says proper book formulas used in calculations and specific gravity reported to nearest 0.001 (or
	reported to nearest 0.01 for fine aggregate meeting M6 requirements)?
	trote to assessor. The specifically rejers to fine aggregates used in hydrautic cement and concrete applications.
COM	MENTS (T84 / C128): (T84 / C128)

Revised 2014-04-10

SPECIFIC GRAVITY AND ABSORPTION OF COARSE AGGREGATE

(T85)	
(C127)	

	APPARATUS Date:
1.	Sample container, Wire basket of 3.35-mm (No. 6) mesh or finer?
	4 to 7 L, for up to 37.5-mm (1 ½-in.) material OR a larger container that prevents trapping air when submerged for
2.	plus 37.5-mm (1 ½-in.) material. Water teals:
۷.	Water tank: (a) Capable of completely submerging the sample container?
	(a) Capable of completely submerging the sample container:
3.	Suspension apparatus:
	(a) Of suitable design and in good condition?
	(b) Center of suspension apparatus properly located with respect to center of balance pan or other point of contact with balance?
	(c) AASHTO only: Wire suspending the container is of smallest practical size?
4.	Immersion water, temperature is 23.0±1.7°C (73.4±3°F) [ASTM: 23±2.0 ℃]?
5.	Large absorbent cloth (paper towels or several small cloths NOT acceptable)?
6.	Balance: AASHTO: Class G5?
	ASTM: Sensitive, readable, and accurate to 0.05% of sample weight or 0.5 g (greater)?
7.	Sieves, 4.75 mm (No. 4) or other sizes as needed?
8.	Oven, maintains 110±5°C (230±9°F)?
	<u>PROCEDURE</u>
D	1
Proce	
1. 2.	Sample obtained by (T248 / C702)?
2. 3.	Sample mass as follows: 1/2 in. or less - 2 kg; 3/4 in 3 kg; 1 in 4 kg; 1 ½ in 5 kg?
<i>3</i> . 4.	Washed to clean surfaces of particles?
5.	Dried to constant mass at 110±5°C (230±9°F) and cooled to room temperature for 1 to 3 hours (for up
J.	to 1 ½-in. nominal maximum size, longer for larger sizes)?
	Note: Oven drying not necessary if naturally moist condition is desired.
6.	Covered with water for 15 to 19 hours [ASTM 20 to 28 hours]?
7.	Rolled in cloth to remove visible films of water? (A moving stream of air may be used to assist in the drying operation.)
8.	Larger particles wiped individually and evaporation avoided?
9.	AASHTO: If sample dries past SSD, sample immersed in water for 30 minutes and drying re-started?
10.	SSD sample mass in air determined? (B)
	(a) AASHTO: All masses determined to nearest 1 g or 0.1% of sample mass (whichever is greater)?
	(b) ASTM: All masses determined to nearest 0.5 g or 0.05% of sample weight (whichever is greater)?
11.	Sample immediately placed in sample container?
12.	Entrapped air removed before weighing by shaking container while immersed?
13.	Mass determined in water at 23.0±1.7°C (73.4±3°F) [ASTM: 23±2.0 °C]? (C)
14.	Dried to constant mass at 110±5°C (230±9°F) and cooled to room temperature for 1 to 3 hours (or
1.5	until aggregate has cooled to comfortable handling temperature, approximately 50°C)?
15.	Oven dried sample mass determined? (A)
16.	AASHTO: Bulk specific gravity calculated using the following formulas and reported to the nearest 0.001 (or
	nearest 0.01 for coarse aggregate meeting M80 requirements)?
	Note to assessor: M80 specifically applies to coarse aggregates used in concrete applications.
	Bulk specific gravity = $A/(B-C)$
	Bulk sp gr (by SSD mass) = $B/(B-C)$
	Apparent specific gravity = $A/(A-C)$
	Absorption = $[(B-A)/A] \times 100$
COM	MENTS (T85 / C127): (T85 / C127)

COMMENTS (T96 / C131):

RESISTANCE TO DEGRADATION OF SMALL-SIZE COARSE AGGREGATE BY ABRASION AND IMPACT IN THE LOS ANGELES MACHINE

(T96)	
(C131)	

		<u>APPARATUS</u>	Date	:		
Los A	ngeles machine					
(a)	Horizontal cylindrical drum, inside diameter 711 ± 5 mm (28 ± 0.2 in.), inside length 508 ± 5 mm (20 ± 0.2 in.), and wall thickness 12.7 ± 3.2 mm ($\frac{1}{2}\pm1/8$ in.) [ASTM: 12.7 mm, no tolerance]?					
(b)						
(c)	Cover for opening has dust-tight gasket and is securely fastened to drum?					
Interio	or Shelf requirements:					
(d)	Shelf projects inward 89±2 mm	(3.5±0.1 in.) or 152 x	102 x 12.7 mm (6 x 4 x ½in.)	?		
(e)	ASTM only: Interior surface of and steel spheres (except for the	f the cylinder free of proceedings	rotrusions disrupting the path	n of sample		
(f)	Shelf firm, rigid, and in good pl					
(g)	Shelf extends [AASHTO only: t					
(h)	Shelf located such that the char					
(i)	ASTM only: Distance from she	elf to the opening is 12	70 mm (50 in.) or more in the	?		
	direction of rotation?	••••••	••••••	•••••		
	on requirements:					
(j)	Uniform peripheral speed (± 1 .					
(k)	AASHTO only: Machine equipp					
(1)	Cylinder rotates at 30 to 33 rev	olutions per minute ove	r 5 minutes period?	•••••		
	Counter reading (Start):		Counter reading (End):			
	Elapsed time (minutes and seco					
	Average speed = $60 * (\# \text{ of revo})$	olutions) / time in secon	ds: RPM			
Chaus						
Charg		No. and a second	1i 200 44	F		
(a)	Number of spheres tested:	Number of spin	eres having a mass of 390-44	5 g:		
\	Mass of charge: Range	ge	Charge available?			
	A 12 spheres 49°	C				
	B 11 spheres 455					
	C 8 spheres 33	10 to 3350 g?				
	D 6 spheres 248	85 to 2515 g?				
(b)	All grading charges possible?					
Sieves	s. 1.70 mm (No. 12) and other size	s as needed?				
Balan	ce, AASHTO: Class G5, ASTM:	Accurate to 0.1% of te	st load?			
	ce, AASHTO: Class G5, ASTM: maintains 110 ± 5 °C $(230 \pm 9$ °F)?					

(T96 / C131)

RESISTANCE TO DEGRADATION OF SMALL-SIZE COARSE AGGREGATE BY ABRASION AND IMPACT IN THE LOS ANGELES MACHINE

(T96)_	
(C131)	

PROCEDURE	Date:

- 3. Mass determined to nearest 1.0 g?
- 4. Specimen masses conform to the table below?

SIEVE SIZE	GRADING A	GRADING B	GRADING C	GRADING D
1 to 1 ½ in	1250 ± 25 g			
3/4 to 1 in	1250 ± 25 g			
½to 3/4 in	$1250 \pm 10 \text{ g}$	$2500 \pm 10 \text{ g}$		
3/8 to ½in	$1250 \pm 10 \text{ g}$	$2500 \pm 10 \text{ g}$		
1/4 to 3/8 in			$2500 \pm 10 \text{ g}$	
No. 4 to 1/4 in			2500 ± 10 g	
No. 8 to No. 4				$5000 \pm 10 \text{ g}$
Total Mass	$5000 \pm 10 \text{ g}$	$5000 \pm 10 \text{ g}$	$5000 \pm 10 \text{ g}$	5000 ± 10 g

5.	Sample and spheres put in machine and tumbled 500 times?
	Note: Loss after 100 revolutions may be determined, and then entire sample returned to drum for final 400 revolutions.
6.	Contents of drum separated on a sieve coarser than a 1.70 mm (No. 12) according to T27/C136?
7.	Finer material separated on a No. 12 sieve?
	Note to assessors: Mechanical sieving or hand sieving is acceptable for this test.
8.	Material coarser than No. 12 washed and dried to constant mass at 110±5°C (230±9°F)? (See Note)
	Note: If material is essentially free of adherent coatings and dust, the requirement for washing is optional.
	For referee testing, the washing procedure must be performed.
9.	Mass of material coarser than No. 12 determined to nearest 1 g?
10.	Percentage of wear calculated as: % wear = (original - final mass) / original mass?

COMMENTS (T96 / C131):

(T96 / C131)

SOUNDNESS OF AGGREGATE BY USE OF SODIUM SULFATE OR MAGNESIUM SULFATE

(T104)	
(C88)	

		<u>APPARATUS</u>	Date:
			method?
		n be written here if the oven was checked otes under R18 (if applicable).	d incorrectly or if it has never been checked.
Piease	e write ali recora interval n	otes unaer K18 (ij applicable).	
Sieve	<u>s:</u>		
63.0 r	nm (2 ½ in.)?	_ 16.0 mm (5/8 in.)?	2.36 mm (No. 8)?
ľ	nm (2 in.)?		1.18 mm (No. 16)?
37.5 r	nm (1 ½ in.)?	9.5 mm (3/8 in.)?	.600 mm (No. 30)?
i	nm (1 1/4 in.)?		.300 mm (No. 50)?
25.0 r	nm (1 in.)?	_ 4.75 mm (No. 4)?	.150 mm (No. 100)?
19.0 r	mm (3/4 in.)?	_ 4.00 mm (No. 5)?	_
(a)	Is the laboratory pres	senting both coarse and fine method	s?
		e an informational Observation if the la	
		y	- emy pergerma ent memoral
Samn	le containers:		
(a)		(8-in) diameter sieves in good phy	vsical condition?
(4)		s are acceptable if they permit free acce	
		thout loss of aggregate. Sieves are requ	
<i>(b)</i>			ne agg - 250 μm (No. 60) size sieves?
(c)		are perforated to allow solution ac	
(0)			
	wunoui sampie ioss.	••••••	•••••
Culfor	to colution.		
	te solution:		
(a)	Solution containers:		16 110
			tion and foreign material?
1			depth of at least 12.5 mm (1/2 in.)?
			discolored solutions should be filtered)?
			ume of all samples immersed at one time?
(b)	Temperature regulati		
		controlling temperature of sulfate so	
	(2) Solution ter	nperature 20.3 to 21.9°C (68.5 to 71	1.5°F) [ASTM: 21±1°C (68 to 71.6°F)]?
(c)	Suitable device for n	neasuring specific gravity of solution	n to within ±0.001:
(d)			STM: 1.151 to 1.174]?
(e)	For Magnesium Sulf	ate, specific gravity is 1.297 to 1.30	6 [ASTM: 1.295 to 1.308]?
D -1 -			
$\frac{\text{Balan}}{(a)}$		a 0.1% of sample mass, or hotter?	
(<i>a</i>) (b)			
(0)	ASIM. FOR JULE US	greguie, accurate to 0.1 g	••••••
Domina	m ahlarida salutian.		
	m chloride solution:	(116 a RaCl non liter of colution)	
(a))
(b)		% barium chloride solution prepar	
Тогт	2 0 2	nL aisuuea waier?	
	erature recorder:	7	10
(a)		lution temperature a minimum of or	
	for duration of t	est and accurate to 0.3 \mathcal{C} (0.5 \mathcal{F})? .	
(b)	ASTM: accuracy of	at least 1°F (0.5°C) and can record	l temperature at least every 15 minutes?
AASH	ITO only: <u>Thermometer</u>	covers temperature range of solution	on and readable to 0.1 ${\mathcal C}$ (0.2 ${\mathcal F}$)?
	(T101 / G00)		
ENTS	(T104 / C88):		(T10

SOUNDNESS OF AGGREGATE BY USE OF SODIUM SULFATE OR MAGNESIUM SULFATE

(T104)	
(C88)	

		SAMPLE PREPARATION	<u>N</u> Date:		
Fine A	Aggregate:				
1.					
2.					
3.					
4.	Sample rough graded to obtain 11		ving sizes, if possible:		
	9.5 to 4.75 mm (3/8 in. to				
	4.75 to 2.36 mm (No. 4 to	,			
	2.36 to 1.18 mm (No. 8 to				
	1.18 to 0.600 mm (No. 10				
_	0.600 to 0.300 mm (No. 1		10		
5.			tested?		
6.	Each size sieved a second time to	rerusar/			
7.					
8.	100±0.1 g of each size weighed of	ut and put in separate containers	?		
Coore	se Aggregate:				
1.	Material finer than 4.75 mm (No.	1) removed?			
2.	•		°C (230±9°F)?		
2. 3.	By sieving to refusal, sample sepa		C (230±3 T)?		
٥.	63 to 37.5 mm (2 ½to 1 ½				
	37.5 to 19.0 mm (1 ½ to				
	19.0 to 9.5 mm (3/4 to 3/				
	9.5 to 4.75 mm (3/8 in. to				
4.	Weight of each fraction present as follows:				
	63 to 37.5 mm :		3000±300 g?		
	$(2 \frac{1}{2} \text{ to } 1 \frac{1}{2} \text{ in.})$	(2 ½ to 2 in.)	3000±300 g?		
	(2 /2 to 1 /2 m.)		2000, 200		
		50 to 37.5 mm			
		(2 to 1/2m.)			
	37.5 to 19.0 mm:		1000±50 g?		
	$(1 \frac{1}{2} \text{ to } 3/4 \text{ in.})$	$(1 \frac{1}{2} \text{ to } 1 \text{ in.})$			
		25.0 to 19.0 mm	500±30 g?		
		(1 to 3/4 in.)			
	19.0 to 9.5 mm:	19.0 to 12.5 mm	670±10 g?		
	(3/4 to 3/8 in.)	(3/4 to 1/2 in.)	070=10 g <u></u>		
	,	· · · · · · · · · · · · · · · · · · ·	330±5 g?		
		(1/2 to 3/8 in.)	330±3 g?		
	0.5.4.75	·			
	9.5 to 4.75 mm :		300±5 g?		
	(3/8 in. to No. 4)				
_	If annula annual as less than 500 of		44-19		
5.	ii sample contains less than 5% of	any specified size, that size not	tested?		
COM	MENTS (T104 / C88):		(T104 / C88		
COM	1VILITID (110+/ C00).		(1104 / C00		

SOUNDNESS OF AGGREGATE BY USE OF SODIUM SULFATE OR MAGNESIUM SULFATE

(T104)	
(C88)	

	PROCEDUR	<u>E</u> Date: _	
Proced	<u>edure</u>		
1.	Salt cake in bottom of solution container broken up and stirr	ed?	
2.	Specific gravity of solution checked?		
3.	Each sample immersed in depth at least 12.5 mm (½ in.) about	ve its top?	
4.	Samples kept immersed for 16 to 18 hours?		
5.	After removal from solution, each sample drained 10 to 20 i	ninutes?	······
6.	Dried to constant mass at 110±5°C (230±9°F)?		
7.	Cooled to room temperature [AASHTO only: 20 to 25 °C (68	to 77 °F)]?	
8.	AASHTO only: Temperature of aggregate checked by thermoplacement in sulfate solution?		
9.	Re-immersed and process continued until required number of		
	Note, AASHTO only: If test must be interrupted, samples should be	*	
10.	AASHTO only: Temperature records from recording unit rev		<u>.</u>
	temperature limits were not exceeded?		
11.	After final cooling, sample washed by circulating water at 4	3 ± 6 °C (110 ±10 °F) through the	
	samples inside their containers?		
12.	Hot water introduced near bottom and allowed to pass throu	gh samples and overflow?	
13.	Impact or abrasion of samples avoided during washing oper	ation?	
14.	Barium chloride used to check completeness of washing?		
	Note: If barium chloride reacts with lab water, completeness of wa		
15.	Each fraction dried to constant mass at 110±5°C (230±9°F)		
16.	<u>Fine Aggregate</u> : Sieved over same sieves used before test a originally, should be hand sieved at the end)?		
17.	Coarse Aggregate: Hand sieved over:		
	31.5 mm sieve for 63 to 37.5 mm?	8.0 mm sieve for 19.0 to 9.5 mm	n?
	(1 1/4 in. sieve for 2 ½ to 1 ½ in.)	(5/16 in. sieve for 3/4 to	o 3/8 in.)
	16.0 mm sieve for 37.5 to 19.0 mm?	4.00 mm sieve for 9.5 to 4.75 m	m?
	(5/8 in. sieve for 1 ½ to 3/4 in.)	(No. 5 sieve for 3/8 in.	to No. 4)
18.	Mass of material retained on each sieve determined?		<u></u>
COMM	IMENTS (T104 / C88):		(T104 / C88)

CLAY LUMPS AND FRIABLE PARTICLES IN AGGREGATE

(1112)	
(C142)	

	<u>APPARATUS</u>	Date:
1. 2. 3. 4.	Sample containers, rust resistant, shape such that sample can be spread in thin layer on be Sieves: 4.75 mm (No. 4), 1.18 mm (No. 16), and other sizes as needed?	······
	<u>PROCEDURE</u>	
Sample 1. 2. 3. 4.	PreparationSamples taken from materials left over from (T11 / C117)?Material dried to substantially constant mass at $110 \pm 5^{\circ}$ C ($230 \pm 9^{\circ}$ F)?Fine Aggregate, sample mass at least 25 g, consists of particles coarser than 1.18-mm (NCoarse Aggregate:(a) Consists of particles coarser than 4.75-mm (No. 4)?(b) Sample mass at least:No. 4 to 3/8 in. 1000 g?3/8 to 3/4 in. 2000 g?	o. 16) sieve?
5.	3/4 to 1 ½ in. 3000 g?	
Procedu 1. 2. 3. 4.	Sample mass determined to 0.1% and spread in thin layer on bottom of container?	
5.	Residue of friable particles removed by wet sieving as follows: (a) Fine aggregate on No. 20? (b) No. 4 to 3/8 in. on No. 8? (c) 3/8 to 3/4 in. on No. 4? (d) 3/4 in. and larger on No. 4?	
6. 7.	Residue from each sieving dried to constant mass at 110 ± 5 °C (230 ± 9 °F)?	
COMM	ENTS (T112 / C142):	(T112 / C142)

LIGHTWEIGHT PIECES IN AGGREGATE

(T113)	
(C123)	

		APPARATUS Date:
1.		valiquid AASHTO: One of the following. ASTM: A heavy organic liquid with an appropriate ic gravity and that can readily removed from the aggregate such as one of the following:
	(a)	Solution of zinc chloride in water for materials with specific gravity less than 2.0?
or	(b)	Solution of zinc bromide in water for material with specific gravity less than 2.6 [ASTM: 2.4]?
or	(c)	AASHTO: Mixture of kerosene with 1,1,2,2 tetrabromoethane for materials with specific gravity between 2.4 and 2.95 (must be used in a fume hood)?
		ASTM: Mixture of heavy organic liquids proportioned to achieve the desired specific gravity (suggested: tetrabromoethane, dibromoethane, 26 tetrachloroethance, or dichloromethane)?
	Note:	The chemicals specified in option "c" above are highly toxic – a fume hood, eye protection, and skin protection are needed when working with these chemicals. Zinc chloride and zinc bromide do not have dangerous fumes.
2.	Contai (a) (b)	iners: Suitable drying containers?
3.	Specif (a)	Suitable device for measuring specific gravity of heavy liquid within ± 0.01:
4.	Skimn (a) (b)	ner: Piece of 300-µm (No. 50) sieve cloth? Suitable size and shape?
5.	Hot pl	ate or oven, capable of maintaining temperature of 110±5°C (230±10°F)?
6.	Sieves	<u>s</u> : 300-μm (No. 50) and 4.75-mm (No. 4)?
7.	Balanc AASH' ASTM	ce: TO: Readable to 0.1% of sample mass, or better?
8.	Miscel (a) (b)	Ilaneous: Hood in good working condition (if kerosene mixture is used)?
COMN	MENTS ((T113 / C123): (T113 / C12

LIGHTWEIGHT PIECES IN AGGREGATE

(T113)	
(C123)	

	PROCEDURE Date:
Sample	e Preparation
1.	Sample obtained by (T248 / C702)??
2.	Minimum sample mass as follows?
	AASHTO: 4.75 mm (No. 4) - 200 g; 19.0 mm (3/4 in.) - 3 kg; 37.5 mm (1 ½ in.) - 5 kg; 75 mm (3 in.) - 10 kg.
	ASTM: 4.75 mm (No. 4) or smaller - 200 g; 9.5 mm (3/8 in.) – 1.5 kg; 12.5 to 19.0 mm (1/2 to 3/4 in.) - 3 kg;
	25 to 37.5 mm (1 to 1 1/2 in.) – 5 kg; 50 mm (2 in.) or larger – 10 kg.
3.	Aggregate dried to constant mass at 110±5°C (230±10°F) and cooled to room temperature?
	ggregate
1.	Sieved on a 300-μm (No. 50) sieve?
2.	Sieving continued until less than 1% of material on sieve passes in 1 minute of continuous hand sieving?
3.	Mass of plus 300-μm material determined to nearest 0.1 g?
4.	Aggregate brought to saturated surface-dry condition by T84/C128?
or	Amount of water that aggregate will absorb added, covered for 30 minutes, and tested?
	Note, AASHTO only: If material undergoes degradation in water, it does not have to be in SSD condition.
5.	Sample placed in container holding heavy liquid?
6.	Volume of heavy liquid at least 3 times the volume of aggregate tested?
7.	Liquid poured into second container through skimmer?
8.	Only floating particles decanted?
9.	Heavy liquid recovered and poured back into starting container?
10.	Aggregate agitated by stirring?
11.	Steps 7 through 10 repeated until all floaters are removed?
12.	Lightweight particles on skimmer washed free of heavy liquid using alcohol (for tetrabromoethane) or
	water (for zinc chloride or zinc bromide)?
13.	Lightweight particles allowed to air dry or dried to constant mass at no greater than 115°C (240°F)?
14.	Mass of lightweight particles determined to nearest 0.1 g?
15.	Lab says book formulas used in all calculations?
10.	
Coarse	e Aggregate
1.	Sieved on a 4.75-mm (No. 4) sieve?
2.	Mass determined to the nearest 1 g?
3.	Aggregate brought to saturated surface-dry condition by (T85 / C127)?
	Note, AASHTO only: If material undergoes degradation in water; it does not have to be in SSD condition.
4.	Sample placed in container holding heavy liquid?
5.	Volume of heavy liquid at least 3 times the volume of aggregate tested?
6.	Skimmer used to remove the floating particles and particles saved?
7.	Aggregate in container agitated?
8.	All floating particles removed by above process?
9.	Lightweight particles on skimmer washed free of heavy liquid using appropriate solvent?
10.	Lightweight particles allowed to air dry or dried to constant mass at no greater than 115°C (240°F)?
11.	Mass of lightweight particles determined to the nearest 1 g?
12.	Lab says book formulas used in all calculations? {% Lw = (dry mass of floating / sample mass) * 100}
- 4.	240 says cook formulas asca in an encountrions. [70 DW – (ary mass of mounting / sample mass) 100}

COMMENTS (T113 / C127):

(T113 / C127)

(T176)	
(D2419)	

			<u>APPARATUS</u>	Date:
1.	Gradu	ated plastic cylinders (at least three recor	nmended)?	
	(a)	Outside diameter 38.1 mm (1.5 in.).		
	(b)	Inside diameter 31.0 – 32.0 mm (1.25	in.).	
	(c)	Inside height 430 mm (17 in.).		
	(d)	Graduations at 2.54 mm (0.1 in.), mark	ked up to at least 15 in.	
	(e)	Rubber stopper that fits cylinder.		
2.	Satisfa			<u></u>
	(a)	Irrigator tube with an outside diameter	6.4 mm (1/4 in.) and le	ength approximately 510 mm (20 in.).
	(b)	Pinched end with No. 60 holes (1.0 mi	m diameter) drilled in tv	wo places on end.
3.	Weigh	nted foot assembly, weighs 1000 ± 5 g wi	th a guide fixed to the s	haft?
	Note:	Older (1969) model of weighted foot assembly		
	gradua	ated cylinder is acceptable.		
4.	Tin m	easure, diameter approximately 57 mm (2	2 1/4 in.) and capacity of	of 85±5 mL?
5.	Wide-	mouth funnel [AASHTO only: Diameter	approx. 100 mm (4 in.)	[AMRL: 3 to 5 in.] at the mouth]?
6.	Clock	or watch, readable in minutes and second	ds?	
7.	Shake	r (One of the following):		
		AASHTO only: Mechanical shaker required fo	or referee testing. Informa	utional note if mechanical shaker NP.
	(a)	<u>Mechanical</u>		
	//	(1) Operates at 175 ± 2 cycles pe	er minute (127 to 135 c	ycles during testing period)?
		(2) Securely fastened to firm and	l level mount?	
	(b)			
	10	(1) Securely fastened to firm and	l level mount?	
	(c)	Hand method		
	1	(1) Effective method of determin	ing 9 ± 1 in. throw leng	gth?
COM	MENTS ((T176 / D2419):		(T176 / D2419)
~ ~ 111		(, - , 1 - /), ·		(II/O/D211))

(T176)	
(D2419)	

Date: _____

APPARATUS (Continued)

	(a) 454 g (1 lb) technical grade anhydrous calcium chloride, 2050 g (4.515 lb) USP glycerin, and 47 g (0.10 lb) formaldehyde (40% by volume solution); diluted to 3.78 L (1 gallon) with		
	distilled or demineralized water?		
or	(b) 577 g (1.27 lb) A.C.S. grade calcium chloride dihydrate, 2050 g (4.515 lb) USP glycerin, and		
	59 g (0.13 lb) 1,5-pentanedial (glutaraldehyde) (50% solution in water); diluted to 3.78 (1 gallon) with distilled or demineralized water?		
or	(c) 577 g (1.27 lb) A.C.S. grade calcium chloride dihydrate, 2050 g (4.515 lb) USP glycerin, and 63 g (0.14 lb) kathon CG/ICP; diluted to 3.78 L (1 gallon) with distilled or demineralized water?		
	Note: Stock solution may be made without using any biocide (formaldehyde, glutaraldehyde, or kathon), provided the storage time of the stock solution is not sufficient to promote fungi growth.		
	provided the storage time of the stock solution is not sufficient to promote fungi growth.		
	Working calcium chloride solution:		
	(a) One measuring tin full (85±5 mL) of stock calcium chloride solution diluted to 3.78 L (1 gallon) with water?		
	(b) Stored in $\stackrel{?}{4}$ L (1 gallon) bottle on shelf 915 ± 25 mm (36 ± 1 in.) [ASTM: 90 ± 5 cm (36 ± 2 in.)] above work surface?		
	Note: Solution may be stored in larger glass or plastic vat, provided the liquid level is maintained between 915 to 1170 mm (36 and 46 in.) [ASTM: 36 and 45 in. (91 to 114 cm)] above work surface.		
	(c) Temperature of solution is $22 \pm 3^{\circ}\text{C}$ $(72 \pm 5^{\circ}\text{F})$?		
	(d) Solution is free of biological growth [ASTM: fungus]?		
	(e) AASHTO only: Solution discarded if it is not clear and transparent?		
	AASHTO only: Solution discarded if more than 30 days old?		
	ASTM only: Solution discarded if more than 2 weeks old, and fresh solution not added to old solution (Sections 6.6 to 6.8)?		
Э.	Oven, maintains 110 ± 5 °C (230 ± 9 °F)?		
1.	Work surface free of vibration and not exposed to direct sunlight?		
2.	4.75-mm (No. 4) sieve?		
3.	AASHTO only: Straightedge or spatula?		
4.	AASHTO only: Quartering or splitting cloth?		
5.	ASTM only: Flat pan, for mixing?		

COMMENTS (T176 / D2419):

(T176 / D2419)

(T176)	
(D2419)	

	PROCEDURE	Date:
Sample	Preparation Preparation	
AASHT		
1.	Sample obtained by T2, pulverized and passed through 4.75-mm (No. 4) sieve?	
2.	All fines cleaned from +No. 4 particles and included with -No. 4 material?	
3.	Sample split or quartered to yield 500 to 750 g (1.1 to 1.6 lb) of -No. 4 material?	
ASTM o		
1.	Sampled by D75, mixed and reduced according to C702 (splitting or quartering)?	
2.	Sample sieved on No. 4 (4.75-mm) sieve until not more than one weight percent of resi	
	passes the sieve during one minute?	
<i>3</i> .	Any +No. 4 lumps pulverized to pass No. 4 sieve?	
4.	All fines cleaned from +No. 4 particles and included with -No. 4 material?	
<i>5</i> .	Sample is at least 1500 g of -No. 4 material?	
Method	1 - Air Dry	
	necessary, material may be dampened before splitting or quartering to avoid segregation or loss of	fines.
AASHT	O only:	
1.	Enough -No. 4 material split or quartered to fill the 85-mL (3-oz) tin slightly rounded ab	ove brim?
2.	While filling, bottom edge of tin tapped on hard surface to consolidate material?	
3.	Tin struck off level full with spatula or straightedge?	
ASTM o	only (Procedure A):	
1.	Measuring tin filled four times by dipping from sample?	
2.	When a measure full is dipped, bottom edge tapped on hard surface at least 4 times to o	
3.	Measure level full or slightly rounded above the brim?	
4.	Amount of material in four measures determined by weight or by volume, using plastic	
5.	This material returned to sample?	
6.	Sample quartered or split according to C702 to obtain the predetermined weight or volve	ume?
7.	Sample split or quartered two more times to obtain specimens?	
8.	Each specimen dried at 230 \pm 9°F (110 \pm 5°C) and cooled to room temperature before	testing?
	2 - Pre-Wet (AASHTO and ASTM Procedure B)	
1.	ASTM only: Material dampened sufficiently to prevent segregation or loss of fines?	
2.	ASTM only: 1000 to 1500 g of material split or quartered out?	
3.	ASTM only: Material mixed thoroughly with hand trowel in circular pan by scooping a	
	middle of pan while rotating it horizontally?	
4.	ASTM only: Mixing continued for at least one minute?	
5.	Moisture condition checked by tightly squeezing small portion in palm of hand, forming	
6.	Sample at proper water content (cast permits careful handling without breaking)?	
	(a) If too dry (cast crumbles easily), water added and remixed?	
	(b) If too wet (shows free water), sample drained and air dried, mixing frequently?.	
7.	If either (a) or (b) above occurred, sample placed in pan, covered with lid or damp cloth	

(not touching sample), and allowed to stand for at least 15 minutes?.....

COMMENTS (T176 / D2419):

(T176 / D2419)

(T176)	
(D2419)	

Date: _____

PROCEDURE (Continued)

Method	2 - Pre-Wet (AASHTO and <i>ASTM Procedure B</i>) (Continued)		
1.	AASHTO: Sample placed on splitting cloth and mixed by alternately lifting each corner of cloth		
and pulling it over sample toward diagonally opposite corner, causing material to be rolled?			
	ASTM: Sample remixed for 1 minute after minimum curing time, without water, and		
	formed into a cone with a trowel?		
2.	AASHTO only: When material appears to be homogeneous, mixing finished with sample		
	in a pile near center of cloth?		
3.	Tin measure pushed through base of pile with free hand against pile opposite the measure?		
4.	Material fills tin to overflowing?		
5.	Material compacted into tin with palm of hand?		
6.	Tin struck off level full with spatula or straightedge [ASTM: or with trowel using a sawing motion]?		
Method	3 – AASHTO only Reference / Referee Method		
1.	AASHTO only: If using referee method (mechanical shaker), sample obtained by either Method 1 or 2, then		
	dried to constant mass at 110 ± 5 °C (230 ±9 °F), and cooled to room temperature before testing?		
Procedu	u <u>re</u>		
1.	101.6±2.5 mm (4±0.1 in.) of working calcium chloride solution siphoned into plastic cylinder?		
2.	Prepared sample poured from measuring tin into cylinder, using funnel to avoid spillage?		
3.	Bottom of cylinder tapped sharply on heel of hand several times to release air bubbles?		
4.	Wetted sample allowed to stand undisturbed for 10±1 minutes?		
5.	Stopper placed in cylinder and material loosened from bottom by shaking?		
6.	Mechanical Shaker Method (Referee Method):		
٠.	(a) Stoppered cylinder placed in mechanical shaker and timer set?		
	(b) Cylinder and contents shaken for 45 ± 1 seconds (127 to 135 cycles during testing period)?		
	(b) Cylinder and contents shaken for 13 ± 1 seconds (127 to 133 cycles during testing period)		
	Manual Shaker Method		
	(a) Stoppered cylinder secured in hand shaker and stroke counter reset to zero?		
	(b) Fingertips pushed against right hand spring steel strap, and smooth oscillating motion maintained?		
	(c) Tip of pointer reverses direction within marker limits?		
	(d) Shaking action continued for 100 strokes in 45 ± 5 seconds?		
(-)			
	Hand Method		
	(a) Cylinder held horizontally and shaken vigorously in horizontal linear motion from end to end?		
	(b) Cylinder shaken 90 cycles (one cycle is a complete back and forth motion) in approx. 30 seconds		
	[AMRL: ± 3 s.], using throw of 229±25 mm (9 ± 1 in.)?		
7.	Following shaking, cylinder set upright on work table and stopper removed?		
8.	Irrigator tube inserted in cylinder and material rinsed from cylinder walls as irrigator is lowered?		
9.	Irrigator forced through material to bottom of cylinder by gentle stabbing and twisting action		
<i>7</i> .	while solution flows from tip?		
10.	Stabbing and twisting motion applied until cylinder filled to 381-mm (15-in.) [ASTM: 38.0-cm] mark?		
11.	Irrigator raised slowly without shutting off flow so liquid level is maintained at about 15 in.?		
12.	Final level adjusted to 15 in. (381 mm) before irrigator is removed from cylinder		
	[AASHTO only: between top 2 graduations, but not above the 381-mm level]?		
13.			
14.	Timing started immediately after withdrawal of irrigator?		

COMMENTS (T176 / D2419):

(T176 / D2419)

(T176)	
(D2419)	

	PROCEDURE (Continued) Date:
Procedu	ure (Continued)
15.	After sedimentation, level at top of clay suspension (clay reading) recorded?
16.	If no clear line of demarcation, sample allowed to stand undisturbed until clay reading can be obtained, and total sedimentation time recorded?
17.	If sedimentation time exceeds 30 minutes, test rerun using 3 individual samples of same material, and clay reading requiring shortest sedimentation time recorded?
18.	Weighted foot assembly gently lowered into cylinder, without hitting mouth of cylinder?
19.	When foot rests on sand, assembly tipped toward cylinder graduations until indicator touches cylinder?
20.	254 mm (10 in.) subtracted from level indicated by extreme top edge of indicator, and this value recorded as sand reading?
21.	If clay/sand readings fall between 2.5-mm (0.1-in.) graduations, is level of higher graduation recorded?
<u>Calculat</u>	tions Sand equivalent calculated to 0.1 using following equation?
	Sand Reading x 100 Clay Reading
2. 3.	If sand equivalent is not a whole number, reported as next higher whole number? If desired to average sand equivalent values, and average is not a whole number, reported as next higher whole number?
	/// ANDI
COMM	ENTS (T176 / D2419): (T176 / D2419)

(1210)	
(D3744)	

		<u>APPARATUS</u>	Date:
1.	Mach	anical washing vessel (pot):	
1.		anical wasning vessel (pot): Flat-bottomed, straight-sided, flared-edge, and cylindrical, approx	vimataly 4 litar (2 gallon) canacity?
	(a)	Pot is 0.9 mm (20-gage) stainless steel with a gasket that is 3.2-m	
	(b)	Three trunk clamps, placed at 1/3 intervals and clamps attached to	
	(c) (d)	Lid forms watertight seal with flared edge of pot with gasket and	
	(u)	Lid forms watertight sear with mared edge of pot with gasket and	nd clamped in place:
2.		ction pan:	
	(a)	Round, with vertical or nearly vertical sides?	
	(b)	AASHTO: At least 250 mm (10 in.) in diameter and at least 100 m	nm (4 in.) deep?
		ASTM: At least 9 in. (230 mm) in diameter and approximately 4	4 in. (100 mm) deep?
	(c)	Holds wire mesh of 203.2-mm (8-in.) diameter sieve at least 76 n	nm (3 in.) above bottom?
		Note: A sieve frame resting on the bottom of the pan may be used.	
3.	Agita	tor, mechanical device capable of lateral reciprocating motion of 285	5 ± 10 complete cycles/minute
		a length of stroke 44.5 \pm 0.6 mm (1.75 \pm 0.025 in.) [ASTM: 45 \pm 6 m	
1.		C176 / D2419) equipment (covered in T176 / D2419 worksheets)?	
	(a)	Method C only: at least two sand equivalent cylinders?	
5.	Sieve	s: 19.0 mm (3/4 in.), 12.5 mm (½ in.), 9.5 mm (3/8 in.), 4.75 mm (N	
		 .18 mm (No. 16), and 75 μm (No. 200)?	
5.		ce, class G2, readable to 0.1g [ASTM: GP5, readable to 1 g, min. c	
7.		um chloride solutions, stock and working solutions as specified in (T	
	for re	feree testing the temperature of working solution is 22±3°C (72±5°F)?
3.	Distil	led or demineralized water, for referee testing the temperature of wat	ter is 22±3°C (72±5°F)?
9.		I only: Graduated cylinder, 1000 mL capacity?	
Note t	o Assesso	rs: It is preferable to observe Procedure A or Procedure C during the on-sit	te assessment.
		SAMPLE PREPARATION	
nitial	Sample	Preparation (all methods)	
l.		le obtained in accordance with (T2 / D75) (Sampling Aggregates)?	
2.		egate dried at temperature not exceeding 60°C (140°F), sufficiently to	
		75-mm (No. 4) sieve and to develop free-flowing condition in the por	
3.	If san	apple contains appreciable clay, aggregate turned frequently during dry	ying process?
1.	Hard clods broken up, fine coatings removed from coarse aggregate particles?		
5.	Gradi	ng determined by sieving in accordance with (T27 / C136) on 19.0, 1	12.5, 9.5, 4.75, 2.36 and
	1.18-1	mm (3/4 in., ½ in., 3/8-in., No. 4, No. 8, and No. 16) sieves?	···············
5.	Material retained on 19.0-mm (3/4-in.) sieve discarded?		
7.	Test F	Procedure (A, B or C) determined based on grading of aggregate?	
	(a)	If less than 10% aggregate passes 4.75 mm (No. 4), tested by Pro	
	(b)	If less than 10% aggregate is coarser than 4.75 mm (No. 4), tested	d by Procedure B only?
	(c)	If both coarse and fine aggregate fractions are each present in qua	antities $\geq 10\%$:
		(1) If percent passing 1.18 mm (No. 16) is greater than 10%	, both Procedures A and B
		used on appropriate aggregate sizes?	
		(2) If percent passing 1.18 mm (No. 16) is less than or equal	
		used on appropriate aggregate sizes?	······
	(d)	If most aggregate (75 - 80%) is between 9.5 and 1.18 mm (3/8 in Procedure C only?	and No. 16), tested by
COM	MENTS	(T210 / D3744):	(T210 / D37

Sample Preparation - Coarse Aggregate

AGGREGATE DURABILITY INDEX

(T210) _	
(D3744)	

Date:

PROCEDURE A – COARSE AG

Sam	pic i reparation	- Coarse Aggreg	<u>zaic</u>						
1.	Preliminary	test sample hav	ring mass of 2	$2550 \pm 25 \text{ g}$ (a	air-dry) prep	ared using	following table?	·····	

Aggregate Size	Air-Dry Mass, g
19.0 to 12.5 mm (3/4 to 1/2 in)	1070 ± 10
12.5 to 9.5 mm (1/2 to 3/8 in)	570 ± 10
9.5 to 4.75 mm (3/8 toNo.4)	910 ± 5

Note: If material has less than 10% of any size fraction, above masses adjusted to the actual percentage of the original grading, and sizes proportioned accordingly.

- 2. AASHTO only: Sample dried to constant mass at 110±5 ℃ (230±9 F), allowed to cool, and mass recorded?.... Note, AASHTO only: If an adjustment of the test specimen mass or volume of wash and test water, or both, is not required, it is not necessary to oven-dry the test sample prior to the initial wash.
- 3. Sample placed in mechanical washing vessel, and 1000 ± 5 mL distilled or demineralized water added?
- 4. Vessel lid clamped in place and vessel secured in sieve agitator?.....__________
- 5. When all aggregate is not completely inundated by water:
 - Material not inundated is washed and added to test sample?.....
 - Adjusted sample masses and water volumes used in testing when washed material is used? (b)
 - (c) Bulk, oven-dry specific gravity, and percentage of absorption of aggregate determined in accordance with (T85 / C127)?....
- 6.
- 7.
- 8. Vessel removed from agitator, lid unclamped, and contents poured onto 4.75 mm (No. 4) sieve?......
- 9. Remaining fines from vessel rinsed onto sieve and water (from a flexible hose attached to a faucet) directed onto the aggregate until water passing through the sieve is clear?......
- Material retained on 4.75 mm sieve dried to constant mass at 110±5°C (230±9°F) and mass determined?....... 10.
- If loss in mass due to washing is equal to or less than 75 g, skip to Procedure for Coarse Aggregate?..... 11.
- 12. If loss in mass exceeds 75 g:
 - (a) Preliminary test sample retained and combined with a second washed sample (by above washing procedure) according to the specified masses to provide the desired test sample?.....
 - (b) Grading for preliminary test sample determined using the following table – if each of the aggregate sizes listed in following table represents 10% or more of the 19.0 to 4.75-mm (3/4-in. to No. 4) portion, as determined from masses recorded in Step 6 of <u>Initial Sample Preparation</u> (previous page), are following oven-dry masses used in preparing the preliminary test sample?.....

Aggregate Size	Oven-Dry Mass, g
19.0 to 12.5 mm (3/4 to 1/2 in)	1050 ± 10
12.5 to 9.5 mm (1/2 to 3/8 in)	550 ± 10
9.5 to 4.75 mm (3/8 to No. 4)	900 ± 5
Total	2500 ± 25

- 2500 ± 25 g preliminary test sample prepared using the prescribed grading?.... (c)
- Test sample dried to constant mass at 110±5°C (230±9°F)?..... (d)
- Preliminary sample mechanically washed as in the same manner as the first specimen?..... (e)
- Steps repeated, if necessary, to obtain sufficient material to yield a washed test (f)
- sample of 2500±25 g and contain each size fraction in quantity specified in table above?..... After oven-dried material is allowed to cool, washed coarse aggregate separated on 12.5, 9.5 and (g)
- 4.75-mm (½ in., 3/8-in., and No. 4) sieves?.... (h)
- Material passing 4.75-mm (No. 4) sieve discarded?

(i)	Washed test sample prepared using masses specified in Step (b) above from representative
	portions of each size of washed material?

COMMENTS (T210 / D3744):

(T210 / D3744)

PROCEDURE A – COARSE AGG

(T210)	_
(D3744)	

Date: _____

Procedu	re for Coarse Aggregate
1.	Sand equivalent test cylinder placed on work surface free of vibration?
2.	7 mL (0.24 oz) of stock solution poured into cylinder?
3.	4.75 and 75-μm (No. 4 and No. 200) sieves placed in collection pan, with 4.75 mm on top?
4.	Washed test sample placed in mechanical washing vessel?
5.	Distilled or demineralized water added, lid clamped in place, and vessel secured in agitator?
6.	Agitation started 60 seconds after introduction of wash water?
7.	Vessel agitated for 600 seconds (10 minutes) ±15 seconds, immediately taken from agitator and lid removed?.
8.	Contents of vessel agitated by moving upright vessel vigorously in horizontal circular motion 5 or 6
	times to bring fines into suspension?
9.	Contents immediately poured over nested 4.75 and 75-µm (No. 4 and No. 200) sieves in collection pan?
10.	Material retained on 4.75 mm (No. 4) sieve discarded?
11.	All wash water and material passing 75-µm (No. 200) sieve collected in collection pan?
12.	To ensure all minus 75-μm material is washed through the sieve:
	(a) Jarring action applied to sieve by lightly bumping side of sieve frame with heel of hand as wash water drains through 75-µm sieve?
	(b) When concentration of material is retained on 75-µm sieve, fine material re-rinsed by pouring
	wash water through sieve again as follows:
	(1) Wash water allowed to stand undisturbed in collection pan for short time to permit
	heavier particles to settle to bottom?
	(2) Upper portion of wash water poured into another container?
	(3) Wash water poured back through 75-µm sieve, and all wash water and minus
	75-µm material collected in collection pan again?
13.	
	ASTM only: Wash water transferred to 1000-mL graduated cylinder?
14.	Distilled or demineralized water added to bring volume of dirty wash water to 1000±5 mL?
15.	Wash water transferred to container suitable for stirring and pouring?
16.	Funnel placed in sand equivalent cylinder?
17.	Wash water stirred by hand to bring fines into suspension?
18.	While water is still turbulent, enough wash water poured into cylinder to bring level of
	liquid to 381-mm (15-in.) mark?
19.	Funnel removed, stopper placed in end of cylinder, and contents mixed immediately?
20.	Contents mixed by alternately turning cylinder upside down and right side up, allowing bubble to
	completely traverse the length of the cylinder 20 times in approximately 35 [AMRL: \pm 5 s.] seconds?
21.	Cylinder placed on work table, stopper removed?
22.	Cylinder allowed to stand undisturbed for 1200 seconds (20 minutes) ±15 seconds?
23.	Height of sediment column immediately read and recorded to nearest 2.5 mm (0.1 in.)?
Calculat	tions for Procedure A – Coarse Aggregate
1.	Durability index calculated to nearest whole number using the following equation, or from Table 1
	$D_c = 30.3 + 20.8 \text{ cot } (0.29 + 0.15 \text{ H}) \text{ for H in inches.}$
	$D_c = 30.3 + 20.6 \text{ cot } (0.25 + 0.13 \text{ H}) \text{ for H in lineaes.}$ OR
	$D_c = 30.3 + 20.8 \cot (0.29 + 0.0059 \text{ H}) \text{ for H in mm.}$
	$D_{\rm c} = 30.3 \pm 20.0$ Cot (0.27 ± 0.0037) H) for H in limit.
COMM	ENTS (T210 / D3744): (T210 / D3744)

(T210)	
(D3744)	

	PROCEDURE B – FINE AGG Date:
Sample	e Preparation – Fine Aggregate
1.	Representative portion of 500±25 g obtained from minus 4.75 mm (No. 4) sieve oven-dry material?
2.	Preliminary test sample dried to constant mass at 110±5°C (230±9°F) and cooled to room temp.?
3.	Sample placed in mechanical washing vessel and 1000±5 mL distilled or demineralized water added?
4.	Vessel lid clamped in place and vessel secured in agitator?
5.	Agitation started 600 seconds (10 minutes) ±30 seconds after introduction of wash water?
6.	Vessel agitated for 120 seconds (2 minutes) ±5 seconds, vessel removed from agitator, and lid unclamped?
7.	Contents poured over 4.75 and 75-µm (No. 4 and No. 200) sieve nest?
8.	Any remaining fines rinsed from vessel onto sieve using water (from flexible hose attached to faucet)
	directed onto the aggregate until water passing through sieve is clear?
9.	If clayey or silty samples need to be flooded prior to pouring them over the sieve (to prevent clogging
	of the 75-µm (No. 200) sieve), flooded by adding water to vessel following agitation period?
10.	After rinsing, material transferred from sieve to drying pan?
11.	Pan left in slanted position until clear water can be decanted?
12.	Large shallow pans used and sample spread as thin as possible to speed drying?
13.	Sample dried to constant mass at 110±5°C (230±9°F)?
14.	After oven-dried material allowed to cool, sufficient amount of washed material split or quartered to fill
	85-mL (3-oz.) measuring tin to overflowing?
15.	Bottom of tin tapped on hard surface while filling?
16.	Tin struck off level full using straightedge?
17.	ASTM only: Mass of the material determined?
D 1	
<u>Proced</u> 1.	ure for Fine Aggregate (Procedure B)
1.	Procedure followed for Sand Equivalent Test (T176 / D2419), except agitator used to
	continuously shake cylinder and contents for 600 seconds (10 minutes) ±15 seconds?
Coloul	ation for Procedure P. Fine Aggregate
Caicui 1.	ation for Procedure B – Fine Aggregate Durability index calculated to nearest 0.1 using the following equation:
1.	Durability index calculated to hearest 0.1 using the following equation.
	$D_f = \underline{\text{sand reading}} \times 100$
	clay reading
2.	If D _f is not a whole number, is it reported as next higher whole number?
3.	If average series of values are desired, are whole number values averaged?
4.	If average of whole number values is not a whole number, rounded to next higher whole number?
COMN	MENTS (T210 / D3744): (T210 / D3744)

(1210)	
(D3744)	

	PROCEDURE C – Not Fine Not Coarse AGG Date:
Sample 1	Preparation – Not Fine Not Coarse Aggregate
1.	Sample contained between the 9.5 and 1.18-mm (3/8-in. and No 16) sieves?
2.	
۷.	Sample preparation followed as in Procedure B (Fine Aggregate)?
Procedu	re for Not Fine Not Coarse Aggregate
1.	Sand equivalent cylinder filled to 102.0±2.5 mm (4±0.1 in.) level with distilled or demineralized water?
2.	Prepared test sample poured into cylinder using funnel, avoiding spillage?
3.	Bottom of cylinder tapped sharply with heel of hand?
4.	Cylinder allowed to stand undisturbed for 10±1 minutes?
5.	Stopper placed on cylinder, material loosened from bottom, and cylinder placed in
	mechanical sand equivalent shaker?
6.	Contents agitated for 30±1 minutes?
7.	Water and passing 75-µm (No. 200) material transferred to another cylinder as follows:
	(a) 7 mL stock calcium chloride solution added to an empty second cylinder?
	(b) Nos. 8 and 200 sieves nested into a funnel that empties into the second cylinder?
	(c) Cylinder containing specimen and water held inverted over the nested sieves (with stopper in place)?
	(d) Stopper removed and contents poured over sieves?
	(e) Remaining fines rinsed from first cylinder onto sieves with small amount of fresh distilled water?
	(f) Material retained on the sieves rinsed with additional fresh distilled water until all minus 75-μm
	material passes through the sieve?
	(g) Care taken not to fill the second cylinder above the 380-mm (15-in.) mark?
	(h) Fresh distilled water added to bring level of liquid to 380-mm (15-in.) mark?
8.	Stopper placed on second cylinder and contents mixed by inverting 20 times in 35 seconds?
9.	Cylinder allowed to stand undisturbed for 1200 seconds (20 minutes) ±15 seconds?
10.	Top of clay suspension read to nearest 2.5 mm (0.1 in.)?
Calculat	ion for Procedure C – Not Fine Not Coarse Aggregate
1.	Durability index calculated to nearest whole number using the following equation, or from Table 1?
	$D_c = 30.3 + 20.8 \cot (0.29 + 0.15 \text{ H}) \text{ for H in inches.}$
	OR
	$D_c = 30.3 + 20.8 \text{ cot } (0.29 + 0.0059 \text{ H}) \text{ for H in mm.}$
COMP	CNTEG (TO10 / DO744).
COMMI	ENTS (T210 / D3744): (T210 / D3744)

REDUCING SAMPLES OF AGGREGATE TO TESTING SIZE

(T248)	
(C702)	

	<u>PROCEDURE</u>	Date:
Selection	on of Method	
1.	Fine aggregate, drier than saturated surface-dry – use Method A (Splitter)?	
2.	Fine aggregate, with free moisture present – use Method B (Quartering) or Method C (N	Miniature Stockpile)?
3.	Coarse Aggregate or Mixture of coarse and fine – use Method A (Splitter) or Method B	
	Method B or C is desired but sample does not have free moisture present, sample may be moistene	d and
	nly mixed before sample reduction.	
	moist sample is very large, preliminary split may be made using wide chute openings 38 mm (1 $\frac{1}{2}$	in.) or larger to
reduce s	ample to at least 5 kg. That portion should then be dried before sample reduction by Method A.	
Observe	e one of the following methods:	
	I A – Splitting	
1.	Material spread uniformly on feeder?	
2.	For fine aggregate (< 3/8 in.), maximum chute width is ¾ in. (19 mm)?	
3.	For coarse aggregate, minimum chute width is at least 50% bigger than maximum partic	cle size?
4.	Rate of feed slow enough so that sample flows freely through chutes?	
5.	Material in one pan re-split until desired weight is obtained?	······
Method	B - Quartering	
1.	Sample placed on clean, hard, and level surface?	
	Note: The sample may be placed upon a canvas quartering cloth or tear resistant tarp and a stic	
	placed under the cloth to divide the pile into quarters.	
2.	Mixed by turning over at least 3 times with shovel or by raising canvas or tarp and pulli	
3.	Thoroughly mixed conical pile formed?	
4.	Pile flattened to uniform thickness and diameter?	
5.	Diameter about 4 to 8 times thickness?	
6.	Divided into 4 equal portions with shovel or trowel?	<u></u>
	Note to assessors: The two unused quarters can be used for other testing.	
7.	Two diagonally opposite quarters, including all fine material, removed?	······
8.	Cleared space between quarters brushed clean?	
9.	Process continued until desired sample size is obtained?	
Method	C - Miniature Stockpile Sampling (Fine Aggregate Only)	
1.	Sample placed on clean, hard, and level surface?	
2.	Material thoroughly mixed by turning over three times?	
3.	Small stockpile formed?	
3. 4.	At least 5 grab samples taken at random with sampling thief, small scoop, or spoon?	······
-т.	11. Icase 5 grad samples taken at random with sampling tiner, sman scoop, or spoon?	······
COMM	IENTS (T248 / C702):	(T248 / C702)

COMMENTS (T255 / C566):

TOTAL EVAPORABLE MOISTURE CONTENT OF AGGREGATE BY DRYING

(T255)	
(C566)	

					<u>APPA</u>	<u>RATUS</u>		Date	e:	
1.		Heat sou	rce:							
		(a)	Ventilated oven,							
	or	(b)	If close temp cor	ntrol is not req	uired: hot plat	e, heat lamp	o, or ventilated	microwave of	oven?	
2.		Sample containers, not affected by heat source, of sufficient volume to hold samples, and has a shape so that the depth of the sample does not exceed 1/5 of least lateral dimension?					···			
3.		Stirrer, r	netal spoon or spa	atula of conve	nient size?					
4.		Balance.	readable to 0.1%	of sample ma	ss [ASTM: te	est load], or	better?			···
					PROC	<u>EDURE</u>				
1.		Represen	ntative test sample	e obtained with	h mass confor	ming to the	following table	e?		···
		No. 4	3/8 in.	½ in.	3/4 in.	1 in.	1 ½ in.	2 in	2 ½ in.	7
		0.5 kg	1.5 kg	2 kg	3 kg	4 kg	6 kg	8 kg	10 kg	
2. 3. 4. 5. 6. 7.		Loss of a Sample of If heated overheat Sample of Moisture		prior to determe heat source? Than a controlle ional for micromass and mass and by followin bisture = ori	nining the ma ed temperatur owave use)s determined t g equation?: . ginal sample dried sar	e oven, is sa o nearest 0.1 mass - dried	ample stirred to	avoid locali	zed	
		Note: If h	ot plate is used, de	natured alcohol	may be used to	burn off moi	sture.			

(T255 / C566)

UNCOMPACTED VOID CONTENT OF FINE AGGREGATE

(1304)	
(C1252)	

		APPARATUS Date:	
1.	Cylin	ndrical measure, approximately 100-mL capacity:	
1.	(a)	Volume calculated to nearest 0.1 mL [AMRL: calibrated capacity is 99.0 mL to 101.0 mL]?	
	(a) (b)	Inside diameter approximately 39 mm, inside height approximately 86 mm?	
	(c)	Made of drawn copper water tube, bottom at least 6 mm thick and firmly sealed to tubing?	
	(d)	Bottom provided with means for aligning axis of cylinder with axis of funnel?	
	(e)	Calibrated according to Section 8 with freshly boiled, deionized water at 18 to 24°C?	
2.	Funne	el, lateral surface of right frustum of a cone sloped $60\pm4^{\circ}$ from the horizontal:	
	(a)	Made of metal, smooth on inside and at least 38 mm high?	
	(b)	Opening diameter 12.7±0.6 mm?	
	(c)	At least 200 mL capacity or provided with supplemental glass or metal container to increase vol	
		Pycnometer top C9455 is satisfactory for funnel section, except size of opening has to be enlarged and any apparent burrs or lips should be removed by filing or sanding. Pycnometer top must be used with suitable glass jar with bottom removed.	
3.	Funne	el stand, 3 or 4 legged and holds funnel firmly in position:	
٥.	(a)	Aligns funnel with axis of cylindrical measure (within a 4° angle and a displacement of 2 mm)?	
	(b)	Funnel opening 115±2 mm above top of cylinder?	
4.	Glass	s plate for calibration of measure, at least 4 mm thick, approximately 60 by 60 mm?	
_	Elat a		
5.		metal or plastic pan, of sufficient size to contain the funnel stand and to prevent loss of material in filling the measure, and sufficiently flat to remain steady during testing?	
6.	Metal	ıl spatuala, straight edged:	
	(a)	Straight edge of blade approximately 100 mm long [AMRL: 3 to 6 in. long] and at least 20 mm	wide?
	(b)	End cut at right angle to edges?	
7.	Scale	e or balance, accurate and readable to ±0.1 g	
	1		
CO14	A CENTRO	(T204 / C1050)	004 / 01050

COMMENTS (T304 / C1252):

(T304 / C1252)

UNCOMPACTED VOID CONTENT OF FINE AGGREGATE

(T304)	
(C1252)	

				(C1252)
	PROCEDURE		Date:	
Sampling 1. Sample obtain	ined by splitting and quartering (C702), sieve analysis	s (C136), or from a	an extraction s	sample?
2. Methods A a (a) Sam (b) Sam		00) sieve in accord	lance with C1	17?
3. Method C: A	A split of the as-received sample dried in accordance	with the drying pro	ocedure of C1	36?
Sample Preparation				
1. Method A - S	Standard Graded Sample: Aggregate combined accor	ding to the followi	ng table?	
	Method A - Individual Size Fractions	Mass, g	OK?	
	2.36 to 1.18 mm (No. 8 to No.16)	44 ± 0.2		
	1.18 mm to 600 μm (No. 16 to No. 30)	57± 0.2		
	600 to 300 μm (No. 30 to No. 50)	72 ± 0.2		
	300 to 150 μm (No. 50 to No. 100)	17 ± 0.2		
	Total	190 ± 0.8		
2. Method B - I	Individual Size Fractions: 3 separate 190-g samples of	of aggregate tested	(see table)?	
	Method B - Individual Size Fractions	Mass, g	OK?	
1 51	2.36 to 1.18 mm (No. 8 to No. 16)	190 ± 1		
	1.18 mm to 600 μm (No. 16 to No. 30)	190 ± 1		
	600 to 300 μm (No. 30 to No. 50)	190 ± 1	(V-	[
	Note: These size fractions are tested separately. They sh	ould not be mixed di	uring testing.	
(a) Sam	As Received Grading hiple (dried in accordance with C136) passed through 90±1-g sample of material passing the 4.75-mm sieve			
COMMENTS (T304)	/ C1252):			(T304 / C125

UNCOMPACTED VOID CONTENT OF FINE AGGREGATE

(T304)	
(C1252)	

	PROCEDURE (Continued) Date:	_
Specifi	c Gravity of Fine Aggregate	
1.	If bulk dry specific gravity of aggregate from the source is unknown, specific gravity determined on material passing 4.75-mm (No. 4) sieve in accordance with C128?	
2.	This value used in subsequent calculations unless some size fractions differ by more than 0.05 from the specific gravity typical of the completed sample (in which case the specific gravity of the fraction(s) being tested must be determined)?	_
3.	If specific gravity differences exceed 0.05: (a) Specific gravity of the individual 2.36-mm (No. 8) to 150-μm (No. 100) sizes determined for	
	use with Method A or the individual size fractions for use with Method B?	
Proced	ure.	
1.	Each test sample mixed with spatula until it appears to be homogeneous?	_
2.	Jar and funnel section positioned in stand and cylindrical measure centered?	_
3.	Finger used to block opening of funnel while test sample is poured into funnel?	
4.	Material in funnel leveled with spatula?	
5. 6.	Finger removed and sample allowed to fall freely into cylindrical measure?	
7.	Spatula used with the blade width vertical and using the straight part of its edge in light contact with	_
<i>,</i> .	[ASTM: both sides of] the top of the measure?	
8.	Care used to avoid any disturbance that could cause compaction of aggregate into cylindrical measure?	
	Note: After strike-off, measure may be tapped lightly to make it easier to transfer without spilling any of the sample.	
9.	Adhering grains brushed from outside of container?	
10.	Mass of cylindrical measure and contents determined to nearest 0.1 g?	_
11. 12.	All aggregate particles retained and recombined for second test run?	
12. 13.	Sample from retaining pan and cylindrical measure recombined and procedure repeated? Mass of empty measure recorded?	
		_
Calcula	tions	
1.	Uncompacted voids for each determination calculated as follows:	-
	$U = \frac{V - (F/G)}{V} \times 100$	
	where:	
	V = volume of cylindrical measure, mL	
	F = mass of aggregate in measure	
	G = bulk dry specific gravity of aggregate	
	U = uncompacted voids in material, %	
2.	For Methods A and C, average uncompacted voids determined?	
3.	For Method B, average uncompacted voids for each size fraction determined and mean determined?	
COMM	IENTS (T304 / C1252): (T304 / C1252)	2)

RESISTANCE OF COARSE AGGREGATE TO DEGRADATION BY ABRASION (T327) _____ IN THE MICRO-DEVAL APPARATUS (COARSE AGGREGATE) (D6928) _____

	APPARATUS Date:
1.	<u>Micro-Deval Abrasion Machine</u> , a jar rolling mill similar to Fig 1, capable of running at 100 ± 5 rpm?
2.	Micro-Deval abrasion jars: (a) Stainless steel, 5 L capacity, with a rubber ring in the rotary locking cover?
3.	Abrasive Charge. (a) A total charge of 5000 ± 5 g presented for use in an abrasion jar?
	9.5 ± 0.5 mm 1 2 3 4 5 6 7 8 9 10
	Diameter ok?
4. 5. 6.	Sieves, 19.0 mm (3/4 in.), 16.0 mm (5/8 in.), 12.5 mm (1/2 in.), 9.5 mm (3/8 in.), 6.3 mm (1/4 in.), 4.75 mm (No. 4), and 1.18 mm (No. 16)?
	AASHTO <u>USE OF REFERENCE AGGREGATE</u>
1. 2. 3. 4. 5. 6.	Reference aggregate, mean loss between 15 to 25 %? 10 initial samples of reference aggregate taken at random and tested? Mean loss and sample standard deviation determined from the 10 initial samples? One reference aggregate sample tested for every 10 normal samples tested, but at least one reference sample tested in every week in which a regular sample is tested? For continued acceptance, individual test results of reference aggregate falls within ±2 standard deviations of the established mean percent loss 95% of the time? If the results fall outside of this range, is an investigation into the cause conducted? Percent loss of last 20 samples of calibration aggregate plotted on trend chart? After 10 weekly samples of reference aggregate are plotted, the interval may be increased to one reference sample per month.
	ASTM CALIBRATION
<u>Calibi</u> 1. 2.	Ration Supplies Brechin Quarry No. 2 aggregate, test data falls between 17.5 to 20.7 % loss for 95 % of the time?
1. 2. 3.	ation Procedure 10 samples of calibration aggregate taken at random and tested?
4. 5.	Caubration procedure conducted for new supplies of caubration aggregate, batched according to Section 8? Control sample tested every 10 samples, but at least every week in which a sample is tested?
5. 6 .	Percent loss of last 20 samples of calibration aggregate plotted on trend chart?

COMMENTS (T327 / D6928):

(T327 / D6928)

RESISTANCE OF COARSE AGGREGATE TO DEGRADATION BY ABRASION IN THE MICRO-DEVAL APPARATUS (COARSE AGGREGATE)

(T327)	
(D6928)	

Date: ____

SAMPLE	PREPAR	ATION
--------	--------	-------

Sample	Prepar	ration
--------	--------	--------

- 1. Test Sample washed and oven-dried to constant mass at 110±5°C?.....
- 2. Sample separated into individual size fractions in accordance with Test Method (T27 / C136)?.....
- 3. For material passing the 19.0 mm (3/4 in.) sieve:

Table A

Passing	Retained	Mass	OK?
19.0-mm (3/4 in.)	16.0-mm (5/8 in.)	375g	
16.0-mm (5/8 in.)	12.5-mm (1/2 in.)	375g	
12.5-mm (1/2 in.)	9.5-mm (3/8 in.)	750g	
Tot	al	$1500 \pm 5g$	

4. In a case where the nominal maximum size of the coarse aggregate is: less than 12.5 mm (1/2 in.):

Table B

Passing	Retained	Mass	OK?
12.5-mm (1/2 in.)	9.5-mm (3/8 in.)	750g	
9.5-mm (3/8 in.)	6.3-mm (1/4 in.)	375g	
6.3-mm (1/4 in.)	4.75-mm (No. 4)	375g	
Tot	al	$1500 \pm 5g$	

5. In a case where the nominal maximum size of the coarse aggregate is less than 9.5 mm (3/8 in.):

Table C

Passing	Retained	Mass	OK?
9.5-mm (3/8 in.)	6.3-mm (1/4 in.)	750g	
6.3-mm (1/4 in.)	4.75-mm (No. 4)	750g	
Tot	al	$1500 \pm 5g$	

Note to assessors – The 6.3-mm sieve may be replaced with a 6.7-mm sieve if desired.

COMMENTS (T327 / D6928):

(T327 / D6928)

RESISTANCE OF COARSE AGGREGATE TO DEGRADATION BY ABRASION IN THE MICRO-DEVAL APPARATUS (COARSE AGGREGATE)

(T327)	_
(D6928)	

-	n	\sim				T T		
ы	ĸ	()	Cl	Η'I	ווו	ш	ĸ	Η .

Procedu	ure
1.	Prepared sample weighed to nearest 1 g?
2.	Sample immersed in 2.0 ± 0.05 L of tap water either in Micro-Deval container or other suitable device?
3.	Temperature of tap water 20 ± 5°C and immersed for a minimum of 1 hour?
4.	Sampled placed in Micro-Deval abrasion container with 5000 ± 5 g of steel balls and the same
	water used to saturate the sample?
5.	Cover installed and Micro-Deval container placed on the machine?
6.	If machine is capable of recording total number of revolutions, machine run at 100 ± 5 rpm:
	(a) For $12,000 \pm 100$ revolutions for grading in Table A above?
	(b) For $10,500 \pm 100$ revolutions for grading in Table B above?
	(c) For $9,000 \pm 100$ revolutions for grading in Table C above?
7.	If machine is not capable of recording total number or revolutions, machine run at 100 ± 5 rpm:
	(a) For $2 h \pm 1$ min for grading in Table A above?
	(b) For 105 min \pm 1 min for grading in Table B above?
	(c) For 95 min ± 1 min for grading in Table C above?
8.	Sample and steel balls carefully poured over a 4.75-mm (No. 4) nested on a 1.18-mm (No. 16) sieve?
9.	Care taken to remove entire sample from the stainless steel jar?
10.	Retained material washed and manipulated using a held hand water hose and hand?
11.	Washed until all washing are clear and all material smaller than 1.18-mm (No. 16) passes that sieve?
12.	Stainless steel balls removed using a magnet or other suitable means?
or	ASTM only: Preferred method – sample and charge dried to constant mass before removal of charge?
13.	Material retained on the nest of sieves combined, with care taken not to lose any material?
14.	Sample dried to constant mass at 110 ± 5 °C and weighed to nearest 1 g?
15.	Micro-Deval abrasion loss calculated as follows?

Percent Loss = (A - B) / A * 100

where:

A = Initial sample mass

B = Final sample mass

COMMENTS (T327 / D6928):

(T327 / D6928)

DETERMINING THE PERCENTAGE OF FRACTURED PARTICLES IN COARSE AGGREGATE

(T335)	
(D5821)	

	APPARATUS Date:	
1. 2. 3.	Balance, accurate and readable to within 0.1% of sample mass? Sieves, conforming to M92 / E11? Sample splitter, as specified in T248 / C702?	
	<u>PROCEDURE</u>	
Sample 1. 2. 3. 4. 5.	Aggregate sampled in accordance with T2 / D75 and reduced in accordance with T248 / C702?	······
	ASTM: Minimum mass of sample is as indicated in the table below OR at least large enough so that the largest particle is not more than 1% of the sample mass (whichever amount is smaller)?	
6.	(Optional) ASTM only optional additional procedure For aggregate with nominal maximum size of 19.0 mm (3/4 in.) or larger, where the fracture particle content is to be determined for material retained on the 4.75-mm (No. 4) or smaller sieve: (a) Sample separated on the 9.5-mm (3/8-in.) sieve? (b) Portion passing 9.5-mm (3/8-in.) sieve further reduced by C702, to a minimum of 200 g (0.5 lb Note: This will reduce the number of particles to be separated during the procedure. (c) Percent fractured particles determined on each portion? (d) Weighted average percentage of fractured particles calculated based on mass of each of the portions to reflect total percentage of fractured particles in the entire sample?	o)?
Sample 1. 2. 3.	e Preparation (AASHTO Method 2 – Individual size fractions) Aggregate sampled in accordance with T2 and reduced in accordance with T248? Sample dried sufficiently to obtain clean separation of fine and coarse material in sieving operation? Sample meets on of the following: (a) A previously washed sample from the gradation determination (T11 and T27)? (b) Aggregate is sieved according T27 over the sieves listed in the specification for the material? Representative portion from each sieve selected by splitting or quartering to size specified in table below?	······

AASHTO	Method 1	AASHTO	Method 2	ASTM	
Nominal maximum	Minimum	Nominal maximum	Minimum	Nominal maximum	Minimum
particle size	mass g (lb)	particle size mm (in.)	mass g (lb)	particle size mm (in.)	mass g (lb)
37.5 (1 ½)	2500 (6)	31.5 (1 1/4)	1500 (3.5)	9.5 (3/8)	200 (0.5)
25.0(1)	1500 (3.5)	25.0(1)	1000 (2.2)	12.5 (1/2)	500 (1)
19.0 (3/4)	1000 (2.2)	19.0 (3/4)	700 (1.5)	19.0 (3/4)	1500 (3)
12.5 (1/2)	700 (1.5)	16.0 (5/8)	500 (1.0)	25.0(1)	3000 (6.5)
9.5 (3/8)	400 (0.9)	12.5 (1/2)	300 (0.7)	37.5 (1 ½)	7500 (16.5)
4.75 (No. 4)	200 (0.5)	9.5 (3/8)	200 (0.5)	50.0(2)	15,000 (33)
<u>-</u>	-	6.3 (1/4)	100 (0.2)	63.0 (2 ½)	30,000 (66)
		4.75 (No. 4)	100 (0.2)	75.0 (3)	60,000 (132)
		2.36 (No. 8)	25 (0.1)	90.0 (3 1/2)	90,000 (198)
		2.00 (No. 10)	25 (0.1)		

COMMENTS (T335 / D5821):

(T335 / D5821)

COMMENTS (T335 / D5821):

DETERMINING THE PERCENTAGE OF FRACTURED PARTICLES IN COARSE AGGREGATE

(T335)	
(D5821)	

	APPARATUS Date:
AASH	TO Procedure
1. 1511	Fractured face defined as follows?
	(a) An angular, rough, or broken surface of an aggregate particle.
	(b) Face considered fractured if <u>one-half or more of the projected area</u> is fractured.
	(c) Fractured face has sharp and well-defined edges (excluding small nicks).
2.	Fractured particle defined as a particle having at least the minimum number of fractured faces specified
•	(typically 1 or two fractured faces are required, depending on specification)?
?.	Sample dried and cooled on a clean flat surface large enough to permit careful inspection of each particle?
	Particle held so that the face is viewed directly (normal to the surface of the face)?
	Sample separated into three categories to aid in making the fracture determination?
•	(a) Group 1 - fractured particles meeting the above criteria.
	(b) Group 2 - particles not meeting specification criteria.
	(c) Group 3 - questionable or borderline particles.
	Mass of particles in each of the three categories determined?
'. '.	If on any of the determinations more than 15 percent of the total mass of the sample is placed in the
•	questionable category, is the determination repeated until no more than 15 percent is present in that category?
	questionable category, is the determination repeated until no more than 15 percent is present in that category?
	Mass percentage of the fracture faces calculated to the peacest I percent according to the following equation?
•	Mass percentage of the fracture faces calculated to the nearest 1 percent according to the following equation? $P = \left[\frac{(F+Q/2)}{(F+Q+N)} \right] \times 100$
STM	$P = [(F+Q/2)/(F+Q+N)] \times 100$ I Procedure
STM	$P = [(F+Q/2)/(F+Q+N)] \times 100$
STM	$P = [\ (F+\ Q/2)/(F+Q+N)\] \times 100$ $\frac{1\ Procedure}{Fractured\ face\ defined\ as\ follows?}$
STM	$P = [\ (F+Q/2)/(F+Q+N)\] \times 100$ I Procedure Fractured face defined as follows?
1. STM	$P = [(F + Q/2)/(F + Q + N)] \times 100$ I Procedure Fractured face defined as follows?
1. STM	$P = \left[\frac{(F+Q/2)}{(F+Q+N)} \right] \times 100$ $\frac{I \ Procedure}{Fractured face defined as follows?}$ $(a) An \ angular, \ rough, \ or \ broken \ surface \ of \ an \ aggregate \ particle.}$ $(b) Face \ considered \ fractured \ if \ \underline{one-fourth} \ or \ more \ of \ the \ \underline{maximum \ projected \ area}} \ is \ fractured.}$ $(c) Fractured \ face \ has \ sharp \ or \ slightly \ blunt \ edges \ (excluding \ small \ nicks).}$ $Fractured \ particle \ defined \ as \ a \ particle \ having \ at \ least \ the \ minimum \ number \ of \ fractured \ faces \ specified$
. <u>STM</u>	$P = [(F+Q/2)/(F+Q+N)] \times 100$ $I Procedure$ $Fractured face defined as follows?$
	$P = [(F+Q/2)/(F+Q+N)] \times 100$ $I \ Procedure$ $Fractured face defined as follows?$
. <i>STM</i>	$P = [(F+Q/2)/(F+Q+N)] \times 100$ $IProcedure$ $Fractured face defined as follows?$
	$P = [(F+Q/2)/(F+Q+N)] \times 100$ $IProcedure$ Fractured face defined as follows?
	$P = \left[\frac{(F+Q/2)}{(F+Q+N)} \right] \times 100$ $Exact Procedure$ $Fractured face defined as follows?$
	P = [$(F+Q/2)/(F+Q+N)$] × 100 I Procedure Fractured face defined as follows?
	P = $[(F+Q/2)/(F+Q+N)] \times 100$ Procedure Fractured face defined as follows?
	$P = [(F+Q/2)/(F+Q+N)] \times 100$ $IProcedure$ Fractured face defined as follows?
	P = $[(F+Q/2)/(F+Q+N)] \times 100$ I Procedure Fractured face defined as follows?
A.S.T.M.	P = $[(F+Q/2)/(F+Q+N)] \times 100$ Procedure Fractured face defined as follows?
3. 4.57.4. 5. 6. 7. 10.	P = $[(F+Q/2)/(F+Q+N)] \times 100$ I Procedure Fractured face defined as follows?

(T335 / D5821)

RESISTANCE TO DEGRADATION OF LARGE-SIZE COARSE AGGREGATE BY ABRASION AND IMPACT IN THE LOS ANGELES MACHINE

(C535)

m (28±0.2 in.), inside length 508±5 mn in.)?
in.)?
in.)?
in.)?
102 x 12.7 mm (6 x 4 x ½in.)?
counter reading (End): Elapsed time (seconds): RPM heres having a mass of 390-445 g:
counter reading (End): Elapsed time (seconds): RPM heres having a mass of 390-445 g:
of] full length of the cylinder?
of] full length of the cylinder?
counter reading (End): Elapsed time (seconds): RPM heres having a mass of 390-445 g:
ge suggested)?
ge suggested)?
Counter reading (End): Elapsed time (seconds): nds: RPM heres having a mass of 390-445 g:
Counter reading (End): Elapsed time (seconds): nds: RPM heres having a mass of 390-445 g:
Counter reading (End): Elapsed time (seconds): nds: RPM heres having a mass of 390-445 g:
heres having a mass of 390-445 g:
heres having a mass of 390-445 g:
heres having a mass of 390-445 g:

RESISTANCE TO DEGRADATION OF LARGE-SIZE COARSE AGGREGATE BY ABRASION AND IMPACT IN THE LOS ANGELES MACHINE

(C535)

	PROCEDURE	Date:
1.	Sample obtained by C702?	······
2.	Sample washed and oven-dried to constant mass at 110±5°C (230±9°F)?	
3.	Mass determined to nearest 1.0 g?	
4.	Specimen masses conform to the table below?	

SIEVE SIZE	GRADING 1	GRADING 2	GRADING 3
3 to 2 ½ in	$2500 \pm 50 \text{ g}$		
2 ½ to 2 in	$2500 \pm 50 \text{ g}$		
2 to 1 ½ in	$5000 \pm 50 \text{ g}$	$5000 \pm 50 \text{ g}$	
1 ½ to 1 in		$5000 \pm 25 \text{ g}$	$5000 \pm 25 \text{ g}$
1 to 3/4 in			$5000 \pm 25 \text{ g}$
Total Mass	$10,000 \pm 100 \text{ g}$	$10,000 \pm 75 \text{ g}$	$10,000 \pm 50 \text{ g}$

5.	Sample and spheres put in machine and tumbled 1000 times?	
	Note: Loss after 200 revolutions may be determined, and then entire sample returned to drum for final 800 revolutions.	
6.	Contents of drum separated on a sieve coarser than a 1.70 mm (No. 12)?	•
7.	Finer material separated on a No. 12 sieve?	•
8.	Material coarser than No. 12 washed and dried to constant mass at 110±5°C (230±9°F)? (See Note)	•
	Note: If material is essentially free of adherent coatings and dust, the requirement for washing is optional.	
	For referee testing, the washing procedure must be performed.	
9.	Mass of material coarser than No. 12 determined to nearest 1 g?	•
10.	Percentage of wear calculated as: % wear = (original - final mass) / original mass?	•
COMM	MENTS (C535):	(C535)

FLAT PARTICLES, ELONGATED PARTICLES, OR FLAT AND ELONGATED PARTICLES IN COARSE AGGREGATE

~	-04\
$(\mathbf{D4}')$	791)

			<u>APPAR</u>	<u>ATUS</u>	Date:	
1.	Note: Other dev	ur to Figures 2 or 3 or ices may also be accep	ptable if they can be ve	erified using a machin	ned block, micrometer, oed block, micrometer	
2.	Balance, accura	nte to 0.5% of sampl s: accurate to 5 g for	e mass?smallest sample size, a	a G20 / GP10.		
3.	Oven, maintain	as 110 ± 5 °C (230 \pm	9°F) [if determination	on by mass is requir	ed]?	<u></u>
Samp 1. 2.	Test sample ma	uss when dry conform	ns to following table s for D4791	approximately the e?		testing?
-	9.5 mm (3/8 in.) 1 kg (2 lb)	12.5 mm (1/2 in.) 2 kg (4 lb)	19.0 (3/4 in.) 5 kg (11 lb)	25.0 mm (1 in.) 10 kg (22 lb)	37.5 mm (1.5 in.) 15 kg (33 lb)	50 mm (2 in.) 20 kg (44 lb)
F	1 kg (2 lb)	2 Kg (4 IU)	3 kg (11 lb)	10 kg (22 lb)	13 kg (33 lb)	20 kg (44 lb)
	63 mm (2.5 in.)	75 mm (3 in.)	90 mm (3.5 in.)	100 mm (4 in.)	125 mm (5 in.)	150 mm (6 in.)
	35 kg (77 lb)	60 kg (130 lb)	100 kg (220 lb)	150 kg (330 lb)	300 kg (660 lb)	500 kg (1100 lb)
3.	Reduction to ex	xact predetermined r	mass not permitted?.	<i>//</i> [
COM	IMENTS (D4791):	/ /			$ \top $	(D4791)

FLAT PARTICLES, ELONGATED PARTICLES, OR FLAT AND ELONGATED PARTICLES IN COARSE AGGREGATE

(D4791)

	PROCEDURE (Continued) Date:
Procedu	re
1.	If determination by mass, sample oven-dried to constant mass at 110±5° C (230±9° F)?
3.	Using material retained on 9.5 mm (3/8 in.) or 4.75 mm (No. 4), as required, each size fraction present in amount of 10% or more of original sample reduced according to C702 until approximately 100 particles obtained for each size fraction required?
4.	Size fractions containing less than 10% by mass of the original total sample not tested (can be discarded)?
	Terminology Note: Length is defined as the biggest dimension of the particle. Thickness is the small dimension of the particle. Length $>$ Width $>$ Thickness or $L > W > T$
Method	${f A}$
1.	Each particle in each size fraction tested and placed in one of four groups:
	(1) Flat, (2) Elongated, (3) meeting the requirements of groups 1 and 2, and (4) neither Flat nor Elongated?
2. 3.	Proportional caliper device positioned at proper ratio?
 4. 	Flat particles determined by setting larger opening equal to particle width? ————————————————————————————————————
4.	Example: At a ratio of 2:1 an AASHTO test method book is flat. $W >> T$
5.	Elongated particles determined by setting larger opening equal to particle length?
6.	Particle is elongated if width can be placed within the smaller opening?
7.	Proportion of sample in each group determined by count or by mass, as required?
Method	B – for Superpave
1.	Each particle in each size fraction tested and placed into one of two groups:
_	(1) Flat & Elongated or (2) not Flat & Elongated?
2.	Proportional caliper device positioned at proper ratio?
3.	Larger opening set equal to particle length?
4.	Particle is <u>flat and elongated</u> if the <u>thickness</u> can be placed in the smaller opening?
5.	Proportion of sample in each group determined by count or by mass, as required?
Calculat	ion
1.	Percent particles in each category calculated to nearest 1% for each sieve size tested?
2.	Report shows original gradation of aggregate sample, number/mass of particles tested for each sieve size, percent particles in each category, and dimension ratio used?
COMMI	ENTS (D4791): (D4791)

SPECIFIC GRAVITY AND ABSORBTION OF FINE AGGREGATE USING INFRARED

(D7172)

	APPARATUS Date:
1.	Large neck volumetric flask, capacity 500-mL?
2.	Automatic Volumetric Mixer:
	(a) Orbital mixer capable of holding a 500 mL volumetric flask?
	(b) Clamp and clamping rod capable of securely holding the neck of the flask?
	(c) Vacuum pump capable of removing entrapped air?
	(d) Hose and stopper capable of joining the vacuum pump and mouth of the flask?
3.	Infrared Unit:
	(a) Capable of detecting saturated surface-dry (SSD) condition using an infrared sourced and detector?
	(b) Calibrated monthly? (Record S/N: and check records)
	(c) Consists of an orbital mixer, water pump, infrared source, infrared detector, and mixing bowl?
	(d) Lid for mixing bowl, consists of two sapphire lenses and an injection nozzle?
4.	Distilled Water?
5.	Thermometer:
	(a) Range of 0 to 50°C (0° to 122°F)?
	(b) Readable to 0.5°C (1°F)?
6.	Balance, readable to within 0.1% of the test sample mass at any point within the range of use?
7.	Timer, capable of measuring at least 5 minutes?
	CALIBRATION
	ration of Water Pump:
1.	Calibration performed monthly?
2.	Infrared device filled with distilled water?
3.	Mass of empty water-collection container determined?
4.	Water-collection container positioned to minimize splashing?
_	Note: Placing the container under the nozzle in the lid is a good way to minimize splashing.
5.	Manufacturers' instructions followed to determine the number of injections into the test flask
_	(usually 3000 total injections)?
6.	Mass of water-collection container determined?
7.	Mass of water injected determined by subtracting mass of empty container from mass of full container?
Calibi	ration of Infrared Unit:
1.	Calibration performed monthly?
2.	Infrared water unit full?
3.	Unit turned on and allowed to warm up per manufacturers' instructions?
4.	Initiate calibration routine per manufacturers' instructions?
5.	500.0 g (no tolerance) of Ottawa Silica sand inserted into mixing bowl?
6.	After calibration, unit displays results?
7.	Silica sand dried to constant mass at 110 ± 5 °C (230 ± 9 °F) and stored for future calibrations?
COM	MENTS (D7172): (D717

SPECIFIC GRAVITY AND ABSORBTION OF FINE AGGREGATE USING INFRARED

(D7172)

	PROCEDURE Date:
	g Procedure:
1.	Sample obtained by C702?
2.	Approximately 1.5 kg ± 10 g of fine aggregate obtained?
3.	Dried to constant mass at 110 ± 5 °C $(230 \pm 9$ °F)?
4.	Allowed to cool to $23 \pm 2.0^{\circ}\text{C}$ (73 ± 3°F)?
5.	Sample split into two 500 ± 5 g samples?
6.	Excess sample discarded?
Film Co	efficient Determination:
1.	Approximately 250-mL of water at 23 ± 2.0°C (73 ± 3°F) placed in calibrated pycnometer?
2.	Pycnometer placed on balance and zeroed?
3.	Approximately 500.0 ± 0.1 g sample transferred to pycnometer with water and mass determined?
4.	Water in pycnometer completely covers the sample but does not overflow or exceed calibration line?
5.	Sample allowed to stand for 5 minutes?
	Note: Paper towel or isopropyl alcohol may be used to remove air bubbles if necessary.
6.	Pycnometer filled with 23 ± 2.0 °C (73 ± 3 °F) water to the calibration mark?
7.	Mass determined to the nearest 0.1 g?
8.	Pycnometer with rubber stopper inserted in AVM unit?
9.	Mixer agitates pycnometer for three minutes, then the vacuum pump engages at a level of 56 cm (22 in.) Hg
	for another 3 minutes, and last 5 minutes engages at a level of 69 cm (27 in.) Hg (automatic setting)?
10.	AVM unit stops automatically when testing is complete (approximately 11 minutes)?
11.	Isopropyl alcohol or paper towel used to remove air bubbles?
12.	Pycnometer filled to calibration mark with water and mass determined?
13.	Proper book formulas used to calculate film coefficient?
	/
	Gravity and Percent Absorption Determination:
1.	Infrared unit allowed to warm up for 30 minutes?
2.	Sample weighing 500 ± 0.1 g placed into the bowl and the mass determined?
3.	Bowl with aggregate placed in infrared unit with the notch in front aligned with metal mounting plate?
4.	Ring on the bowl fastened by pressing down and turning the ring one quarter turn until tight?
5.	Lid closed and latched with notch lined up in front of the bowl?
6.	Reservoir unit full of distilled water?
7.	Film coefficient entered?
8.	After the test, film coefficient on display compared with the measured film coefficient for the material?
9.	Mass of the bowl immediately determined after removing the lid?
10.	Percent absorption determined?
11.	Lab says proper book formulas used in calculations?
COMM	ENTS (D7172): (D7172)

COMMENTS (D7370):

RELATIVE DENSITY AND ABSORPTION OF AGGREGATE USING VACUUM SATURATION AND RAPID SUBMERSION

(D7370)_	
----------	--

Date: _____

APPARATUS

Balance	readable and accurate to 0.1 g, equipped with a suitable apparatus for suspending the sample in water?
Water B	
1.	Equipped with an overflow outlet?
2.	Sample completely submerged when suspended?
3.	Maintains temperature of 25 ± 1 °C (77 ± 2 °F)?
4.	<u>Square:</u> Length of 610 mm by Width of 460 mm (18 in.) by Depth of 460 mm (24 x 18 x 18 in square)?
or	Cylindrical: Minimum diameter and minimum depth of 460 mm (18 in.)?
Sample :	
1.	Having no sharp edges, for displacement of the sample?
Vacuum	<u>Chamber:</u>
1.	Equipped with a pump capable of evacuating chamber to 6 mm Hg (at sea level)?
2.	Automatically seals bag?
3.	Exhausts air back into chamber in a controlled manner to ensure plastic conforms to specimen?
4.	Air exhaust and vacuum operation time set at factory prior to initial use?
Vacuum	Measurement Gauge:
1.	Independent of the vacuum sealing device?
2.	Capable of being placed directly inside chamber to verify vacuum performance and sealing of unit?
3.	Capable of reading pressure down to 3 mm Hg?
4.	Readable to $\pm 1 \text{ mm Hg}$?
Plastic E	
1.	Made of puncture-resistance plastic, impermeable to water, minimum thickness of 0.127 mm (0.005 in.)?
2.	Apparent specific gravity of bags provided by the manufacturer?
3.	One of the following sizes used:
	(a) <u>Smaller bags:</u> Opening in bag 235 – 260 mm (9.25 – 10.25 in.)?
	(b) <u>Larger bags:</u> Opening in bag 375 – 394 mm (14.75 – 15.5 in.)?
	letal Pycnometer (for testing fine aggregate):
1.	Inner diameter of $137 \pm 0.2 \text{ mm} (5.375 \pm 0.008 \text{ in.})?$
2.	Height of 89 ± 0.40 mm $(3.5 \pm 0.016$ in.)?
3.	Machined smooth on all surfaces?
4.	Inside of lid machined at 5° angle to create an inverted conical surface?
5.	Equipped with a graduated temperature strip to monitor temperature during testing?
6. -	Lid has a 3 mm (1/8 in.) hole on its surface?
/.	Equipped with a fixture for holding and securing lid in place and equipped with a leveling indicator?
_	letal Pycnometer (for testing coarse and blended aggregate):
1.	Inner diameter of $198 \pm 0.2 \text{ mm}$ (7.776 ± 0.008 in.)?
2.	Height of $114 \pm 0.8 \text{ mm} (4.5 \pm 0.03 \text{ in.})$?
3.	Machined smooth on all surfaces?
4. -	Inside of lid machined at 5° angle to create an inverted conical surface?
5.	Equipped with a graduated temperature strip to monitor temperature during testing?
5.	Lid as a 3 mm (1/8 in.) hole on its surface?

(D7370)

RELATIVE DENSITY AND ABSORPTION OF AGGREGATE USING VACUUM SATURATION AND RAPID SUBMERSION

(D1310)	(D7370)	
---------	---------	--

	APPARATUS (Continued) Date:
Access	
1.	Knife or scissors for cutting plastic bags?
2.	Spray bottle filled with isopropyl alcohol?
3.	A bucket large enough to allow the pycnometer to be fully submerged in water?
4.	Water containers to dispense water during testing?
5.	Syringe with a needle no larger than 3 mm (0.125 in.)?
6.	Small paint brush?
7.	Metal spatula 25 ± 5 mm $(1 \pm 0.2 in.)$ wide?
	r Sheets:
1.	For protecting plastic bags against sharp edges of the aggregate sample?
2.	Apparent specific gravity provided by the manufacturer?
3.	Thermometric device, for monitoring temperature to within \pm 1°C (\pm 1.8°F)?
	<u>VERIFICATION</u>
Systen	n Standardization:
1.	Vacuum settings of chamber verified every 12 months, after major repairs, and after shipment / relocation?
2.	Vacuum measurement gauge placed inside vacuum chamber?
3.	Setting recorded while vacuum is operating?
4.	Gauge indicates a pressure of 6 mm Hg (6 TORR) or less?
5.	Unit not used if gauge reads above 6 mm?
6.	Vacuum Measurement Gauge, standardized for accuracy once a year?
	Note: In-line vacuum gauges are not suitable for use in enclosed chambers and shall not be used.
Calibra	ation of Small and Large Pycnometer:
1.	Pycnometer re-calibrated before each day of use?
2.	Pycnometer conditioned to $25 \pm 1^{\circ}$ C ($77 \pm 2^{\circ}$ F) by placing inside a bucket of water maintained at that temp.?.
3.	Small Pycnometer: Lid-holding fixture leveled during conditioning (using attached or separate level ok)?
4.	Pycnometer removed from water bucket and dried with a towel?
5.	Small Pycnometer: Pycnometer placed in the fixture and pushed back until contact is made with the stops?
6.	Large Pycnometer: Pycnometer set on a level surface?
7.	Pycnometer filled with water at $25 \pm 1^{\circ}$ C (77 $\pm 2^{\circ}$ F) to approximately 10 mm (0.375 in.) from the top?
8.	Surface of the water sprayed with isopropyl alcohol spray bottle to remove bubbles?
9.	Lid gently placed on pycnometer (Small Pycnometer: and fixture clamps closed)?
10.	Syringe filled with water at 25 ± 1 °C (77 ± 2 °F)?
11.	Pycnometer filled through the large fill hole on the lid post with the syringe?
12.	Syringe tip kept below the water level while filling?
13.	Formation of air bubbles avoided?
14.	Pycnometer filled until water comes out of the 3 mm hole on the lid?
15.	Any remaining water on the top of the lid wiped with a towel?
16.	Pycnometer (Small Pycnometer: Entire Fixture) placed on the scale and the mass recorded to nearest 0.1 g?
17.	Pycnometer cleaned and calibration repeated two more times?
18.	Average of three calibration masses obtained?
19.	Small Pycnometer: Range of three calibration masses is within 0.5 g?
20.	Large Pycnometer: Range of three calibration masses is within 1 g?
21.	If the range of calibration masses is not within 0.5 g (small pycnometer) or 1 g (large pycnometer), steps
	taken to ensure that the calibration is done correctly?
22.	Calibration repeated until 0.5 g (small pycnometer) or 1 g (large pycnometer) requirement is met?
COMN	MENTS (D7370): (D7370)

COMMENTS (D7370):

RELATIVE DENSITY AND ABSORPTION OF AGGREGATE USING VACUUM SATURATION AND RAPID SUBMERSION

(D7370)_	
----------	--

	SAMPLE PREPARATION Date:
	regate Samples (Method A):
1.	Sample thoroughly mixed before reducing?
2.	Reduced to one 1000 ± 5 g (for apparent density) and two 500 ± 3 g samples (for bulk density)?
3.	Reduction done in accordance with ASTM C702?
Coarse	ggregate Samples (Method B):
1.	Aggregate dried to constant mass at $110 \pm 5^{\circ}$ C ($230 \pm 9^{\circ}$ F)?
2.	Thoroughly mixed before reducing?
3.	Reduced to one 2000 \pm 10 g (for apparent density) and two 1000 \pm 10 g samples (for bulk density)?
4.	Reduction done in accordance with ASTM C702?
5.	If the sample is tested in two or more size fractions, sample is graded in accordance with ASTM C136?
6.	Grading must include the sieves used for separating the size fractions?
	Method A (Fine Aggregate) - PROCEDURE
	A, Fine Aggregate - Bulk Density Determination:
1.	Water temperature maintained at 25 ± 1 °C (77 ± 2 °F) throughout bulk and apparent density determination?
2.	Pycnometer conditioned to $25 \pm 1^{\circ}$ C (77 ± 2°F) by placing inside a bucket of water maintained at that temp.?.
3.	Samples dried to constant mass and allowed to cool to room temperature?
4.	Pycnometer removed from water bucket and dried with a towel?
5.	Pycnometer placed in the fixture and pushed back until contact is made with the stops?
6. 7.	A 500 \pm 3 g sample at 25 \pm 1°C (77 \pm 2°F) weighted and mass recorded?
7. 8.	Approximately 500 mL of water (halfway full) placed in pycnometer?
8. 9.	Care taken to ensure that aggregate is not lost in the filling process?
9. 10.	Brush used to sweep away any remaining fines into the pycnometer, if necessary?
11.	If any aggregate is lost during the filling process, is the test started over?
12.	Metal spatula pushed to the bottom of the pycnometer against the circumference?
13.	Spatula slowly and gently dragged to the center of the pycnometer and removed after reaching center?
14.	Steps 10 & 11 repeated 7 more times (8 times total) around the sample in 45° increments
	until the start point is reached?
15.	Squeeze water bottle used to rinse sample residue off of the spatula and into the sample, if necessary?
16.	Pycnometer filled with water to approximately 10 mm (0.375 in.) from the rim of the container?
17.	Surface of the water sprayed with isopropyl alcohol spray bottle to remove bubbles?
18.	Lid gently placed on pycnometer and fixture clamps closed?
19.	Syringe filled with water at 25 ± 1 °C (77 ± 2°F)?
20.	Pycnometer filled through the large fill hole on the lid post with the syringe?
21.	Syringe tip kept below the water level while filling?
22.	Formation of air bubbles avoided?
23.	Pycnometer filled until water comes out of the 3 mm hole on the lid?
24.	Any remaining water on the top of the lid wiped with a towel?
25.	Entire fixture, including pycnometer, placed on the scale and the mass recorded to nearest 0.1 g?
26.	Procedure repeated with a second 500 ± 3 g sample?
27.	Average masses calculated for all determinations made for the duplicate samples?

Revised 2014-04-10

(D7370)

RELATIVE DENSITY AND ABSORPTION OF AGGREGATE USING VACUUM SATURATION AND RAPID SUBMERSION

(D7370) _	
-----------	--

Method A (Fine Aggregate) – PROCEDURE (continued)

Date:

Method	A, Fine Aggregate - Apparent Density Determination:
1.	Pressure level set on vacuum device according to manufacturer's instructions?
2.	Small plastic bag inspected for holes, stress points, or side seal discontinuities (discarded if bag is damaged)?
3.	Plastic bag mass recorded?
4.	The 1000 ± 5 g sample of oven dried aggregate mass recorded and sample placed in bag?
5.	Bottom of the bag supported by a smooth tabletop while pouring to prevent puncture and impact?
6.	The bag containing sample placed in vacuum chamber?
7.	Sample spread flat by grabbing the bag from two sides and shaking gently?
8.	Pressing down on the sample from outside of the bag avoided?
9.	If the aggregate sample contains a large amount of minus 75-µm (No. 200) material, sample lightly
	misted to keep dust down during sealing?
10.	Open end of the bag placed over the seal bar in the chamber?
11.	Chamber door closed and vacuuming and sealing process begins?
12.	After sealing, chamber door opened and sample removed?
13.	Sample immediately submerged into the water tank?
	Note: It is extremely important the sample be submerged immediately after vacuum sealing to prevent air from
4.4	slowly entering the bag. This can result in low apparent density results.
14.	One corner of the plastic bag, approximately 25 to 50 mm (1 to 2 in.), cut from one side?
15.	Bag is completely submerged at least 2 in. below water surface while cutting the bag?
16.	Cut portion of bag held open for 45 seconds to allow water to freely enter?
17.	Any small residual air bubbles allowed to escape?
18.	Shaking and squeezing the sample avoided (may cause fines to escape)?
19.	Second corner of the bag cut and any residual air bubbles removed by running fingers across the top the bag?
20.	Bag placed in weighing basket in the water?
21.	If bag is folded to place it in the basket, is the bag unfolded to allow water to freely flow into the sample once it is in the basket?
22.	Sample and bag kept underwater at all times?
23.	Care is taken to ensure that the bag and sample are not touching the bottom or sides of the tank, or floating
23.	out of the water tank?
24.	Sample allowed to stay in the water bath for a minimum of 15 minutes?
25.	Underwater mass of sample and bag recorded to nearest 0.1 g?
26.	Data entered into PC using manufacturer's software or equations given in method?
COMMI	ENTS (D7370): (D7370)

RELATIVE DENSITY AND ABSORPTION OF AGGREGATE USING VACUUM SATURATION AND RAPID SUBMERSION

(D7370)	
---------	--

Method B (Coarse or Combined Aggregate) - PROCEDURE Date:

Method	B, Coarse and Combined Aggregate Samples - Bulk Density Determination
1.	Water temperature maintained at $25 \pm 1^{\circ}$ C ($77 \pm 2^{\circ}$ F) throughout bulk and apparent density determination?
2.	Pycnometer conditioned to $25 \pm 1^{\circ}$ C (77 $\pm 2^{\circ}$ F) by placing inside a bucket of water maintained at that temp.?.
3.	Samples dried to constant mass and allowed to cool to room temperature?
4.	Pycnometer removed from water bucket and dried with a towel?
5.	A 1000 ± 10 g sample at 25 ± 1 °C (77 ± 2 °F) weighted and mass recorded?
6.	Approximately 1000 mL of water (halfway full) placed in pycnometer?
7.	Sample slowly and evenly poured into the pycnometer?
8.	Care taken to ensure that aggregate is not lost in the filling process?
9.	Brush used to sweep away any remaining fines into the pycnometer, if necessary?
10.	If any aggregate is lost during the filling process, is the test started over?
11.	Metal spatula pushed to the bottom of the pycnometer against the circumference?
12.	Spatula slowly and gently dragged to the center of the pycnometer and removed after reaching center?
13.	Steps 10 to 11 repeated 7 more times (8 times total) around the sample in 45° increments until
	the start point is reached?
14.	Squeeze water bottle used to rinse sample residue off of the spatula and into the sample, if necessary?
15.	Pycnometer filled with water to approximately 10 mm (0.375 in.) from the rim of the container?
16.	Surface of the water sprayed with isopropyl alcohol spray bottle to remove bubbles?
17.	Lid gently placed on pycnometer?
18.	Syringe filled with water at 25 ± 1 °C (77 ± 2 °F)?
19.	Pycnometer filled through the large fill hole on the lid post with the syringe?
20.	Syringe tip kept below the water level while filling?
21.	Formation of air bubbles avoided?
22.	Pycnometer filled until water comes out of the 3 mm hole on the lid?
23.	Any remaining water on the top of the lid wiped with a towel?
24.	Pycnometer placed on the scale and the mass recorded to nearest 0.1 g?
25.	Procedure repeated with a second 1000 ± 10 g sample?
26.	Average masses calculated for all determinations made for the duplicate samples?
Method	B, Coarse and Combined Aggregate Samples - Apparent Density Determination
1.	Pressure level set on vacuum device according to manufacturer's instructions?
2.	One small plastic bag and one large plastic bag selected, inspected for holes, stress points, or side seal discontinuities,
	and any defective bags discarded?
3.	Both plastic bags weighed and the mass recorded?
4.	Two rubber sheets weighed and the mass recorded?
5.	Mass of the 2000 \pm 10 g sample of oven dried aggregate recorded and sample placed in small bag?
6.	Bottom of the bag supported by a smooth tabletop while pouring to prevent puncture and impact?
7.	Large bag placed in the vacuum chamber?
8.	Rubber sheet placed inside large plastic bag?
9.	Rubber sheet is flat, centered, and pushed all the way to the back of the bag?
10.	Small plastic bag containing sample placed into large plastic bag on top of rubber sheet?
11.	Sample manually spread inside small plastic bag?
COMM	ENTS (D7370): (D7370)

RELATIVE DENSITY AND ABSORPTION OF AGGREGATE USING VACUUM SATURATION AND RAPID SUBMERSION

(D7370)	

PROCEDURE (Continued)

	PROCEDURE (Continued) Date:	
Method	B, Coarse and Combined Aggregate Samples - Apparent Density Determination (Continued)	
12.	If the aggregate sample contains a large amount of minus 75-µm (No. 200) material, sample	
	lightly misted to keep dust down during sealing?	
13.	Another rubber sheet placed on top of small bag, inside large bag?	
14.	Is the small bag completely contained within the area between the two rubber sheets?	_
15.	Open end of the large bag placed over the seal bar in the chamber?	
16.	Rubber sheets are not over the seal bar?	_
17.	Chamber door closed and vacuuming and sealing process begins?	
18.	Chamber door opened and sample removed?	
19.	Sample immediately submerged into the water tank?	_
	Note: It is extremely important the sample be submerged immediately after vacuum sealing to prevent air from slowly	_
	entering the bag. This can result in low apparent density results.	
20.	One corner of the large plastic bag, approximately 70 to 100 mm (3 to 4 in.), cut from one side?	
21.	Bags are completely submerged below water surface while cutting the bag?	
22.	Cut portion of large bag held open for 25 seconds to allow water to freely enter?	
23.	Any small residual air bubbles allowed to escape?	
24.	Second corner of the large bag cut?	
25.	Any residual air bubbles removed by running fingers across the top the large bag?	
26.	Bags placed in weighing basket in the water?	
27.	If bag is folded to place it in the basket, is the bag unfolded to allow water to freely flow into the sample once it is in the basket?	
28.	Sample and bag kept underwater at all times?	
29.	Care is taken to ensure that the bag and sample are not touching the bottom or sides of the tank, or	
	floating out of the water tank?	
30.	Sample allowed to stay in the water bath for a minimum of 20 minutes?	
31.	Underwater mass of sample and bags recorded to nearest 0.1 g?	
32.	Data entered into PC using manufacturer's software or equations given in method?	
COMM	ENTS (D7370): (D7370	0)

COMMENTS (D7428):

RESISTANCE OF COARSE AGGREGATE TO DEGRADATION BY ABRASION (D7428) _____ IN THE MICRO-DEVAL APPARATUS (FINE AGGREGATE)

		<u>APPARATUS</u>		Date:	
1.	Micro-Deval Abrasion Machine, a j	ar rolling mill similar to Fig	1, capable of running	at 100 ± 5 rpm?	
2.	(b) External diameter of 194 to	y, with a rubber ring in the rope 202 mm and internal heights smooth and have no observ	t of 170 to 177 mm?.		
3.		250 ±5 g presented for use in 0.5 mm in diameter? (Meas			
	9.5 ± 0.5 mm 1 2 Diameter ok?	3 4 5	6 7	8 9 10	
 4. 5. 	Note: It is recommended that separate Sieves, 6.3-mm (1/4 in.), 4.75-mm (300- μ m (No. 50), 150- μ m (No. 100 Note: A 6.7-mm sieve may be used in p Oven, maintains $110 \pm 5^{\circ}$ C?	(No. 4), 2.36-mm (No. 8), 1.), 75-μm (No. 200)? place of a 6.3-mm sieve.	18-mm (No. 16), 600-	-µm (No. 30),	
6.	Balance, accurate to 0.1 g?				
<u>Calibra</u> 1.	tion Supplies Standard Sutherland Micro-Deval F			le)?	···
	Passing	Retained	Mass	OK?	
	4.75-mm (No. 4)	2.36-mm (No. 8)	40g		
	2.36-mm (No. 8)	1.18-mm (No. 16)	115g		
	1.18-mm (No. 16)	600-μm (No. 30)	180g		
	600-μm (No. 30)	300-μm (No. 50)	120g		
	300-μm (No. 50)	150-μm (No. 100)	38g		
2	150-μm (No. 100)	75-µm (No. 200)	7g		
2.	Calibration Aggregate, mean loss be	etween 15 to 25 %?			
Calibra	tion Procedure				
1.	10 samples of calibration aggregate				
2. 3.	10 samples of Standard Sutherland If Standard Sutherland Micro-Deva				
٥.	loss 95% of the time, the mean valu				
4.	Calibration procedure conducted for	r new supplies of calibration	aggregate, batched ac	ecording to Section 8?	
5. 6.	Control sample tested every 10 sam Percent loss of last 20 samples of ca				
0.	1 electic loss of last 20 samples of ea	moration aggregate protted (m ucha chart:	••••••	

(D7428)

Sample prepared as follows:

Sample Preparation

2.

3.

RESISTANCE OF COARSE AGGREGATE TO DEGRADATION BY ABRASION (D7428) IN THE MICRO-DEVAL APPARATUS (FINE AGGREGATE)

	PROCEDURE	Date:
<u>Preparation</u>		
Test Sample washed over 75-µm sieve and ov	ven-dried to constant mass at 110±5°C?	
Sample separated into individual size fraction	s in accordance with ASTM: C136?	

Passing	Retained	Mass	OK?
4.75-mm (No. 4)	2.36-mm (No. 8)	50g	
2.36-mm (No. 8)	1.18-mm (No. 16)	125g	
1.18-mm (No. 16)	600-μm (No. 30)	125g	
600-μm (No. 30)	300-μm (No. 50)	100g	
300-μm (No. 50)	150-μm (No. 100)	75g	
150-μm (No. 100)	75-µm (No. 200)	25g	
Tot	al	$500 \pm 5 \text{ g}$	

Note: It may be practical to test material without preparing for the above grading for routine quality control purposes.

Procedure

11000	uui C
1.	Prepared sample weighed to nearest 0.1 g?
2.	Sample immersed in 0.75 ± 0.05 L of tap water either in Micro-Deval container or other suitable device?
	(a) Temperature of tap water 20 ± 5 °C?
	(b) Immersed for a minimum of 1 h?
3.	Sample placed in Micro-Deval abrasion container?
	(a) With 1250 ± 5 g of steel balls?
	(b) Also with the same water used to saturate the sample?
4.	Cover installed and Micro-Deval container placed on the machine?
5.	If machine is capable of recording total number of revolutions:
	(a) Machine run at 100 ± 5 rpm for 1500 ± 10 revolutions?
or	If machine is not capable of recording total number or revolutions:
	(a) Machine run at 100 ± 5 rpm for $15 \text{ min } \pm 5 \text{ seconds}$?
6.	Sample and steel balls carefully poured over a 6.3-mm (1/4 in.) sieve into a suitable container?
	Note: A 6.7-mm sieve may be used instead of a 6.3-mm sieve.
	(a) Care taken to remove entire sample from the stainless steel jar?
	(b) No aggregate is lost in the process?
	(c) Steel balls retained on the sieve washed to remove adhering aggregate?
7.	Material recovered in the container below the 6.3-mm (1/4 in.) sieve washed in accordance with ASTM C117?
	(a) Washing continued until water runs clear and all material smaller than 75 μm passes through sieve?
8.	Sample dried to constant mass at 110 ± 5 °C?
9.	Sample weighed to the nearest 0.1 g?
10.	Micro-Deval abrasion loss calculated to the nearest 0.1% as follows:

Percent Loss = (A - B) / A * 100

where:

A = Initial sample mass B = Final sample mass

COMMENTS (D7428): (D7428)