HOT-MIX WORKSHEET INDEX

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^{**} NP for Not Presented or use a vertical line.

^{★ -} Indicates the line has been modified since the last version of the worksheets, 2010-09-21.

REDUCING SAMPLES OF HOT-MIX ASPHALT TO TESTING SIZE

<u>APPARATUS</u>

-	_		-	
- (ĸ	4	. / \	

		APPARATUS Date:	
Equipn	nent for	one of the following methods:	
Machai	nical Sn	litter Method	
1.		anical Splitter Type A:	
1.	(a)	Designed so that the HMA field sample will flow smoothly and freely through the divider	
	(u)	without restriction or loss of materials (See Figure 1)?	
	(b)	Splitter has four equal width chutes?	
	(c)	Four appropriate sized containers?	
	(d)	Hopper with release handle?	
	()		
2.	Mech	anical Splitter Type B:	
	(a)	No less than 8 equal sized openings?	
	(b)	The openings minimum width must be at least 50% larger than largest particle to be split?	
	(c)	Hopper or straightedge pan?	
3.	Annro	oved Release Agent (such as non-stick cooking spray)?	
J.	(a)	Does not contain solvents?	
	(b)	Does not contain petroleum based products that affect binder properties?	
	(0)	Note to Assessors: Products such as WD-40 contain solvents and petroleum products, and are not acceptable for this test method.	
<u>Quarter</u> 1.	ring Mer One o	thod: of the following:	
	(a)	Quartering template?	
		(1) Forms a cross forming 90 degree angles at juncture?	
		(2) Sufficient length (1.1 times the diameter of the flattened cone of HMA to be quartered)?	
or	(b)	Straightedges?	
_	\	AASHTO Materials Reference Laboratory	
2.		ottom scoop?	
3.		ge spatula, trowel, or piece of metal to be used as a straightedge?	
4. ~		stick paper or heat resistant plastic?	
5.		oved Release Agent (such as non-stick cooking spray)?	
	(a)	Does not contain solvents?	
	(b)	Does not contain petroleum based products that affect binder properties?	
		acceptable for this test method.	
	ental M		
1.	Flat b	ottom scoop?	
2.	Non-s	stick heavy paper or heat-resistant plastic?	
3.	Large	spatulas, trowels, metal straightedges, or a 12-in drywall taping knife?	
4.	Hot p	late, gloves, buckets, and cans?	
COMM	IENTS	(R47):	(R47)

COMMENTS (R47):

REDUCING SAMPLES OF HOT-MIX ASPHALT TO TESTING SIZE

	<u>PROCEDURE</u>	Date:
	chanical Splitter Method: for a large amount of material, Method A should be	
1.	Optional: Splitter and accessories heated, not to exceed 110°C as d	
_	temperature device?	
2.	Optional: All surfaces coming into contact with HMA coated with	approved release agent?
3.	Mechanical Splitter Method (Type A)	
	(a) Field or laboratory sample placed in hopper avoiding samp	
	(b) Sample containers positioned to receive HMA?	······
	(c) Release handle used dropping HMA through chutes?	
	(d) Samples taken from opposing corners for reintroduction in	
	(e) Split as many times as necessary for appropriate test?	······
4.	Mechanical Splitter Method (Type B)	
	(a) Sample placed in hopper or straightedge pan?	
	(b) Uniformly spread edge to edge?	
	(c) Rate at which sample introduced allows free flow into same	
	(d) Steps repeated until sample size obtained?	
Note:	e: Unlike C702, the half of the split sample normally regarded as trash may be s	et aside for reduction in size for other tests.
	artering Method	
1.	Sample placed on a hard, non-stick, clean, level surface?	
2.	Approved release agent, non-stick paper, or heat resistant plastic ma	
3.	Sample mixed to uniformity by turning over four times?	
4.	Mixed using flat bottom scoop or by alternately lifting each corner of	
	pulling toward the opposite corner?	
5.	During the last turning, entire sample formed into conical pile by de	
	top of previous one or by lifting two opposite corners of the paper o	
6.	Pile flattening into uniform thickness and diameter by pressing dow	
7.	Diameter approximately four to eight times the thickness?	
8.	A visual check is done to ensure that the material is homogenous?	
9.	Flattened mass divided into four quarters using quartering template Quartering template pressed down until it has complete contact with	or straightedges?
10.	Quartering template pressed down until it has complete contact with	surface?
11.	Two diagonally opposite quarters selected as "quartered" material?	
12.	Steps repeated until sample size obtained?	
	remental Method	
1.	Sample placed on a hard, non-stick, clean level surface covered with	
	plastic, or another suitable material?	
2.	Sample mixed to uniformity by turning over four times?	
3.	Mixed using flat bottom scoop or by alternately lifting each corner of	
	pulling toward the opposite corner?	
4.	During the last turning, entire sample formed into conical pile by de	
	top of previous one or by lifting two opposite corners of the paper o	r plastic?
5.	A visual check is done to ensure that the material is homogenous?	
6.	Paper or plastic grasped and material is rolled into a cylindrical roll	
7.	Paper pulled so that at least 1/4 of the length of the loaf is off of the e	
	overhanging the counter sliced off and placed in a container?	
0	or A straightedge used to slice off approximately 1/4 of the loaf and mat	erial placed in a container?
8	Additional material removed as needed to obtain test size?	

(R47)

MECHANICAL ANALYSIS OF EXTRACTED AGGREGATE

(130)	
(D5444)	

	APPARATUS Date:	
1.	Nest of sieves: Upper sieve: 2.00- or 1.18-mm (No. 10 or 16), lower sieve No. 200?	
2.	Mechanical Shaker, sieving accuracy met in a reasonable time period?	
3.	Wetting agent used?	
4.	Oven, capable of maintaining $110 \pm 5^{\circ}$ C $(230 \pm 9^{\circ}F)$?	
5.	Balance, capable of weighing to 0.1 % of sample mass?	
6.	Container, capable of holding sample covered with water and to permit agitation without without loss of sample?	
	<u>PROCEDURE</u>	
Sample	Preparation: circle one Extraction sample Ignition sample	
1. 2. 3. 4. 5. 6. 7.	Sample consists of all aggregate after extraction or ignition oven sample? ASTM only: Gradation analysis only performed on aggregate extracted by ignition method in Test Method D6307 when the correction factor is 1.0 or less? Minimum mass of mix sample based on nominal maximum size? Sample dried to constant mass? Sample weighed to nearest 0.1g (enter mass below)? Extraction samples only: Total mass of aggregate for percent calculation includes mineral matter mass? AASHTO only: If from T308, sample agrees with the mass after ignition from T308 (W _F) to within 0.1 % or else not used for acceptance purposes?	
Wash 1. 2. 3. 4. 5. 6. 7. 8.	After mass is recorded, sample placed in container and covered with water? Small amount of wetting agent added? Contents of container agitated vigorously? Wash water poured through proper nest of two sieves? Decantation of coarse particles onto the sieves avoided as much as possible? Washing continued until wash water is clear? Material retained on nested sieves returned to container? Washed material coarser than 75-µm (No. 200) dried according to T255 [ASTM only: Material can be dried to constant mass at a max temp. = compaction temperature + 9°F (5°C)]? Mass Before Washing Mass After Washing Sample weighed to nearest 0.1g?. Amount of - No. 200 material removed by washing calculated? (T30 / D5444):	

MECHANICAL ANALYSIS OF EXTRACTED AGGREGATE

(130	")
(D5444	Ð

		<u>PROCEDURE</u>	(Continued)	Date:			
Sieve T	esting: circle which type(s) we	re used 8-in sieves	12-in sieves	Other (such as square sieves)			
 Material sieved on specified sieves (including 75-μm)? Sieving continued until not more than 0.5 percent by mass of the total sample passes a given sieve in 1 min. (check by hand with 8 in. diameter sieve)? 							
	Sieve size: Total sample: Mass passing sieve: Percent Passing: (a) Mass retained on any sieve with openings smaller than No. 4 is less than 6 kg/m² (4 g/in.²) (200 g for 8 in. diameter sieve, 438 g for 12 in. diameter sieve)? (b) Mass retained on any sieve with openings larger than No. 4 is less than 2.5 x (sieve opening in mm) x (πr²) [see table below]? Note to assessors: This is not identical to (T27/C136), they are calculated differently.						
	Sieve	Opening (mm)	Mass (g) – 8 in. dia	Mass (g) – 12 in. dia.			
	< #4	< 4.75	200	438			
	#4	4.75	385	867			
	1/4 in.	6.3	510	1149			
	3/8 in.	9.5	770	1734			
	1/2 in.	12.5	1013	2281			
	3/4 in.	19.0	1539	3468			
3. 4.	Each fraction of aggregate weighed, including minus 75-µm (No. 200)?						
5. <i>or</i>	sieving + minus 75-µm by washing?						
6. 7. 8.	Sizes larger than 75-µm (No. 200) reported to nearest 1.0 percent (at least)?						
COMM	IENTS (T30 / D5444):			(T30 / D3	5444)		

MOISTURE OR VOLATILE DISTILLATES IN PAVING MIXTURES

(1110)	
(D1461)	

		<u>APPARATUS</u>	Date:
	Still:		
	(a)	Height: 152.4 ± 6.4 mm $(6.0 \pm 0.25$ in.) by 94.0 ± 5.1 mm (3.7 ± 0.2)	in.) O.D.?
		Note: Stills with a 5 in. O.D. may be used to accommodate larger so	amples.
	(b)	Still head has one hole 25.4 mm (1 in.) in inside diameter?	
	(c)	Clamp for still head satisfactory?	
	(d)	Heavy paper gasket?	
•	Glass	Tube Condenser:	
	(a)	Jacket length not less than 400 mm (15 3/4 in)?	<u> </u>
	(b)	Inner tube O.D. 9.5 to 12.7 mm (3/8 to 1/2 in)?	
		Note: It may be necessary to supplement one condenser with a secon	nd of the same dimensions.
	Receiv	ver, made of well-annealed glass? [ASTM: conforming to Section 4.3?]	
	Solven	nts:	
	(a)	Xylene? or 20% toluene and 80% xylene?	
	(b)	Petroleum distillate?	<u> </u>
	(c)	Sodium carbonate (Na ₂ CO ₃) (For Volatile Distillates)?	·····
	Heatin	g device	
•	Houtin	<u>g de 100</u>	

COMMENTS (T110 / D1461): (T10 /

(T110 / D1461)

MOISTURE OR VOLATILE DISTILLATES IN PAVING MIXTURES

(T110)	_
(D1461)	

	<u>PROCEDURE</u>	Date:
Comple	Preparation:	
1.	Sample thoroughly mixed?	
2.	Weighed sample should be not less than 500 g for normal mixtures?	
3.	Weighed sample broken up to avoid large lumps and placed in the still?	
4.	Remainder of the sample kept in a tightly covered container?	
••	remainder of the sample rept in a against covered container.	
	ination of Moisture	
1.	Sample placed in the still?	
2.	200 mL of solvent added to the sample and stirred?	
3.	Gasket moistened with water?	
4.	Satisfactory assembly of components (all connections vapor or liquid tight, caution is	
5.	Loose cotton plug inserted in top of condenser?	
6.	Cold water circulated in jacket of condenser?	······································
7.	Heat applied and refluxing starts within 5 to 10 minutes?	······································
8.	Drip rate adjusted from 85 to 95 drops of distillate per minute?	······································
9.	Distillation time does not exceed 1.5 hours?	······
or	Distillation continued until three successive 15 minute intervals show no increase in	water?
10.	Contents of receiver allowed to reach room temperature and read to the nearest scale	division?
11.	Volume of water recorded and calculated in accordance with the method?	······
	% Water = 100 (Volume of water in receiver / Mass of sample)	
	ination of Volatile Distillates	
1.	Sample placed in the still?	
2.	350 mL of water and approx. 3 g [AMRL: ± 1 g] of sodium carbonate added to the sa	
3.	Gasket moistened with solvent?	
4.	Receiver used is the dilution trap specified in Section 4.3.2 and Figure 6?	······· <u> </u>
5.	Satisfactory assembly of components?	
6.	All connections vapor or liquid tight (caution if the apparatus leaks)?	manorarony
7.	Loose cotton (or similar) plug inserted in top of condenser?	······································
8.	Cold water circulated in jacket of condenser?	······································
9.	Heat applied and refluxing starts within 5 to 10 minutes?	
10.	Drip rate adjusted from 85 to 95 drops of distillate per minute?	
	Note: It may be necessary to add a second condenser or to reduce the rate of distilla prevent escape of the solvent.	tion somewnat to
11.	Distillation continued until three successive 15 minute intervals show no increase in	upper and
11.	lower levels of the diluent?	1.1
12.	Heat removed and solvent allowed to stand for 0.5 hours?	
13.	Contents of receiver read to the nearest scale division?	
13. 14.	Volume of dilute recorded and calculated in accordance with the method?	
17.	volume of antice recorded and calculated in accordance with the method?	
	% Diluent = 100 [(Volume of dilute in receiver * Sp. G. of dilute at 25°C) / M	[ass of sample]
COMM	ENTS (T110 / D1461):	(T110 / D1461)

(T164)	_
(D2172)	

		<u>APPARATUS</u>	Date:
Annoro	tua Common to all Mothoda (A. D. D. and E)	
<u>Appara</u> 1.	tus Common to all Methods (Solvent:	A, B, D, and E	
1.		/ reused solvents should not be used for testing.	
	(a) Trichloroethylene?		
		grade conforming to ASTM D4080?	
	(2) AASHTO	only: Reagent grade is required if running T17	70 using Trichloroethylene)?
		e, technical grade?	
	(c) normal-Propyl Bro	omide, conforming to ASTM D6368?	
		pene extractant, AASHTO Method A or E?	
		ctants that gel when exposed to water are not accept	
2.		ace exhaust system in a well-ventilated area?	
3.		ng 110 ± 5 °C (230 ± 9 °F) for warming sample?	
4.	AASHTO only: <u>Oven</u> , can m	naintain 149 to 163 $^{\circ}\!$	g (if moisture not determined)?
5.	Pan of appropriate size [AS	TM: 12 x 8 x 1 in.]?	
6.	Spatula or trowel?		
7.	AASHTO only: Balance, re	adable to 0.1% of sample mass, conforms to M	1231?
	ASTM only: Balance: acci	uracy of at least 0.01 percent of sample mass?	······
<u>Appara</u>	tus for Determining Mineral I	<u>Matter</u>	
	Ashing Method		
		r: capacity 1 or 2 L or 100 mL cylinder?	
	(b) Ignition dish, mini	mum capacity of 125 mL?	
	(c) Steam bath or hot	plate?	
	(d) Desiccator, large e	nough to contain ignition dish?	
	(e) Ammonium carbon	nate, reagent grade (as saturated solution or salt	
	(f) Class P (0.001 g) s	onelytical belongs available?) !
	(f) Class B (0.001 g) a	analytical balance available?nition furnace or Bunsen burner?	nce Laboratory
	(g) AASHTO only: Igi	ntion jurnace or Bunsen burner?	
or	Centrifuge Method		
	(a) Centrifuge capable	of 3000 r/min or greater?	
		/pe?	
		balance available [ASTM: Class GP1]?	
or	Volumetric Method		
	(a) Flask large enough	to hold extract?	
	(b) Constant temperatu	ure bath?	<u> </u>
		able to 0.2°F (0.1°C)?	
Additio	onal Apparatus for Method A		
1.	Centrifuge Extractor:		
		ith cover?	
	(b) Can be rotated at v	ariable speeds up to 3600 r/min?	
	(c) Apparatus set up sa	affely (not prone to explosions and installed in fi	
2.	Filter rings felt or naner to	fit rim of bowl?	
∠. or		ash content less than 0.2 percent?	
01	20 " usii paper iiitei iiiigs. t	2011 Content 1000 than 0.2 percent.	
COMM	IENTS (T164 / D2172):		(T164 / D2172)

(T164)	_
(D2172)	

			APPARATUS (Continued)	Date:
A ddit	ional An	paratus for Method B		
<u>Auun</u> 1.		x Extractor:		
1.	(a)		s jar free of scratches, cracks, or flaws?	
	(b)	Cylindrical metal	frames, one or two, each containing a conical filte	er cone support?
	(0)	(1) Bottom o	f lower cone above level of solvent?	
		(2) If two fra	mes, the upper frame can be supported on the low	ver?
2.	Conde	enser fits top of reflu	c extractor?	
3.	Filter	paper: medium grade	, fast filtering, completely lines cone?	
	(a)	Electric hot plate.	thermostatically controlled?	
	(b)	AASHTO only: Th	nermal-distributing protective pad of adequate thi	ckness, approx. 3 mm thick?
	(-)	ASTM only: The	rmal-distributing protective pad made of heat-res	sistant?
Addit	ional Anı	paratus for Method D		
1.		ction Kettle:		
	(a)		ate glass?	
	(b)			
	(c)	Condenser top, fit	ed to basket?	
2.		filter sacks with elast	ic hem for lining basket?	
A ddit	ional An	paratus for Method E		
Addii 1			ally as shown in Fig. 6a, b, and c):	
1.	(a)		eferably vane type)?	
	(a) (b)	Filter support plate		
	(0)		d with holes small enough to support the filter pap	per and numerous
			ensure adequate suction?	
		(2) Overlaps	or fits just inside "O" ring?	
			ng?	
2.	Filter	naner: medium grade	fact filtering 330 mm diameter?	·····
2. 3.	1 11tCl	TO: Sample contain	, fast filtering, 330 mm diameter? er, 3.8 L (4 qt.) capacity or greater?	ce laboratory
<i>J</i> .	ACTA	10. Stainless steel hee	ker, 9 qt. capacity?	<u></u>
4.	Two f	lasks 4000 mL canad	ity each?	······
5.	Glass	oraduate olass with a	capacity of 500 mL?	
6.	Wash	bottle filled with wat	er?	······
7.	Spatu	la and large mixing sr	oon?	······
8.	Stiff-l	oristle brush approxir	nately 1 in. wide?	······
9.	Dial t	hermometer 50 to 18	0°F (10 to 82°C)?	
10.	Ontio	nal· Ultrasonic clear	er, 4 qt. minimum capacity with insert tray?	
11.	Ethyl	alcohol denatured [A	ASHTO only: optional]?	······
12.	Ontio	nal AASHTO only: 1	2 in. diameter No. 16 and No. 200 sieves?	
13.			ratus for Method E-II (Method for slow-filtering)	
	(a)		pacity?	
	(b)		a 4 in. diameter?	
	(c)	Metal tongs for ha	ndling watch glass?	
	(d)	Diatomaceous silie	ca filtering aid, conforming to requirements of AS	TM D604 - Type B?
	(-)		<i>G</i> ,	J.F

COMMENTS (T164 / D2172):

(1164)	
(D2172)	

				PROCEI	<u>DURE</u>	Da	te:
Sar	nple	Preparation:					
1.	-	If necessary, mix	cture warmed in pan	at $230 \pm 9^{\circ}F$ (110	± 5°C) until it can	be handled?	
2.		Particles of mixture separated with spatula or trowel?					
3.	Sample obtained by splitting or quartering, conforms to minimum sample mass table below?						
		Table of minimum sample masses for T164/D2172 conforms to minimum sample mass table below					
		No. 4		1/2 in.		1 in.	1.5 in.
		1/2 kg	1 kg	1.5 kg			
4.				or multiple extracti	ons if necessary?	<u> </u>	
5.		If necessary test	specimen for moist	ure determination of	obtained?		······
٠.		11 1100000011, 0000	op comicinate and the second				
Wa	iter D	Determination					
1.			on test portion deteri	mined? (W_i)			
2.			tumen is NOT requir				
							r to extraction?
		(u) ANISITI	Dried to constant	mass prior to extra	ection at 230 + 0° F	7 (110 + 5°C)2	
	or	(b) AASHT	O only: Moisture de	etermined by T3202	Tmin 1000 a sampl	le over at either IM	F mixing range or
	O1						minute intervals]?
	or	(c) Moistur	re determined by T1	, <i>urteu joi 90 ± 5 ii</i> 10/D1/61 (apparat	unuies, unu inen v us must ha availah	le must be used fo	or recovery)?
3.	OI	Mass of water of	alculated by multiply	ing percent water.	as must be avamab	of extraction test	portion? (W_2)
٥.		Note to Assessors	: If the sample is dried	to constant mass W	is the dry mass and	d W. is 0 (the mass of	of the water)
		11010 10 2133033013.	ij ine sampie is arieu	to constant mass, m	is the ary mass, and	i m 2 is 0 (the mass c	y the water).
Evi	tracti	on Procedure by I	Method A (Centrifu	ge Method)			
1.	nacti				$at 230 \pm 0$ °F ()	$110 \pm 5\%$)] and we	eighed?
2.		Sample covered	with solvent and all	owed to disintegrat	e for not more than	: 10 ± 5 C)] and wi	
2. 3.		Bowl with colve	nt and cample place	I in extraction appo	e 101 1101 11101e tilai oratus?	II 1 III. (
<i>3</i> .		Weighed test no	rtion placed into box	1111 CXII action appo	natus:		
- . 5.		Dry filter ring fit	tted around edge of 1	owl and cover cla	mned tightly on bo	19	
5. 6.		Container place	lunder drain to colle	oot extract?	inped tightiy on oo	W1:	······
0. 7.		Centrifuge starte	d revolving clowly	and eneed increased			······
8.		Maximum spaed	not greater than 260	mu specu mercasei	i gradually (ence Lab	oratory —
o. 9.		Centrifuge conti	nued until solvent ce	oces to flow?			
э. 10.		Centrifuge conti	ad and 200 ml or m	ases to now!	d9		·····
11.		Stens (6) through	h (10) repeated addi	ng at least 3 increr	nents of solvent?	•••••	
12.		Last extract clea	r and not darker ther	light etropy color?	ichts of solvent!	•••••	
13.		Contents dried in	anu not uarker thai	til fimes dissinate	ງ		······
13. 14.		Sample can be d		itii tuilles dissipate			······
14.			ng and aggregate tra	neferred to tored m	etal non than drie	d to constant mass	in on oven at
			C (230 ± 9 F)?	······································	1 4 1 . 4 4	110 + 500 (220 + 1	00F) ("It
	or		ate dried to constant				
		dried se	eparately to constant	mass in an oven at	110 ± 5°C (230 ±	9°F)?	
	or	(c) Aggreg	ate and filter ring dr	ied in the bowl to o	constant mass in ar	oven at $110 \pm 5^{\circ}$	C (230 ± 9°F)?
		(0 : 1) 701					
15.			ash filter paper is u				
		with aggregate to	o avoid loss?				
16.			of filter ring subtract				
		mass of extracte	d aggregate? (W_3)				
17.		Mineral matter is	n the extract determi	ned by one of the s	specified procedure	es?	
~ -							/ma
CO	MM	ENTS (T164 / D2	2172):				(T164 / D2172)

Revised 2011-03-25

(T164)	
(D2172)	

	PROCEDURE (Continued) Date:
Extra	ction Procedure by Method B (Reflux Method)
1.	Filter paper(s) dried to constant mass in oven [AASHTO only: at $230 \pm 9 \%$ ($110 \pm 5 \%$)] and weighed?
2.	Mass of each frame with filter paper determined to the nearest 0.5 g and recorded?
3.	Place test portion in frame(s)?
4.	Mass of each loaded frame determined to the nearest 0.5 g and recorded?
5.	Solvent poured into glass jar?
6.	Frame with supporting legs placed in jar?
7.	Solvent level below tip of cone in lower frame?
8.	Optional: denatured alcohol used to wet filter paper?
9.	If multiple frames, upper frame(s) stacked on bottom frame?
). 10.	Insulating pad and cylinder placed on hot plate [AASHTO only: optional]?
11.	Gentle steady flow of cold water circulated through covered condenser?
12.	Heat adjusted so that solvent boils gently?
13.	Steady flow drips into cone?
14.	If necessary, adjust temperature of hot plate to maintain the solvent stream at a rate to keep
14.	test portions completely covered, but not overflowing the filter cones?
15.	Extraction continued until running from tip of lower cone appears a light straw color when
13.	viewed against white background?
16.	Heat shut off, but not condenser water?
10. 17.	Apparatus allowed to stand until cool enough to handle?
18.	Condenser turned off and removed from cylinder?
10. 19.	Loaded frame(s) removed from jar and dried in air?
19. 20.	Dry frame(s) to constant mass in oven at $230 \pm 9^{\circ}F$ ($110 \pm 5^{\circ}C$)?
20. 21.	Mass of extracted aggregate determined? (W_3)
21. 22.	Mineral matter in solution determined by one of the procedures specified?
22.	witherai matter in solution determined by one of the procedures specified?
Evtra	ction Procedure by Method D (Kettle Method)
1	ction Procedure by Method D (Kettle Method) Filter sack placed in extraction basket and mass determined with the tare pan and total tare mass determined?
2.	Test portion placed in filter sack and total mass determined, then mass of test portion calculated?
3.	Suspension rod attached to loaded basket and assembly set into the extraction kettle?
<i>3</i> . 4.	Approximately 600 mL [AMRL: ± 100 mL] of solvent poured over the test portion?
5.	Condenser cover placed on kettle?
6.	Cold water started through condenser cover?
7.	Basket raised to immersion level, approx. 1/2 in. above bottom of kettle?
7. 8.	Extractor placed on hot plate and heated to a gentle boil?
9.	Heating continued for 15 to 30 minutes in immersion position?
10.	Basket raised [AASHTO only: to refluxing level]?
11.	Heat increased to maintain active boiling until solvent dripping from the basket appears light straw color?
12.	Extractor removed from plate and allowed to cool?
13.	Basket removed from kettle and filter sack removed from basket and contents distributed into tared pan?
13. 14.	Filter sack placed on top of aggregate and dried on a steam bath and then in an oven at
14.	
15	$230 \pm 9^{\circ}F$ (110 ± 5°C) to constant mass?
15.	Extraction transferred to a 1000-mL graduate?

Extractor washed clean with solvent then washings added to the extract?

Mass of extracted aggregate determined? (W₃).....

Mineral matter in the extract determined by one of the procedure specified?.....

COMMENTS (T164 / D2172):

16.

17.

18.

(T164)	_
(D2172)	

	PROCEDURE (Continued) Date:
Evtro	ction Procedure by Method E (Vacuum Method)
1.	AASHTO: Filter paper (more than 1 may be used) dried to constant mass in oven at 230 ± 9 °F (110 ± 5 °C)?
1.	ASTM: Filter paper dried to constant mass?
2.	Sample mass determined and recorded?
3.	Cooled below 130°F (54°C)?
4.	200 mL alcohol added with care [AASHTO only: optional, should not be need for if using terpene]?
5.	Approx. 700 mL methylene chloride [AASHTO only: or other approved solvent] added?
6.	Stirred until bitumen visually in solution or ultrasonic cleaner used?
7.	Extractor assembled with dry filter paper (finger tight) centered?
8.	Optional AASHTO only: No. 16 and No. 200 sieves used? [ASTM only: sieves are not acceptable]
9.	Optional: Additional steps for Method E-II (for slow-filtering samples)
9.	Note: A correction may be necessary to account for any diatomaceous silica that is washed through the filter.
	See note 23 in AASHTO T164.
	(a) 50 to 200 g pre-dried filtering aid and 500 mL of solvent added to 1000-mL flask?
	(b) Flask swirled until filtering aid is completely in suspension?
	(c) Solvent and filtering aid mixture poured over filter?
	Note, AASHTO only, optional: Two pre-dried filters separated by an additional 50 to 100 g of
	filtering aid may be used to facilitate flow of liquid.
10.	Vacuum started and solution decanted onto filter or through sieves?
11.	Watch glass used if filtering aid needed [AASHTO only: and sieves not used]? (E-II)
12.	ASTM only: Vacuum stopped when all solvent is removed from filter paper?
13.	Sample remaining in the container covered with solvent and stirred?
14.	Steps 10 through 13 repeated until the solution is a light straw color and aggregate is visually clean?
15.	AASHTO only: If terpene was used, aggregate rinsed with water, preferably above 43 °C (110 °F), in the
	same manner that the bitumen was rinsed off with solvent?
16.	All aggregate carefully transferred onto the filter [AASHTO only: if terpene used. If not, skip to (20)]?
17.	Any aggregate clinging to the container washed onto the filter with the solvent?
18.	Aggregate dried using vacuum?
19.	Aggregate scraped towards center from funnel ring?
20.	Vacuum stopped and aggregate transferred to tared drying pan?
21.	Funnel ring and filter paper brushed?
22.	Aggregate and filter paper dried to constant mass in an oven at $230 \pm 9^{\circ}F$ ($110 \pm 5^{\circ}C$)?
23.	AASHTO only: If low-ash filter paper is used, is it burned in the pan with aggregate to avoid loss?
24.	Mass of aggregate and filter in the pan recorded?
25.	Mass of filter and pan subtracted to determine mass of extracted aggregate? (W_3)
26.	Mineral matter in solution determined by one of the procedures specified?
	(Note to Assessor: The mineral matter determination is not required for laboratories performing Method E for
	plant control testing only. In all other cases, the mineral matter determination must be demonstrated during the

assessment. Add standard Observation finding for Plant Control to final report.)

COMMENTS (T164 / D2172):

COMMENTS (T164 / D2172):

QUANTITATIVE EXTRACTION OF BITUMEN FROM HMA

(T164)	_
(D2172)	

	PROCEDURE (Continued) Date:
	ineral Matter Determination by Ashing Method
1.	Volume [AASHTO only: or mass] of total extract and washings recorded? $(W_I \text{ or } V_I)$
	Note to Assessors: Watch out, AASHTO labels both this volume and the original sample mass as W_1 .
2.	AASHTO only: Ignition dish conditioned in furnace or on Bunsen burner, then cooled in a desiccator?
3.	Ignition dish mass determined to 0.001 g?
4.	Extract thoroughly agitated, approx. 100 mL [AASHTO: or 100 g] immediately measured into ignition dish?
5.	ASTM only: Volume after removing ignition dish portion determined? (V ₂)
6.	Ignition dish evaporated to dryness on steam bath or hot plate?
7.	Residue ashed at dull red heat [500 - 600°C (932 to 1112°F)] and cooled?
8.	Mass of the ash determined?
9.	5 mL of saturated ammonium carbonate solution added per 1 g of ash?
10.	Digested at room temperature for 1 hour?
11.	Dried in oven to constant mass at 110 ± 5 °C (230 ± 9 °F) [ASTM: constant mass at 100 °C]?
12.	Cooled in desiccator and net mass of ash determined on analytical balance to the nearest 0.001 g? (G)
13.	Mass of mineral matter calculated $\{AASHTO: G \times (W_1/100)\} \{ASTM: G \times (V_1/(V_1-V_2))\}? (W_4) \dots $
	ineral Matter Determination by Centrifuge Method
1.	Empty centrifuge cup mass determined to 0.01 g?
2.	Centrifuge cup placed in centrifuge?
3.	Container positioned at appropriate spout to catch effluent?
4.	Extract transferred to container suitably equipped with feed control (clamp or valve)?
5.	Extract container rinsed several times with clean solvent, and washings added to feed container?
6.	Centrifuge started and allowed to reach a constant speed?
7.	Feed line opened and extract fed into centrifuge at a rate of 100 to 150 mL/min.?
8.	Feed mechanism rinsed several times with clean solvent until effluent is essentially colorless?
9.	Centrifuge allowed to stop and cup (or bowl) removed?
10.	Outside of cup cleaned with solvent and allowed to evaporate in a funnel or steam hood?
11.	Cup dried in oven at $230 \pm 9^{\circ}F$ ($110 \pm 5^{\circ}C$)?
12.	Cup cooled and mass of the cup with residual material determined to nearest 0.01 g immediately?
13.	Increase in mass of cup reported as the mass of the mineral matter? (W_4)
T . 13.6	
	ineral Matter Determination by Volumetric Method
1.	Flask calibrated and tared mass recorded?
2.	Extract placed in tared flask?
3.	Flask brought to within \pm 0.1°C (0.2°F) of calibration temperature in controlled-temperature bath?
4.	Flask filled with solvent at same temperature?
5.	Level of liquid in flask brought up to the neck of the flask?
6.	Stopper inserted and flask dried?
7.	Flask weighed to nearest 0.1 g?
8.	Calculations made according to the book (AASTHO Section A1.3.2.2, ASTM Section 11.6.3.2) to
	determine mineral matter? (W_4)
C 1 1 1	' (A 1 1/P) 1 (C / /
	ion of Asphalt Binder Content
1.	Asphalt binder content percentage calculated?
	% Asphalt Binder Content = $(W_1 - W_2) - (W_3 + W_4)$ x 100
	where:
	where: $(W_I - W_2)$ $W_I = \text{mass of test parties}$ $W_I = \text{mass of systematical mineral aggregate}$
	W_1 = mass of test portion W_3 = mass of extracted mineral aggregate W_2 = mass of water in test portion W_3 = mass of mineral matter in the extract
	W_2 = mass of water in test portion W_4 = mass of mineral matter in the extract

BULK SPECIFIC GRAVITY OF COMPACTED HMA USING SATURATED SURFACE-DRY SPECIMENS

(T166) _	
(D2726)	

		<u>APPARA</u>	TUS	Date:
Commo	on Annaratı	s for all methods		
<i>1</i> .		Balance conforms to M231 for class required	l for principal sample mass of	
	samples t	ested (G2 balance for specimens over 200 g)?		
	ASTM:	Balance conforms to D4753, sensitive to 0.1 g	for 100.1 - 999.9 g sample (GP.	2)?
		Balance standardized according to interval in		
2.	Water bat	h:		
	(a)	Maintains $25 \pm 1^{\circ}\text{C} (77 \pm 1.8^{\circ}\text{F})$?		
	(b)	Equipped with overflow outlet to maintain cons	stant water level?	
3.		Drying oven at $52 \pm 3 \%$ (125 $\pm 5 \%$)?		
	ASTM:	Drying oven at 110 ± 5 °C (230 ± 9 °F)?		······
		Orying oven standardized according to interva		
4.	<i>AASHTO</i>	only: Room temperature: $25 \pm 5 \%$ (77 $\pm 9 \%$)	?	
5.	Cloth tow	el, damp (considered damp when no water can	be wrung from it)?	
1.	Water bat (a) (b) (c) (d) (e) (f)	tus for AASHTO Method A and ASTM h suspension: AASHTO only: Suspension from center of balan Holder and sample completely immersed? AASHTO only: No trapped air bubble exists un Can determine constant mass of specimen to 0.0 Water bath deep enough to completely immerse Suspension wire of smallest practical size? Note to Assessors: Ropes, strings, and sash cords a	oder specimen?	
		tus for <mark>A</mark> ASHTO Method B		
1.	Thermom	C (19 to 27 °C, graduated to 0.1 °C)?	ls Reference La	boratory
or		F (66 to 80°F, graduated to 0.2°F)?		
		ly: temperature measuring device presented		
2	D3000 (0	months), see records?	າ	······································
2.	Calibrate	d volumeter and tapered lid with capillary bor	27	
COMM	MENTS (T1	66 / D2726):		(T166 / D2726)

Revised 2011-03-25

BULK SPECIFIC GRAVITY OF COMPACTED HMA USING SATURATED SURFACE-DRY SPECIMENS

(T166)	

AASHTO PROCEDURE	Date:
$rac{1}{3}$ $^{\circ}$ C (125 \pm 5 F) to constant mass (0.05 percent)?

<i>AASH</i>	TO Method A (sequence of steps is optional)
1.	Dry sample used or sample dried at $52 \pm 3 \%$ (125 $\pm 5 \%$) to constant mass (0.05 percent)?
	Note to Assessors: Sample can either be checked for constant mass at successive 2 hr. intervals or dried overnight.
2.	Sample cooled to room temperature, $25 \pm 5 ^{\circ}\text{C}$ (77 $\pm 9 ^{\circ}\text{F}$), and dry mass recorded? (A)
<i>3</i> .	Sample immersed for 4 ± 1 min.?
4.	Immersion water at $25 \pm 1 $ °C (77 $\pm 1.8 $ °F) (check with AMRL thermometer)?
5.	Each specimen immersed and weighed individually?
6.	Immersed mass determined? (C)
7.	Sample removed from bath and quickly blotted with damp towel (not to exceed 5 s)?
8.	Saturated surface-dry mass determined? (B)
9.	Bulk Specific Gravity calculated as $\{A / (B - C)\}$?
10.	Percent Water Absorbed calculated $\{(B-A)/(B-C)\}$?
AASH	TO Method B (sequence of steps is optional)
1111011	Note: Method B is not acceptable for samples with air voids greater than 6 percent.
1.	Dry sample used or sample dried at $52 \pm 3 \text{C} (125 \pm 5 \text{F})$ to constant mass (0.05 percent)?
	Note to Assessors: Sample can either be checked for constant mass at successive 2 hr. intervals or dried overnight.
2.	Sample cooled to room temperature, $25 \pm 5 $ °C (77 $\pm 9 $ °F), and dry mass recorded? (A)
3.	Sample immersed at least 10 min?
4.	Immersion water at $25 \pm 1 ^{\circ}\text{C}$ (77 $\pm 1.8 ^{\circ}\text{F}$) (check with AMRL thermometer)?
<i>5</i> .	Volumeter filled with distilled water at 25 \pm 1 °C (77 \pm 1.8 °F) and mass determined? (D)
6.	Sample removed from bath and quickly blotted with damp towel (not to exceed 5 s)?
<i>7</i> .	Saturat <mark>ed</mark> surface-dry mass determined? (B)
8.	Sample placed in water-filled volumeter and allowed to stand for 60 s?
9.	Volumeter water temperature brought to $25 \pm 1 \%$ (77 $\pm 1.8 \%$)?
10.	Volumeter covered and some water allowed to escape through the capillary bore of the tapered lid?
11.	Volumeter wiped dry and volumeter and contents weighed? (E)
12.	Volumeter wiped dry and volumeter and contents weighed? (E)
13.	Percent Water Absorbed calculated $\{(B-A)/(B+D-E)\}$ x 100 $\}$?
4 4 S H	TO Method C
2121011	Method C is for samples that are not required to be saved and contain a substantial amount of moisture.
1.	Procedure same as Method A or Method B except for determination of dry mass?
2.	Dry mass determined as follows:
	(a) Sample warmed in oven $110 \pm 5 \text{C} (230 \pm 9 \text{F})$ until soft?
	(b) Broken down to 1/4 in. particles?
	(c) Dried in oven to constant mass (2 hr. change less than 0.05 percent)?
	(d) Cooled to room temp. $25 \pm 5 \text{C}$ (77 $\pm 9 \text{F}$) and weighed? (A)
3.	Calculations same as Method A or Method B?
<i>J</i> .	Culculations same as Method A of Method B:
	Calculations (all methods)
1.	Percent water absorbed determined to be less than 2.0 percent?
2.	If the percent of water absorbed by the specimen exceeds 2.0 percent, T275 used to
2	determine the bulk specific gravity instead?
3.	Bulk Specific Gravity reported to nearest 0.001?
4.	Absorption reported to nearest 0.01?

(T166) COMMENTS (T166):

COMMENTS (D2726):

BULK SPECIFIC GRAVITY OF COMPACTED HMA USING SATURATED SURFACE-DRY SPECIMENS

(D2726)

		ASTM PROCEDURE Date:
Me	ethod	For Laboratory-Prepared Thoroughly Dry Specimens
1.		Specimen allowed to stand in room temperature air for at least 1 hr.?
2.		Mass of dry specimen determined? (A)
3.		Specimen immersed in bath for 3 to 5 min.?
	or	If specimen temperature differs from the water temperature by more than 2°C (3.6°F), is specimen immersed 10 to 15 min.?
4.		Immersion water at $25 \pm 1^{\circ}C$ (77 $\pm 1.8^{\circ}F$) (check with AMRL thermometer)?
<i>5</i> .		Immersed mass determined? (C)
6.		Sample quickly blotted with a damp towel?
<i>7</i> .		Saturated surface-dry mass determined? (B)
<u>Me</u>	ethod	For Specimens That Contain Moisture or Solvent, or Both
1.		Specimen immersed in bath for 3 to 5 min.?
	or	If specimen temperature differs from the water temperature by more than 2°C (3.6°F), is specimen immersed 10 to 15 min.?
2.		Immersion water at $25 \pm 1^{\circ}C$ ($77 \pm 1.8^{\circ}F$) (check with AMRL thermometer)?
<i>3</i> .		Immersed mass determined? (C)
4.		Sample quickly blotted with a damp towel?
5.		Saturated surface-dry mass determined? (B)
6.		Specimen dried in an oven at $110 \pm 5^{\circ}C$ (230 $\pm 9^{\circ}F$) oven to constant mass?
•		Note: Other means of drying (such as microwave) ok as long as specimen is not
		over-heated and documentation exists showing results are equivalent to oven drying.
<i>7</i> .		Specimen cooled and weighed in air? (A)
Fir	nal C	alculations (both methods)
1.		Bulk Specific Gravity calculated $\{A/(B-C)\}$?
<i>2</i> .		Percent Water Absorbed calculated \{ ((B - A) / (B - C)) x 100 \}?
<i>3</i> .		Density of specimen calculated { Bulk Sp. Gr. x density of water (997.0 (kg/m³) or 62.24 (lb/ft³)) }?
4.		Percent water absorbed determined to be less than 2.0 percent?
<i>5</i> .		If the percent of water absorbed by the specimen exceeds 2.0 percent, D1188 used to determine the bulk specific gravity instead?

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(D2726)

COMPRESSIVE STRENGTH OF BITUMINOUS MIXTURES

(T167)	_
(D1074)	

		<u>APPARATU</u>	J <u>S</u>			Da	ite:		
_									
	sting Machine				.				
1.	Maker:			•••••	Capaci	ty:			
2.	AASHTO: capable of controlled speeds of 2.								
	ASTM: having a range of controlled speeds specimens to 0.4 in. (10.2 mm)/min for 8-in.	covering at i	east V.1	in. (2.5	mm)/mi	n jor 2-i	n. (30.8-	mm)	
3.	Upper bearing block:	(203.2-mm) S	ресіте	ns:	•••••	••••••		•••••	···
٥.	(a) Spherically seated?								
	(b) Center of sphere coincides with bear								
	(c) Block rotates freely and can be tilted								
	(d) Diameter of bearing face slightly gre	ater than diar	neter of	largest s	specime	n tested?			
	(e) Bearing face plane to 0.025 mm (0.0	01 in.)?							
4.	Lower bearing block:	,							
	(a) Rests on platen and is seat for specin	nen?							
	(b) Diameter of bearing face greater than	n diameter of	largest s	specime	1 tested?				
	(c) Bearing face plane to 0.025 mm (0.0								
	olds and Plungers								
1.	Standard molds								1
	Note: record diameters	1	2	3	4	5	6	7	8
(a)	Inside diameter of molds:								
	101.60 – 101.73 mm (4.000 – 4.005 in.)?								
(b)	Diameter of plungers < 101.6 mm (4.000 in.)?	Top:				Bottor	n:		
(c)	Max. difference between I.D. of molds and	1000							
(•)	dia. of plungers is 1.27 mm (0.050 in.)?								
	dia. of plungers is 1.27 mm (0.050 m.):								
2			4 4 4:	- 414 - C	21 1		1		
2.	Other molds are acceptable provided the diam								
	(a) Top plunger may be of various desig(b) Bottom plunger may be of various designation	II, lace at leas	loogt 12	7 mm (111.) tilic 1/2 in \ t	K / biole?	orat	:ory:	
	(c) Plunger 50 ± 4 mm $(2 \pm 1/8 \text{ in.})$ in he								
3.	Mold supports capable of supporting mold cy								
4.	ASTM only: 2 steel bars $1 \pm 1/8$ in. (25.4 \pm								
т.	7151111 Only. 2 seed but 5 1 ± 1/0 til. (23.4 ±	3.1 mm) squi	ii C uii u	ui icusi .	<i>J III.</i> (70	. 2 mm) t	ong	••••••	•••
Oth	ner Apparatus								
1.	AASHTO: Oven capable of control within ± 3	3°C (+5°F)?							
	ASTM: Oven controlled within $\pm 5^{\circ}F$ ($\pm 3^{\circ}C$								
2.	Hot plate, equipped with a rheostat, for mixin								
3.	Optional: hot water bath	C							
	(a) Large enough to hold 3 sets of 100 n	nm molds wit	h plunge	ers?					
	(b) Capable of being maintained at just u	ınder boiling	point?						
	(c) Heater, if electric, provided with con	tinuously var	iable co	ntrol?					
4.	Air bath capable of being maintained at 25.0	± 0.5°C (77 ±	1°F) [A	STM: 2	77 ± 1.8	°F (25 ±	<i>0.5℃)</i>]?	·	
5.	Mixing machine for sample preparation?								
6.	Spatulas, one limber and one stiff?								
7.	Ejection device, ejection head has smooth, un	iform rate of	travel? .						
8.	Balance AASHTO: Class G2 balance								
	ASTM: Balances or scales	and weights	meeting	g Specif	ication l	D4753, G	<i>P2?</i>	•••••	•••
9.	ASTM only, Thermometer:			• . •			OF (0 ===	a) .	
	(a) Calibrated liquid-in-glass thermome								
	or (b) conforming to the requirements of s Electronic thermometer (RTD, PRT)	pecification .	E 2231?	ottor as	9	•••••	••••••	••••••	···
	or (b) Electronic thermometer (RTD, PRT,	ii ki) oj eq	uui OF D	ener acc	лису?.	••••••	•••••	••••••	•••

COMMENTS (T167 / D1074):

(T167 / D1074)

COMPRESSIVE STRENGTH OF BITUMINOUS MIXTURES

(1167)	_
(D1074)	

	PROCEDURE Date:
Prenara	tion of Test Mixtures
1.	Initial batch mixed to "butter" mixing bowl and stirrers?
2.	Trial specimen molded to determine proper mass of batch?
3.	Bowl emptied and batch used for trial specimen discarded?
4.	Bowl and stirrers cleaned by scraping with limber spatula?
5.	Wiping with cloth or washing with solvent avoided?
6.	AASHTO: Mixing and compacting temperatures based upon the temperature-viscosity curve for the asphalt
	(170 ± 20 cSt, 280 ± 30 cSt)?
	ASTM: Mixing temperature does not exceed 347°F (175°C)?
7.	Aggregate and mixing bowl heated no more than 28°C (50°F) above the mixing temperature?
8.	Aggregate added to bowl and dry-mixed?
9.	Bituminous material quickly weighed into aggregate?
10.	Asphalt and aggregate mixed with minimal "fanning action"?
11.	Mixing completed within 90 to 120 seconds?
12.	Temperature of mixture after mixing about 3 to 5°C (5 to 9°F) above the compacting temperature?
13.	If necessary, mix reheated using a hot plate, oven, or similar device?
Plant n	ixtures_
1.	Sample reduced according to T248/C136 to slightly more than amount needed to fabricate specimen?
2.	Mass of reduced sample adjusted to required mass by removing and discarding
	a small amount of the mixture?
3.	Both fine and coarse particles discarded to maintain proper gradation?
4.	Mixture placed in appropriate container and heated in an oven to the established mixing temperature?
5.	Sample removed from oven and thoroughly mixed until temperature is about 3 to 5°C (5 to 9°F)
	above the compacting temperature?
6.	Material may be placed into an oven for a short time if multiple samples tested (not more than 1 hour)?
	AASH I O Materials Reference Laboratory

COMMENTS (T167 / D1074):

(T167 / D1074)

COMPRESSIVE STRENGTH OF BITUMINOUS MIXTURES

(T167)	_
(D1074)	

	PROCEDURE (Continued) Date:
Molding	g Test Specimens
1.	Top plungers, bottom plungers, molds, and spatula preheated?
1.	(a) In a water bath just under the boiling point for at least 1 hr.?
or	
2.	Mold and plungers removed and wiped with a clean cloth containing a few drops of oil?
3.	With bottom plunger in place and molding cylinder supported on steel bars:
J.	(a) Approximately half of mixture placed in mold?
	(b) Fifteen blows with spatula struck around inside of mold?
	(c) Ten blows with spatula struck at random over mixture?
	(d) Penetration of spatula as deep as possible?
4.	Remaining mixture quickly placed in molding cylinder?
5.	Step 3, parts (a) through (d) repeated?
6.	Top of mixture left slightly rounded or cone shaped?
7.	Filled mold compressed between top and bottom plungers under about 1 MPa (150 psi) [AMRL: ~1,886 lb]?
8.	Support bars removed from under mold?
9.	Molding load increased to 20.7 MPa (3000 psi) for 2 min. [AMRL: ~37,727 lb]?
	Note: When tested in accordance with T165/D1075, molding load of 20.7 MPa (3000 psi) may be
	increased or decreased to achieve a target air void percentage or density percentage
10.	AASHTO only: Alternate methods of compaction may be used, provided approximately
	7 percent air voids are achieved?
11.	Specimen removed from mold with ejection device?
T . C	
Test Spe	Specimens 102 mm in diameter?
1.	Specimens 102 mm in diameter?
2.	Specimens 101.6 ± 2.5 mm in height?
3.	After ejection, specimens placed in oven for 24 hrs. at 60°C (140°F)?
4.	Specimens placed in air tight containers if NOT tested in compression within 24 hrs. after oven curing?
	AASH IO Materials Reference Laboratory
Procedu	ra
1.	Specimens cooled at room temp. for at least 2 hrs. after removal from curing oven?
2.	Bulk Specific Gravity of specimens determined by Method A or B of (T166 / D2726)?
3.	Specimens brought to $25 \pm 1^{\circ}\text{C}$ (77 ± 1.8°F) by storing in an air bath for at least 4 hrs?
<i>3</i> . 4.	Specimens tested in axial compression without lateral support?
4. 5.	Rate of vertical deformation uniform at 0.05 mm/min·mm (0.05 in./min·in.) of specimen height?
J.	Note: For 101.6 mm (4 in.) specimens, use a rate of 5.08 mm/min (0.2 in./min).
	1.000.1 of 101.0 mm (1 m.) specimens, use a rate of 5.00 mm/mm (0.2 m./mm).

COMMENTS (T167 / D1074):

(T167 / D1074)

RECOVERY OF ASPHALT FROM SOLUTION BY ABSON METHOD

(T170)	
(D1856)	

			<u>APPARATUS</u>	Date:
1.		Centrifu (a)	ge apparatus (either of the following): Batch unit capable of 770 times gravity?	······
		(a)	(1) Wide-mouth bottles [AASHTO only: 250 to 500 mL capacity]?	······
		or	(2) Cylindrical tubes, 6 or 8 in. long, with conical ends; capacity 100 mL?	
	or	(b)	Continuous unit capable of 3000 times gravity?	
			Note to Assessors: RCF (gravities) = $1.118 \times 10^{-5} \times r$ (in cm) $\times (RPM^2)$	
2.		Distillat	ion flasks (2):	
2.		(a)	Wide-mouth, flat bottom, 250 mL extraction flasks?	
		(b)	Cork with holes for delivery tube, aeration tube, funnel and/or thermometer?	
		()	, ,	
3.		Suitable	flask for the receiver?	
		D. 11		
4.		Delivery	v tube: 10 mm I.D., goose-neck shaped glass tube connects flask to condenser?	······
5.		Inlet aer	ation tube: at least 180 mm long having a 10 mm bulb with 6 staggered 1.5 mm l	holes?
6.		Dictillat	ion flask heater:	
0.		(a)	Electric heating mantle with variable transformer?	
	or	(b)	Oil bath with means of measuring temperature of bath?	
	or	(c)	Fluidized sand bath with means of measuring temperature of bath?	<u>-</u>
7.		Water ja	acketed condenser with 200 mm minimum jacket length?	
8.		<u>Thermor</u>	meter: ASTM 7C or 7F?	
0				
9.			w meter capable of indicating flow up to 1000 mL/min. (CO ₂)?	
10.		Separato	ory funnel: 125 mL capacity or larger (required only if Abson apparatus is	boratory
		used for	primary distillation)?	
11.			on solvent:	
	0.14	(a) (b)	Trichloroethylene, reagent grade? Methylene Chloride, reagent grade [AASHTO only: in case of dispute]?	
	or or	(b) (c)	ASTM only: Normal Propyl Bromides (nPB), conforming to Specification De	
	or	(d)	AASHTO only: Technical grade: Methylene Chloride OR Trichlor., type I, Fed.	
	01	(u)	(1) If technical grade solvent used, has blank been run (recommended)?	
			(2) If so, by technique of Note 1 (70 g asphalt with known properties disso	
			into 800 mL solvent then recovered by T170)?	
12.		Supply	of carbon dioxide gas?	
14.		<u> </u>	of Carbon Groving gast	······
CO	MM	ENTS (T	170 / D1856):	(T170 / D1856)

RECOVERY OF ASPHALT FROM SOLUTION BY ABSON METHOD

(1170)	_
(D1856)	

	PROCEDURE Date:
Sample	Preparation:
1.	Is sample a solution from an extraction of sufficient mass to provide approximately 75 to 100 g of recovered asphalt?
2.	AASHTO only: Asphalt mixture heated in covered container until workable at $110 \text{C} (230 \text{F})$ for no longer than 30 minutes?
3.	Extraction apparatus clean and free of petroleum distillates?
4.	Was all of the asphalt in the mixture extracted?
5.	Time extraction started: Extraction method A [AASHTO only: or E] used?
Testing:	
1.	Centrifuging:
	(a) Solution centrifuged at 770g for 30 or more minutes?
or	(b) Centrifuged continuously at not more than 150 mL/minute at not less than 3000g?
2.	Solution concentrated to approximately 200 to 300 mL by:
۷.	(a) Any primary distillation that meets the following criteria:
	(1) Has a flask large enough to hold all the solution from the extraction?
	(2) Solution from primary distillation transferred to Figure 1 assembly using several washes of solvent?
or	(b) Distillation started in assembly shown in Figure 1: (1) Separatory funnel in place through cork?
3.	Bulb of aeration tube lowered to make contact with the bottom of the flask?
4.	AASHTO: Slow introduction of CO ₂ begun at the beginning of the distillation (about 100 mL/min)?
5.	When temperature reaches 157 to 160°C (315 - 320°F), gas rate increased to 900 mL/minute? Time temperature reaches 160°C (320°F):
6.	Temperature at 160 to 166°C (320 - 330°F) and gas flow maintained at 900 mL/minute for 10 minutes (minimum) or until dripping stops?
7.	Gas flow minimum times: (a) Gas flow and heat maintained for an additional 5 minutes after the last drop (CO ₂ flow never
	less than 15 minutes)?
	(b) If residue in flask is highly viscous and the expected penetration is less than 30, maintain the gas flow and temperature for 20 - 22 min., including initial 15 minutes?
8.	CO ₂ flow never less than 15 minutes? Time gas flow and heat cut off:
9.	CO ₂ flow never less than 15 minutes? Time gas flow and heat cut off: Ash content of recovered asphalt determined in accordance with (T111 / D2939)?
10.	Elapsed time between time extraction started and time gas flow and heat cut off 8 hrs. or less?
11.	Residue retained for further testing?

COMMENTS (T170 / D1856):

(T170 / D1856)

COMMENTS (T209 / D2041):

MAXIMUM SPECIFIC GRAVITY OF HMA (RICE TEST)

$(T209)_{-}$	
(D2041)	

				APPARATUS Date:	
1		Contain			
1.		Contair (a)		um bowl (used for weighing in air and water):	
		(u)	(1)	Either metal or plastic?	
			(2)	Diameter approximately 180 to 260 mm (7 to 10.25 in.)?	
			(3)	Height approximately 160 mm (6.3 in.)?	
			(4)	Equipped with a transparent cover with a rubber gasket?	
			(5)	Using a small sample in a large container avoided?	
			(6)	AASHTO only: Capacity between 2,000 and 10,000 mL?	
	or	(b)	Vacuu	um flask (used for weighing in air only):	
		(-)	(1)	Thick-walled volumetric glass flask?	
			(2)	Fitted with a rubber stopper with a connection for the vacuum line?	
			(3)	Using a small in a large container avoided?	
			(4)	AASHTO: Capacity between 2,000 and 10,000 mL?	
				ASTM: Approximately 4000 mL capacity?	
	or	(c)	AASH'	TO only: Pycnometer (for weighing in air only):	
	-	(-)	(1)	Glass, metal, or plastic pycnometer?	
			(2)	Using a small in a large container avoided?	
			(3)	Capacity between 2,000 and 10,000 mL?	
 4. 		Balance (a) (b) (c) Vacuum (a) (b)	Class (For W permit AASH) a pump of Capab or less Note: When	G2 GP2 [ASTM only: readable to nearest 0.1 g]?	ry
			flasks,	or equivalent], is installed between the vacuum source and the vacuum vessel?	
5.				rement Device? (Reads in mm of Mercury)	
		(a)		ected at the end of the vacuum line using an appropriate tube and either a "T" connector	
				of the vacuum vessel or by using a separate opening in the top of the vessel?	
		(L)		The manometer is not to be situated on top of the vessel but adjacent to it, to avoid damage.	
		(b)		TO: Pressure gauge standardized every 12 months to be accurate to a careful of the control of th	
				this is the control of the control o	
				assessor: a mercury manometer does not need to be standardized. It is the standard.	
6.		ASTM in vacu	only: <u>M</u> um line	Aanometer or vacuum gauge, connected either directly to the vacuum source or close to the source?	

Revised 2011-03-25

(T209 / D2041)

COMMENTS (T209 / D2041):

MAXIMUM SPECIFIC GRAVITY OF HMA (RICE TEST)

(T209)	_
(D2041)	

	APPARATUS (Continued) Date:
7.	Mechanical Agitation Device [AASHTO only: Method A only] (a) Capable of applying a gentle but consistent agitation of the sample?
8. or	Thermometric Device: (a) Calibrated liquid-glass thermometer, with subdivisions and maximum scale error 0.5°C (1°F)? (b) Any other thermometric device of equal accuracy, precision, and sensitivity?
9.	Water Bath, maintains a constant temperature between 20 and 30°C [ASTM only: 25 ± 1 °C (77 ± 1.8 °F)]?
10.	Bleeder Valve, attached to the vacuum train?
11.	Drying Oven, AASHTO only: maintaining_135 \pm 5 $^{\circ}$ C (275 \pm 9 $^{\circ}$ F) or 105 \pm 5 $^{\circ}$ C (221 \pm 9 $^{\circ}$ F)?
12. Note to A	Apparatus for Supplemental Procedure for Mixtures Containing Porous Aggregate Not Completely Coated ssessors: Typically laboratories should not be demonstrating this supplemental procedure. It is included for reference. (a) Electric fan?
Standare 1. 2. 3. 4.	ization [ASTM: Calibration] of Container: AASHTO only: Water used to fill the container at a temperature between 20 and 30 °C?
5.	One of the following: Weighing-in-water determination (bowl): (a) Bowl filled with water [ASTM: immersed in water] and mass-in-water determined? (B)
	Weighing-in-air determination (bowl): (a) Bowl filled with water [ASTM: immersed in water]? (b) ASTM only: Lid placed on the bowl while submerged? (c) Outside of bowl dried and mass-in-air determined? (B) (d) ASTM only: Procedure repeated three times and results averaged? (D)
	Weighing-in-air determination (flask): (a) Flask filled with water?

(T209 / D2041)

MAXIMUM SPECIFIC GRAVITY OF HMA (RICE TEST)

(T209)	
(D2041)	

Date: _____

PRO	CEL	DURE	

- 1. Sample obtained by splitting or quartering?....

ASTM: Mass of sample as follows (samples larger than about two thirds the capacity of the container shall be divided into portions that are not less than 1250 g and the results averaged):

Nominal Maximum Aggregate	AASHTO: Min. Sample Size	ASTM: Min. Sample Size
37.5 mm or greater (\geq 1.5 in.)	4000 g	5000 g
19 mm – 25 mm (3/4 to 1 in.)	2500 g	2500 g
12.5 mm or smaller ($\leq 1/2$ in.)	1500 g	1500 g

3.		AASHTO only: Laboratory prepared samples conditioned and dried to constant mass (within 0.1%)
		in an oven at 135 \pm 5 °C for a minimum of 2 hours, or as appropriate to match the mix design?
	or	AASHTO only: Field samples dried to constant mass in oven at $105 \pm 5 ^{\circ}\text{C}$ (221 $\pm 9 ^{\circ}\text{F}$)?
		ASTM: Field samples dried to constant mass $105 \pm 5 \%$ (221 $\pm 9 \%$)?
4.		Particles of sample separated while warm by hand, using care not to fracture mineral fragments?
5.		After separation, fine aggregate particles not larger than 6.3 mm (1/4 in.) [ASTM: 6 mm]?
6.		Sample cooled to room temperature?
Tes	ting:	
1.		Placed in tared flask or bowl weighed and net mass of sample determined? (A)
2.		Water at approx. 25°C (77°F) added to cover sample?
3.		Vacuum increased until manometer reads 27.5 ± 2.5 mm Hg $(3.7 \pm 0.3 \text{ kPa})$?
4.		ASTM only: Vacuum achieved within 2 minutes of turning the vacuum system on?
5.		Container and contents agitated continuously by mechanical device [AASHTO: Method A]?
	or	<u>AASHTO only, Method B:</u> Container and contents agitated during the vacuum period by vigorously
		shaking at intervals of about 2 min.?
6.		Vacuum and agitation continued for 15 ± 2 min after vacuum is achieved?
7.		Vacuum released slowly [AASHTO only: at a rate not to exceed 8 kPa per second]?

COMMENTS (T209 / D2041):

(T209 / D2041)

MAXIMUM SPECIFIC GRAVITY OF HMA (RICE TEST)

(T209)	_
(D2041)	

PROCEDURE (C	ontinued)
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PROCEDURE (Continued)	Date:	

Note to Assessors: The laboratory should demonstrate one of the following methods of determining maximum specific gravity.
AASHTO/ASTM Weighing-in-water determination:
1. Bowl (without lid) and contents suspended in water?
Net mass of contents in water determined after 10 ± 1 min immersion? (C)
3. AASHTO: If temperature is not $25 \pm 1 \text{C}$ (77.0 $\pm 1.8 \text{F}$), mass corrected to 25C (77 $ \text{F}$)?
ASTM: Temperature of the water in bath $25 \pm 1^{\circ}C$ (77.0 $\pm 1.8^{\circ}F$) (temperature recorded)?
4. Theoretical maximum specific gravity calculated $\{A / (A - (C - B))\}$?
Theoretical maximum specific gravity calculated (A / (A – (C - B))):
AASHTO only: Weighing-in-air determination (any):
1. Flask, pycnometer, or bowl filled with water?
2. Contents adjusted to $25 \pm 1 \text{C} (77.0 \pm 1.8 \text{F})$?
3. Mass of filled container determined 10 ± 1 min. after removal of entrapped air completed? (E)
3. Mass of fitted container determined 10 \pm 1 min, after removal of entrapped air completed? (E)
4. Theoretical maximum specific gravity calculated $\{A / (A + D - E)\}$?
ACTM and Wainling in air datamain atom (Land).
ASTM only: Weighing-in-air determination (bowl):
1. Bowl and sample slowly submerged in the 25 ± 1 °C (77.0 ± 1.8 °F) bath for 10 ± 1 min?
2. Lid also placed in water bath at the same time as the bowl?
3. Lid placed on bowl without being removed from the water so as to avoid entrapping any air?
4. Lid pressed firmly down on the bowl?
5. Bowl with lid in place removed from the bath and the bowl and lid carefully dried off?
6. Mass of the bowl, lid and sample determined? (E)
7. Temperature of the water in bowl measured and recorded?
8. Procedure repeated a second time (no need to wait 10 more minutes)?
9. If mas <mark>s v</mark> aries by more than 1.0 g, procedure repeated until two masses are within 1.0 g?
10. Theoretical maximum specific gravity calculated $\{A/(A+D-E)\}$?
<u>ASTM only: Wei<mark>g</mark>hing-in-air determination (flask):</u> 1. Flask slowly filled with water taking care not to introduce air into the sample?
1. Flask slowly filled with water taking care not to introduce air into the sample?
2. Flask and contents placed in water bath for 10 ±1 min?
3. Top of flask should not be submerged?
4. Temperature of the water in flask 25 \pm 1 $^{\circ}$ C (77.0 \pm 1.8 $^{\circ}$ F) (temperature recorded)?
5. Flask completely filled with water using a cover plate?
6. Care taken not to entrap air beneath cover plate?
7. Moisture wiped from the exterior of the flask and cover plate?
8. Mass of the flask, cover plate, and contents determined? (E)
9. Theoretical maximum specific gravity calculated $\{A/(A+D-E)\}$?
Supplemental Procedure for Mixtures Containing Porous Aggregate
Note: This procedure is only performed if the aggregates are not thoroughly sealed.
Note to Assessors: Typically laboratories should not be demonstrating this supplemental procedure. It is included for reference.
1. Water decanted from the container through towel [ASTM: through a 75μm (No. 200) sieve]?
2. Several large pieces of aggregate broken to examine for wetness?
3. If the aggregate has absorbed water, sample spread [ASTM only: on a flat tray] in front of a fan to
remove surface moisture and stirred periodically?
4. Sample weighed at 15 minute intervals until constant mass (less than 0.05%) is reached?
5. Final surface dry mass substituted into the equation for the mass of the dry sample in air?
COMMENTS (T209 / D2041): (T209 / D204

Revised 2011-03-25

									(D69	26 / D	692
		<u>APPARAT</u>	<u>US</u>					Date	e:		
Speci	men Mol	d Assemblies:									
			1	2	3	4	5	6	7	8	9
Inside	diameter	of mold: 3.995 – 4.005 in.?									
Colla	r and base	e plate fit mold?									
Specia (a) (b)		actor eter of disk not less than 100 mm (3.95 in.) ness at least 12.5 mm (1/2 in.)?									
Comp	action Ha	<u>ammer</u>									
(a)	Make	r: Serial No. (or I.D. No.									
(b)	Monu	ol hommore [ASTM: Manual hamman and	maaha	nical	la cerea re		/ 10 010	untat	ina h	asa n	dag
(0)	Manual hammers [ASTM: Manual hammer and mechanical hammer w/ non-rotating base pedestal]: (1) Tamping face: flat, not slanted, 3 7/8 in. (98.4 mm) [ASTM: 98.0 – 100.0 mm] in diameter?										
		Tamping face: flat, not slanted, 3 7/8 in. (98.4 mm) [<i>ASTM</i> : 98.0 – 100.0 mm] in diameter? Sliding mass: 4527 to 4545 g (9.98 to 10.02 lb)?									
	(2)			?							
	(2)		0.02 lb) ⁴							•••••	•••••
	(3)	Sliding mass: 4527 to 4545 g (9.98 to 1 Note to Assessors: <i>Check records for hamn</i> Drop: 455.7 to 458.7 mm (17.94 to 18.0	0.02 lb); ers that 6 6 in.)? .	are di <u>f</u>	ficult	to asse	emble.				
		Sliding mass: 4527 to 4545 g (9.98 to 1 Note to Assessors: Check records for hamm Drop: 455.7 to 458.7 mm (17.94 to 18.0 AASHTO Mechanical compactor: calib	0.02 lb)sers that 6 in.)? .	are di <u>f</u> give	ficult i result	to asse ts com	emble. ipara	ble w	 ith a 1	 manud	 ally-
	(3)	Sliding mass: 4527 to 4545 g (9.98 to 1 Note to Assessors: <i>Check records for hamn</i> Drop: 455.7 to 458.7 mm (17.94 to 18.0	0.02 lb)sers that 6 in.)? .	are di <u>f</u> give	ficult i result	to asse ts com	emble. ipara	ble w	 ith a 1	 manud	 ally-
(a)	(3) (4)	Sliding mass: 4527 to 4545 g (9.98 to 1 Note to Assessors: Check records for hamm Drop: 455.7 to 458.7 mm (17.94 to 18.0 AASHTO Mechanical compactor: calib operated compactor (check records)?	0.02 lb) ^o ers that of the control o	are dif	ficult i	to asse	emble. ipara	ble w	 ith a 1	 manud	 ally-
(c)	(3) (4) <u>Mech</u>	Sliding mass: 4527 to 4545 g (9.98 to 1 Note to Assessors: Check records for hamm Drop: 455.7 to 458.7 mm (17.94 to 18.0 AASHTO Mechanical compactor: calib operated compactor (check records)?	0.02 lb) ers that to 6 in.)? . rated to	are dif	ficult i result	to asse	emble. upara 	ble w	ith a 1	nanud	ally-
(c)	(3) (4)	Sliding mass: 4527 to 4545 g (9.98 to 1 Note to Assessors: Check records for hamm Drop: 455.7 to 458.7 mm (17.94 to 18.0 AASHTO Mechanical compactor: calib operated compactor (check records)? anical hammer for use with rotating base Sliding mass: 4.54 ± 0.01 kg (10.00 ± 0.00)	0.02 lb) ers that 6 in.)? . rated to	are di <u>f</u> give	ficult i	to asse	emble. ipara	ble w	ith a 1	nanud	 ally-
(c)	(3) (4) <u>Mech</u> (1)	Sliding mass: 4527 to 4545 g (9.98 to 1 Note to Assessors: Check records for hamm Drop: 455.7 to 458.7 mm (17.94 to 18.0 AASHTO Mechanical compactor: calib operated compactor (check records)? anical hammer for use with rotating base Sliding mass: 4.54 ± 0.01 kg (10.00 ± 0.00 to Assessors: Check records for hamm	0.02 lb) of ers that to the following that to the following the following the following that the following the following that the following the following that the following that the following that the following that the following the following that the following that the following that the following the	are di <u>f</u> give ! [AS]	ficult in the second se	to asse	emble.	ble w	ith a 1	manud	ally-
(c)	(3) (4) <u>Mech</u>	Sliding mass: 4527 to 4545 g (9.98 to 1 Note to Assessors: Check records for hamm Drop: 455.7 to 458.7 mm (17.94 to 18.0 AASHTO Mechanical compactor: calib operated compactor (check records)? anical hammer for use with rotating base Sliding mass: 4.54 ± 0.01 kg (10.00 ± 0.00)	0.02 lb)? ers that to 6 in.)? . rated to	are di <u>f</u> give ! [AS]	ficult i result <u>FM or</u> icult to	to asse	emble	ble w	ith a 1	manue	ally-
(c)	(3) (4) (4) (1) (2)	Sliding mass: 4527 to 4545 g (9.98 to 1 Note to Assessors: Check records for hamm Drop: 455.7 to 458.7 mm (17.94 to 18.0 AASHTO Mechanical compactor: calib operated compactor (check records)? anical hammer for use with rotating base Sliding mass: 4.54 ± 0.01 kg (10.00 ± 0.00 to Assessors: Check records for hamm Drop: 457.2 ± 1.5 mm (18.00 ± 0.06 in. Hammer face: circular, slanted, & 99.5)	0.02 lb) ers that (6 in.)? . rated to	give give [JAS] re diff	ficult in result FM of the control	to asse	emble.	ble w	ith a 1	manud	ally-
(c)	(3) (4) Mech. (1) (2) (3)	Sliding mass: 4527 to 4545 g (9.98 to 1 Note to Assessors: Check records for hamm Drop: 455.7 to 458.7 mm (17.94 to 18.0 AASHTO Mechanical compactor: caliborated compactor (check records)? anical hammer for use with rotating base Sliding mass: 4.54 ± 0.01 kg (10.00 ± 0 Note to Assessors: Check records for hamm Drop: 457.2 ± 1.5 mm (18.00 ± 0.06 in.)	0.02 lb) ers that (6 in.)? . rated to	give give [JAS] re diff	ficult in result FM of the control	to asse	emble.	ble w	ith a 1	manud	ally-

4. Compaction Pedestal:

(a)	Manuai	com	paction	pedestai	<i>ASTM</i> :	<u>non-rotatin</u>	<u>g base[:</u>
	(1)	Woo	oden no	st: 203 3	x 203 2	x 457 2 mm	(8 x 8 x 1

en post: 203.3 x 203.2 x 457.2 mm (8 x 8 x 18 in.)?.... 1. Post plumb?....._____

2. Attached to solid concrete slab by four angle brackets?....._____

ASTM only: Average dry density of 670 – 770 kg/m³ (42 to 48 lb/ft²)
[oak, yellow pine, etc.]?

Steel cap, at least 304.8 x 304.8 x 25.4 cm (12 x 12 x 1 in.)?..... (2)

Steel cap level?..... Firmly attached to wooden post?....

Specimen holder mounted on pedestal and holds base plate, mold, and collar securely?...... (3)

1. Centers specimen mold over center of post?.....

COMMENTS (T245 / D6926 & D6927):

(T245 / D6926 & D6927)

MARSHALL APPARATUS FOR HMA RESISTANCE TO PLASTIC FLOW (T245) _____ (D6926 / D6927) _____

			APPARATUS (Continued) Date:
		(a)	Compaction pedestal (rotating base) [ASTM only]: (1) Wooden post plumb (vertical with respect to floor?)
			(2) Steel cap level and firmly attached to wooden post?
			(3) Mold holder mounted on pedestal and holds base plate, mold & collar securely?
			(4) Base rotation rate is 18 – 30 rpm?
5.		Breaking	
		(a)	Inside radius curvature in each segment is 2 in.?
		(b)	Ends of curvature lie in chordal plane 5/8 in. from center of curvature?
		(c)	$1/4 \times 1/4$ in. bevels on inside corners of each segment?
			Note to Assessors: The clear plastic template should fit inside the breaking head. The ends of the breaking head should lie inside the two lines.
		(d)	Two guide posts perpendicular to base with minimum diameter of 12.5 mm?
		(e)	Guide sleeves exhibit no appreciable play or friction?
6.		Loading	Device
0.		(a)	
		(b)	Maker: Serial No. (or I.D. No.) Produces uniform movement of 50 ± 5 mm/min $(2.00 \pm 0.15$ in./min)?
		(c)	Load measuring device:
			(1) Capacity: 22.2 kN (5000 lbf) [ASTM: 20 kN (5000 lb)]?
7.		Flow Me	easuring Devices:
		(a)	Guide sleeve and gauge (deformation indicator) graduated to 0.25 mm (0.01 in.) and operates with minimal friction?
	or	(b)	Micrometer dial graduated in increments of 0.25 mm (0.01 in.) or finer?
	or	(c)	Stress-strain recorder (LVDT) capable of indicating flow to 0.25 mm. (0.01 in.)?
8.		Orrana	nd/or Hot Distor presented (for heating aggregate caphelt, molds hommers, etc.)?
٥.			nd/or Hot Plates presented (for heating aggregate, asphalt, molds, hammers, etc.)?
		(a)	If a hot plate, is it provided with a suitable shield, baffle, or sand bath to minimize local overheating?
		(b)	if a not plate, is it provided with a suitable shield, barrie, of sand bath to minimize local overheating?
9.		Water B	ath_
		(a)	Depth at least 152.4 mm (6 in.)?
		(b)	Has perforated false bottom or a shelf at least 50.8 mm (2 in.) above bottom of bath?
		(c)	Bath thermostatically controlled to $60 \pm 1^{\circ}\text{C}$ ($140 \pm 2^{\circ}\text{F}$)?
10.		Mixing A	Apparaturs .
		(a)	Spoon, bowl, or pan for hand mixing or a mechanical mixer [AASHTO: recommended] which can be
		` /	maintained at the required mixing temperature and produce a homogenous mixture?
		(b)	Hot plate, infrared lamp, or any other device for maintaining mixing temperature?
			- · · · · · · · · · · · · · · · · · · ·

COMMENTS (T245 / D6926 & D6927):

(T245 / D6926 & D6927)

APPARATUS (Continued) Date: ermometers Thermometers for aggregates, bitumen, and mixes presented?
Thermometers for aggregates, bitumen, and mixes presented?
Note: Armored-glass, dial type, or digital thermometers with metal stems are recommended. (1) Range: 9.9 to 204°C (50 to 400°F) [ASTM: 50 to 400°F (10 to 200°C)]?
(1) Range: 9.9 to 204°C (50 to 400°F) [ASTM: 50 to 400°F (10 to 200°C)]?
(2) Sensitivity of 2.8°C (5°F) [ASTM: 3°C (5°F)]?
(2) Sensitivity of 2.8°C (5°F) [ASTM: 3°C (5°F)]?
(3) ASTM only: Thermometers calibrated? SN:
mi
Thermometer for water bath [ASTM only; standardized] (if not standardized also note under R18)?
(1) Readable to 0.2°C (0.4°F)?
ASTM only: Temperature measuring device, readable to 2°F (1°C), for checking mixing and compaction temperatures?*
lances
2 kg capacity, sensitive to 0.1 g, for weighing molded specimens?
5 kg capacity, sensitive to 1 g, for batching mixtures?
mperature-Viscosity Curve
Mixing and compaction temperatures based upon the temperature-viscosity curve for the asphalt used, AC or PG binder: 170 ± 20 cSt for mixing and 280 ± 30 cSt for compacting?
Note, for cutbacks: 170 ± 20 cSt for mixing and 280 ± 30 cSt (tested with 50% solvent) for compacting.
Mixing temperature range: to to to to to to
1

COMMENTS (T245 / D6926 & D6927):

(T245 / D6926 & D6927)

AASHTO Materials Reference Laboratory

(1245)	
(D6926)	

	<u>PROCEDURE</u>	Date:
Drenara	ation of Mixture	
1.	AASHTO only: Initial batch prepared for "buttering" the mixing bowl and stirrers and cleaned by scraping, not wiped with cloth or solvent?	
2.	AASHTO only: At least 3 specimens prepared for each combination of aggregates and	bitumen content?
3.	Amount of each aggregate size fraction required for each specimen weighed into a pan	
4.	Pan containing aggregate placed on a hot plate or in an oven and heated to a temperature mixing temperature by more than approx. 28°C (50°F)?	re not exceeding the
	Note, cutbacks: no more than 14°C (25°F).	
5.	AASHTO only: Aggregate dried to constant mass?	
6.	Hot aggregate placed in bowl, mixed with spoon for approx. 5 seconds & crater formed	d?
7.	Required amount of preheated bituminous material added to aggregate?	·······
	Note, cutbacks: weigh the bowl, mixing blade, aggregate, and asphalt to be mixed.	
8.	Temperature of the aggregate and bituminous material still within the established mixing	
9.	Aggregate and bituminous material rapidly mixed until thoroughly coated?	
10.	If hot plate used during mixing, wire mesh (or similar material) used to prevent direct of	
	hot plate and mixing bowl (to avoid localized overheating)?	······························
	Note, cutbacks: cure the mix in a ventilated oven maintained at 11.1°C (20°F) above compacts precalculated mass of 50% solvent loss is obtained. Weigh at 15-minute intervals at first, and than 10 minutes when the desired mass is being approached.	
<u>Conditi</u>	ioning of Mixture [ASTM only]	
Single l	batched samples:	
1.	Samples transferred to covered metal containers and placed into an oven maintained established compaction temperature for 1 – 2 hours?	l at 8 to 11°C above the
Multipl	le batched samples:	
1.	Entire batch placed on clean non-absorptive surface, mixed by hand to ensure unifo into appropriate sample size to yield a height of 63.5 ± 2.5 mm $(2.5 \pm 0.1$ in.)?	rmity and quartered
	into appropriate sample size to yield a height of 63.5 ± 2.5 mm $(2.5 \pm 0.1$ in.)?	
2.	Samples transferred to covered metal containers and placed into an oven maintained the established compaction temperature for $1-2$ hours?	
	Note, cutbacks: Cure in mixing bowl in a ventilated oven maintained at 11° C (20° F) above conuntil the precalculated mass of 50% solvent loss is obtained. Masses should be obtained in 15 -and then at intervals of less than 10 minutes when the desired mass is being approached.	npaction temperature
COMM	MENTS (T245 / D6926 & D6927):	(T245 / D6926 & D6927)

Revised 2011-03-25

(1245)	_
(D6926)	

	PROCEDURE (Continued) Date:
Compa	action of Specimens
1.	Specimen mold assembly and face of the compaction hammer clean?
2.	Mold assembly and hammer heated in boiling water, on a hot plate,
	or in an oven at 93.3 to 148.9°C (200 to 300°F)?
3.	Filter paper or paper toweling placed in bottom of mold?
4.	Entire batch of mixture placed in mold?
5.	Mixture spaded vigorously with heated spatula or trowel?
	(a) Spaded 15 times around perimeter?
	(b) Spaded 10 times over the interior?
6.	AASHTO only: Surface of mixture smoothed to slightly rounded shape?
7.	Temp. of the mixture immediately prior to compaction within the limits of established compacting temp.?
8.	Filter paper placed on mixture and mold assembly placed in mold holder on compaction pedestal?
9.	50 to 75 blows applied unless otherwise specified, with hammer held perpendicular to base of mold?
10.	ASTM only: No mechanical device used to restrict the handle of the manual hammer in vertical position?
11.	Mold and contents reversed?
12.	Same number of blows applied to reversed specimen?
<u>AASHT</u> 1.	<u>FO only, Removal from Mold</u> Sample extractor placed on end of specimen?
2.	Assembly, with collar up, placed in testing machine?
<i>3</i> .	Pressure applied to collar via load transfer bar?
4.	Specim <mark>en f</mark> orced into collar or otherwise extruded up into collar (no free-fall of specimen)?
<i>5</i> .	Collar <mark>lif</mark> ted from sp <mark>e</mark> cimen and specimen transferred to smooth, flat surface?
6.	Allowed to stand overnight at room temperature?
<i>7</i> .	Specimen mass determined and recorded?
8.	Specimen measured, height is 63.5 ± 1.27 mm $(2.5 \pm 0.05 \text{ in.})$?
	AASHTO Materials Reference Laboratory
	only, Removal from Mold
<i>1</i> .	Specimen allowed sufficient time to cool prior to extruding from mold?
	Note to assessors: Cooling specimens in mold may be facilitated by immersing in cold water. To facilitate extraction
•	and reduce specimen distortion, specimens in mold may be briefly immersed in a hot water bath prior to extruding.
2.	Specimen extruded using any suitable device provided the specimen is not distorted during the process?
<i>3</i> .	Specimen transferred to smooth, flat surface and allowed to cool (preferably overnight)?

COMMENTS (T245 / D6926 & D6927):

(T245 / D6926 & D6927)

(T245)_	
(D6927)	

		PROCEDURE (Continued)	Date:
Testing	g of specimens:		
1.		of three replicate specimens tested?	
2.	ASTM only: Bulk speci	fic gravity of each specimen determined by D2726,	D1188, or D6752?
3.		thickness measured according to D3549 (four measured)	
4.		st temperature by immersing in water bath for	,
		placing in an oven for 2 hours?	
5.	Bath or oven maintained	1 at 60 ± 1 °C (140.0 ± 1.8 °F) for asphalt?	
		ns placed in air bath maintained at 25 ± 1 °C (77.0 ± 2°F).	
6.		urfaces of breaking head cleaned?	
7.	Guide rods lubricated?	-	
8.	Temperature of breaking	g head maintained at 21.1 to 37.8°C (70 to 100°F)?	
9.	Specimen removed from bath or oven and placed in lower segment of breaking head?		
10.	ASTM only: Excess water removed with a towel?		
11.	Breaking head and specimen positioned on testing machine?		
12.	Flow meter (if used) pla	ced over guide rod and adjusted to zero?	
13.	Load applied to specimen until maximum load is reached?		
14.	Maximum load applied	within 30 seconds after removal of specimen from be	ath or oven?
15.		value recorded the instant the load begins to decrea	
16.	Load correction:		
	(a) For core specin	nens: load corrected when thickness of specimen is a	not 63.5 mm (2 1/2 in.)
	by multiplying	by factor from table 1?	······
	(b) For <u>lab-molded</u>	specimens: shall conform to thickness requirement	of 63.5 mm (2 1/2 in.)
	by multiplying	by factor from table 1?	
COM	JENTS (T2/15 / D6026 & 1	D6027)·	(T245 / D6026 & D6027)

(T245 / D6926 & D6927)

AASHTO Materials Reference Laboratory

HVEEM APPARATUS FOR RESISTANCE TO DEFORMATION AND COHESION OF BITUMINOUS MIXTURES

(T246)	_
(D1560)	

			<u>APPARATUS</u>	Date:
1.		Specin	nen Preparation Apparatus	
1.		(a)	AASHTO: California Kneading Compactor (from T247) available	??
		()	ASTM: Any compaction device is acceptable	
	or	(b)	Other kneading compactor that will show stabilometer values equi	
			California kneading compactor (the supplier is responsible for furr	
		Note: a	other compactors that show a similar calibration curve to a California Knead	ding Compactor are acceptable.
2.		Resista	ance to Deformation Test Apparatus	
		(a)	Stabilometer in working condition?	
		(b)	Rubber bulb for removing or adding air into stabilometer (during a	adjustment of stabilometer)?
		(c)	Compression testing machine: minimum capacity of 44.5 kN (10,0	000 lbf)?
			Capable of speed of 1.3 mm/min (0.05 in/min)?	
		(d)	Push-out device?	
		(e)	Oven capable of being maintained at 60 ± 3 °C $(140 \pm 5$ °F)?	······
		(f)	Calibration cylinder, hollow metal cylinder:	0.051)1113577 5 . 01 10
			(1) Height: 140 mm (5 1/2 in.) [ASTM: 140 ± 6.4 mm (5.5 \pm	0.25 in.)] [AMRL: 5 to 8 in.]?
		(~)	(2) Outside diameter: 101.47 to 101.73 mm (3.995 to 4.005 in	n.)?
		(g)	Solid-wall metal follower?	5 to 9 :m 19
			(1) Height: 133.35 to 146.05 mm (5.25 to 5.75 in.) [AMRL: Diameter: 101.092 to 101.346 mm (3.980 to 3.990 in.)?	5 to 8 m.j?
			(2) Diameter. 101.092 to 101.340 min (3.980 to 3.990 m.):	
3.		Cohesi	ion Test Apparatus (cohesion portion of the test not needed for "stab	ility only" accreditation)
		(a)	Cohesiometer?	
		(b)	2000 g of steel shot passing 2.00-mm (No. 10) sieve and retained of	on 1.40-mm (No. 14) sieve?
		(c)	Steel shot flows 1800 ± 20 g/min.?	<u></u>
			Note: Other materials may be used provided rate of loading is equivalent	t to that obtained when using steel shot.
		(d)	Oven maintained at $60 \pm 1^{\circ}$ C ($140 \pm 2^{\circ}$ F) [ASTM: $60 \pm 3^{\circ}$ C ($140 \pm 3^{\circ}$ C)	± 5°F)] ?
		(e)	Balance: 5 kg capacity, sensitive to 1 g [ASTM: 10 kg capacity, se	ensitive to 1 g?]?
0.0			T246 (D1560)	(D0.16 / D1.560)
CC	ИMM	ENTS (T246 / D1560):	(T246 / D1560)

HVEEM APPARATUS FOR RESISTANCE TO DEFORMATION AND COHESION OF BITUMINOUS MIXTURES

(T246)	
(D1560)	

		PRO	<u>CEDURE</u>	Date:
A dine	stment of Stabilometer			
1.		at distance from bottom of uppe	r tanered ring to top o	f base is 89 mm (3.5 in.)?
2.	Calibration cylinder	inserted into stabilometer?	t tapered ring to top o	
3.	ASTM only: Cyline	der preheated to 60°C (140°F))	
4.	A horizontal pressur	re of 34.5 kPa (5 psi) applied?		
5.		ontal pressure of 34.5kPa (5 ps		
				chine?
6.				
7.	Pump handle turned	until the stabilometer dial read	s 689 kPa (100 psi)?	
8.	Pump handle turned	at approx. two turns per second	d?	
9.	Turns indicator dial	reads 1.95 to 2.05 turns?		
10.	If not, is the air in the	ne cell adjusted and procedure re	epeated?	
11.	Horizontal pressure	released and calibration cylinder	er removed?	
. .				
	tance to Deformation	100 (4:): ::	(4) (2) (3) (5) (5)	\1: 10
1.	Test specimens are	102 mm (4 in.) in diameter and	$64\pm3 \text{ mm} (2.5\pm0.1 \text{ in.}$) high?
2		e not correct height or diameter th		
2.	Consider and Investment	ed and compacted in accordance	e with (124// D1361)?
3.	Specimen brought to	$60 \pm 3^{\circ}$ C (140 ± 5°F) [ASTM]	: in an oven jor 3 to 4	hours]?
1		to room temperature when desired		noisture is present. it device?
4. 5	Towned and of anoa	d from filloid to stabilometer by	means of the push-ou	it device?
5.	Fallower placed on	top of anoimon?		······
6. 7.	Harizantal praced off	of 24.5 laDo (5 pgi) applied?		·····
7. 8.	Vartical maxament	of proce boom?		
o. 9.				
9. 10.	If looking chime use	d an apharical hand afleading	daviaa shims ramaya	d prior to stabilometer test?
10. 11.	Stabilomator gauge	readings recorded at vertical la	device, similis remove	ence Laboratory
11.				00, 3000, 4000, 5000, and 6000 lbf)?
	AASIII O. 2.23, 4.4	and 26.7 kN (3000, 5000, and	5.7 KW (500, 1000, 200 6000 IbA?	
12.	Vertical movement	of press stopped at 26.7 kN (60)	0000 <i>10j):</i> 00 lbf) load?	
13.	Vertical load immed	liately reduced to 4.45 kN (100)	0 lbf) [ASTM: 4 to 4	9 kN (900 to 1100 lbf)]?
14.				
	Note: this will result i	n a further reduction of the vertica	l load and is normal.	
15.	Pump handle turned	until the stabilometer dial read	s 689 kPa (100 psi)?	<u> </u>
16.	Pump handle turned	at approx. two turns per second	d?	
17.	Number of turns rec	orded as the displacement read	ing? (D)	
18.	Stabilometer value of	calculated correctly?		
19.	If height of specime	n is not 64 ± 3 mm (2.5 ± 0.1) in	.), is stabilometer valu	ue corrected according chart?
	S =	22.2	Where:	S = stabilometer value
		D * D / (D . D) + 0.222		P_h = horizontal pressure (kPa)

 $P_h * D / (P_v - P_h) + 0.222$ P_v = vertical pressure (kPa)

D = displacement

COMMENTS (T246 / D1560):

(T246 / D1560)

HVEEM APPARATUS FOR RESISTANCE TO DEFORMATION AND COHESION OF BITUMINOUS MIXTURES

(T246)	
(D1560)	

		PROCED	URE (Continued)	Date:	
Cohe	sion (if demonstrated)	[AASHTO only: Cohesion testi	ng is optional and not r	equired to determine Stability]	
1.	Specimen in oven at $60 \pm 1^{\circ}$ C ($140 \pm 2^{\circ}$ F) for minimum of 2 hours?				
2.				! hours?	
3.				± 5°F)]?	
4.					
5.	Temperature allow	ed to recover before testing?		·····	
6.				······	
7.	Flow stopped at break or at 13 mm (1/2 in.) deflection?				
8.	Mass of shot used recorded?				
9.	Cohesiometer value calculated from formula in (T246 / D1560)?				
	C =	L	Where:	C = cohesiometer value	
				L = mass of shot	
		$W * (0.20H + 0.044H^2)$		W = diameter, cm (or in.) or width, cm (or in.)	
				D = height, cm (or in.)	

COMMENTS (T246 / D1560):

(T246 / D1560)

Note to Assessors: At all (except ~5) labs across the country you must write the "Cohesion not demonstrated" note. This is never a Nonconformity – it is always an ASTM only Observation.

AASHTO Materials Reference Laboratory

CALIFORNIA KNEEDING COMPACTOR FOR PREPARATION AND TESTING OF HMA SPECIMENS

(T247)	
(D1561)	

							AP	PARA	1103					Da	te:		
	Kneadii	ng Com	nacto	r													
	(a)				ng comp	actor?											
r		Other	knead	ling co	mpacto	r with	calibratio	on curv	ve sim	ilar to	Calif	ornia l	kneadi	ng co	mpact	or?	
	Maker:						·	•••••				S	erial N	lo. (or	I.D. N	No.)?	
	Accesso	ories															
	(a)		trou	gh?													
	(b)	Paddle	to fi	t cross	section	of trou	ıgh?										
	(c)	Paddle to fit cross section of trough?															
	(d)	Mold l	nolde	r?													
	(e)	Steel s	him a	pprox	imately	6.4 x 1	9 x 64 n	nm [AS]	STM: 6	6.4 x 1	9.1 x	63.5 n	nm				
		[AMR	[AMRL: piece of steel of any convenient shape, ½ in. thick (6.4 mm thick)]?														
	(f)	Round	-nose	steel 1	rod, 9.5	mm (3	/8 in.) ir	ı diame	eter by	406 ı	mm (1	6 in.)	long?				
	(g)	Heavy	pape	r disks	s, 102 m	ım [<i>AS</i>	TM: 101	1.6 mm	ı] dian	neter?							
	Molds																
	Wiolus								1	2	3	4	5	6	7	8	9
	Inside	dia : 10	1 47	101 ′	72	(a 00 =											
	misiac	ara ro	1.7/	- 101.	/3 mm ((3.995)	-4.005 i	in.)?									
	Heigh	t approx	. 127	mm (5	5 in.)?		- 4.005 l		ded)?								
	Heigh Does th	t approx le labora tus for A	tory applieressionetal He Ou	mm (5 nave the cation of n testin followeright: 1 itside d ight: 3	of Static ng machers: 40 mm liameter 8.1 mm	ble mo c "Leve nine: m (5 1/2 r; 101.0		" Load capaci "TM: 13	ty 222 3 9.7 m m (3.9	2 kN (: um] [A 80 to	50,000 MRL 3.990) lbf)? : min. in.) [/	5 in.,	no ma	nx.] hiş 2 mm]	gh?	ry
	Heigh Does th Appara (a)	tus for A Compi Two m (1) (2) aneous	tory tory tory tory tory tory tory tory	mm (5 nave the eation of n testin followeright: 1 ttside d ight: 3	of Static ng maclers: 40 mm liameter 8.1 mm	ble mo "Levenine: m (5 1/2 r: 101.0 (1 1/2 r: 101.0	lds (reco	TM: 13.	tty 222 39.7 m (3.9 m (3.9	2 kN (: mm] [A 80 to 80 to	50,000 MRL 3.990 3.990) lbf)? : min. in.) [/ in.) [/	5 in., 4STM	no ma: 101.	nx.] hig 2 mm]	gh? ?	ry
	Heigh Does th Appara (a) (b) Miscell (a)	t approx te labora tus for A Compi Two m (1) (2)	tory tory tory tory tory tory tory tory	mm (5 nave the cation of n testing followers ight: 1 straight: 3 straight: 3 etside of the cation of	of Stations machine in the station of Stations machines: 40 mm liameter 8.1 mm liameter eferably	ble mo "Levenine: m (5 1/2 r: 101.0 1(1 1/2 r: 101.0	lds (reco	TM: 13.31 mm	ty 222 39.7 m (3.9 m (3.9	2 kN (: am] [A 80 to 80 to	50,000 MRL 3.990 3.990) lbf)? : min. in.) [/ in.) [/	5 in., 4 STM 4 STM	no ma: 101:	mx.] hi 2 mm]	gh? ?	гy
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	Heigh Does th Appara (a) (b) Miscell (a) (b) (c)	tus for A Composition Two m (1) (2) aneous Therm Trowe Metal Balanc AASH	ttory Applications Applicati	mm (5 nave the cation of nave th	of Staticeng maclers: 40 mm liameter 8.1 mm liameter and secondary, secondary	ble mo c "Levenine: m (5 1/2 r: 101.0 (1 1/2 r: 101.0 / armoroops? ensitive	lds (reco	TM: 1.31 mm	tty 222 39.7 mm (3.9 mm (3.9)?	2 kN (: nm] [A 80 to 80 to	5%,000 MRL 3.990 3.990) lbf)? : min. in.) [/ in.) [/	5 in., 4STM 4STM	no ma : 101. : 101. : 101.	2 mm] 2 mm]	gh? ? ?	ry
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COMMENTS (T247 / D1561):

(T247 / D1561)

CALIFORNIA KNEEDING COMPACTOR FOR PREPARATION AND TESTING OF HMA SPECIMENS

(T247)	
(D1561)	

ROCEDURE	Date:
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Sample Preparation:

- 4. Aggregate recombined to 1200 g of specified grading?

5. Asphalt and aggregate at correct temperature when mixing begins (see table below)?

Mixing Temperature Table

Temperature Range, °C (°F)

Asphalt Grade	AASHTO min.	ASTM min.	maximum
AC-2.5, AR 1000, or 200-300 Pen	107 (225)	99 (210)	121 (250)
AC-5, AR 2000, or 120-150 Pen	121 (250)	110 (230)	135 (275)
AC-10, AR 4000, or 85-100 Pen	135 (275)	121 (250)	149 (300)
AC-20, AR 8000, or 60-70 Pen	149 (300)	132 (270)	163 (325)
AC-40, AR 16000, or 40-50 Pen	149 (300)	132 (270)	163 (325)

- 6. Asphalt and aggregate rapidly and thoroughly mixed?
- 7. ASTM only: Mix transferred to flat pan and cured for 2 to 3 h at $146 \pm 3^{\circ}C$ (295 $\pm 5^{\circ}F$) or for 15 to 18 h at $60 \pm 3^{\circ}C$ (140 $\pm 5^{\circ}F$) in an oven equipped with air circulation?.....
- 8. Mixture and molds brought to correct temperature [110°C (230°F) for paving grade asphalt]?.....

Compaction

- 3. Mass of mixture for one specimen placed in preheated trough? ______

- 7. Rest of mixture placed in mold and rodding repeated?8. Compactor foot heated?
- 9. Mold holder and mold placed in compactor?
- 11. Shim removed and mold tightening screw released?

- 19. AASHTO only: If specimens to be tested according to (T246 / D1560), testing completed within 3 hours of returning specimens to the oven?......

COMMENTS (T247 / D1561):

(T247 / D1561)

COMMENTS (T269 / D3203):

		PERCENT AIR VOIDS IN COMPACTED BITUMINOUS PAVING MIXTURES	(T269) (D3203)
		APPARATUS Date:	
1.	or or	Equipment for one of the following: (a) Method T166 / D2726 (Bulk Specific Gravity)? (b) Method T275 / D1188 (Bulk Sp. G by Paraffin Coating)? (c) Method T331 / D6752 (Bulk Sp. G by Vacuum Sealing)?	
2.	or	Equipment for one of the following: (a) Method T209 / D2041 (Maximum Specific Gravity by Rice Method)? (b) ASTM only: Method D6857 (Max Sp. G by Vacuum Sealing)? Note to Assessors: AMRL currently does not assess for D6857.	
		<u>PROCEDURE</u>	
For 1. 2.	r Der	Bituminous Paving Mixtures Bulk specific gravity determined by (T166 / D2726), (T275 / D1188), or (T331 / D6752)? Theoretical maximum specific gravity determined by (T209 / D2041) [ASTM only: or D6857]?	
1. 2. 3.	r Ope	Density of bituminous mixtures (10% air voids or higher) Density of bituminous mixture determined from its dry mass and its volume? Height of specimen determined? Volume of specimen determined based on average height and diameter measurement?	
4. 5.		Density converted to bulk specific gravity?	
<u>Ca</u>	lcula	AASHTO Materials Reference Laborat tions Percent air voids calculated in accordance with method?	-
1,		Percent Air Voids = 100 * [1 - (bulk sp gr / theoretical max sp gr)]	······································
No	te to A	Assessors, alternative terminology:	
110		Gmm = maximum specific gravity (Rice Test, T209/D2041) Gmb = bulk specific gravity (Bulk Sp G Test, T166/D2726) Percent Air Voids = 100 * [1 - (Gmb / Gmm)] Percent Air Voids = 100 * (Max – Bulk) / Max	

(T269 / D3203)

BULK SPECIFIC GRAVITY OF HMA USING PARAFFIN-COATED OR PARAFILM-COATED SPECIMENS

(T275)	_
(D1188)	

	<u>APPARATUS</u>	Date:
4 40117		
<u>AASH1</u> 1.	<u>O METHOD A</u> Balance (M231), can determine constant mass of specimen to 0.1 percent?	
2.	Bath for immersed weighing with overflow outlet, thermostatically controlled to 25.0 \pm 0	
<i>3</i> .	Suitable suspension and holder for immersed weighing:	.5 ((/ / .0 ± 0.5 1):
٥.	(a) Suspended from center of pan?	
	(b) Holder and specimen completely immersed?	
	(c) Wire suspending holder of smallest practical size?	
4.	Paraffin (Specific Gravity known)?	
5.	Room temperature: $25 \pm 5 \%$ (77 $\pm 9 \%$)?	
AASHT	TO METHOD B	
<i>1</i> .	Balance (M231), can determine constant mass of specimen to 0.1 percent?	
2.	Constant temperature water bath:	
	(a) Thermostatically controlled to $25.0 \pm 0.5 \%$ (77.0 $\pm 0.9 \%$)?	
	(b) ASTM 17C or 17F thermometer?	
<i>3</i> .	Calibrated volumeter with a tapered lid and capillary bore?	
<i>4</i> .	Drying oven at $52 \pm 3 $ $^{\circ}$ C $(125 \pm 5 $ $^{\circ}$ F)?	
5.	Paraffin (Specific Gravity known)?	
6.	Room temperature: $25 \pm 5 \%$ (77 $\pm 9 \%$)?	
	METHOD AND REL	
1.	Balance (D4753) with ample capacity and with sufficient sensitivity to determine	CD29
2.	bulk specific gravity to four significant figures? (i.e. 0.1 g for 100.1 to 999.9 g) such as Bath for immersed weighing:	
2.	Bath for immersed weighing: (a) Constant level overflow?	boratory
or		······································
<i>3.</i>	Suspension and holder for completely immersed weighing?	
4.	Parafilm (Specific Gravity known)?	
<i>5</i> .	Polyurethane foam:	
	(a) Mat with a minimum 50 x 50 cm (20 x 20 in) for working surface by 12.5 mm	
	(b) At least one mat with a size approx. equal to the top surface dimensions of spe	ecimen on hand?
6.	Calibration cylinder:	
	(a) Smooth, sided aluminum cylinder?	
_	(b) Approximately 100-mm (4-in) diameter by 60-mm (2.5 in)?	
<i>7</i> .	Sharp knife to cut parafilm?	······································

COMMENTS (T275 / D1188):

(T275 / D1188)

BULK SPECIFIC GRAVITY OF HMA USING PARAFFIN-COATED SPECIMENS (T275)

	AASHTO PROCEDURE Date:
<u>Specim</u>	<u>ens</u>
1.	Recommended size:
	(a) Diameter (or side of sawed specimens) at least 4 times maximum size of aggregate?
	(b) Thickness at least 1 1/2 times maximum size of the aggregate?
<i>2</i> .	Drying to constant mass:
	(a) Distortion, bending, or cracking avoided and free of foreign material?
	(b) Overnight at $52 \pm 3 \%$ ($125 \pm 5 \%$)?
	(c) Additional 2 hr. drying intervals?
	(d) Constant mass (change less than 0.05 percent)?
METHO	
<u>метне</u> 1.	<u>Mass in air determined (dry; see 1 (b) above)? (A)</u>
1. 2.	Allowed to cool in air at room temperature at $25 \pm 5 $ °C (77 $\pm 9 $ °F) for 30 minutes?
2. 3.	Coated with paraffin, filling all voids?
3.	Note: The specimen may optionally be coated with powdered talc before coating with paraffin to facilitate removal.
	Note: The specimen may optionally be cooled to 40 F before coating.
4.	Cooled 30 min., then weighed in air? (D)
5.	Immersed in water at 25 \pm 1 °C (77 \pm 2 °F) and weighed? (E)
6.	Sp. Gr. of paraffin determined (if unknown)? (F)
7.	Bulk Specific Gravity calculated as follows?
	$\stackrel{1}{\sim}$ $\stackrel{2}{\sim}$ $\stackrel{3}{\sim}$ $\stackrel{4}{\sim}$ $\stackrel{1}{\sim}$ $\stackrel{4}{\sim}$ $\stackrel{4}$
	Bulk Specific Gravity =
	D-E-((D-A)/F)
METH	
1.	Mass of dry sample in air determined? (A)
<i>2</i> .	Room temperature $25 \pm 5 ^{\circ}$ C $(77 \pm 9 ^{\circ}$ F)?
<i>3</i> .	Coated with paraffin, filling all voids, and then cooled 30 min.?
4.	Mass of specimen + paraffin determined?
<i>5</i> .	Outside of volumeter wiped dry?
6.	Mass of volumeter + water at $25 \pm 1 \%$ (77 $\pm 2 \%$) determined?
<i>7</i> .	Mass of volumeter + water + specimen at $25 \pm 1 ^{\circ}\text{C}$ (77 $\pm 2 ^{\circ}\text{F}$) determined?
8.	Determine specific gravity of Paraffin (if unknown)?
9.	Bulk Specific Gravity calculated as follows?
	A
	Bulk Specific Gravity =
	D - [E - C + ((C-A)/F)]

COMMENTS (T275): (T275)

COMMENTS (D1188):

BULK SPECIFIC GRAVITY OF HMA USING PARAFILM-COATED SPECIMENS (D1188)

		ASTM PROCEDURE	Date:
Speci	mens		
<u> 1.</u>	Diameter (or side of sawed specimens)	at least 4 times maximum size of ag	gregate?
2.	Thickness at least 1 1/2 times maximum		
3.	Drying to constant mass:	sige of the ugging that	
•		ng of specimen avoided?	<u> </u>
	(c) Dried under fan until constan	t mass achieved?	
<u>Proce</u>	<u>edure</u>		
1.	Mass in air determined (dried under a	fan until constant mass has been ac	hieved)? (A)
<i>2</i> .	On a hard surface, sharp blade used to	cut two 100 x 100 mm and one 100	x 200 mm pieces of parafilm?
<i>3</i> .	Backing taken off one of the 100 x 100		
4.	Opposite sides of film grasped and stre		
	(b) Stretched to an approximately	150 x 150-mm square?	
<i>5</i> .	Stretched film placed over one end of s	pecimen and sides of film pressed a	round sample?
6.	Specimen turned over, placed on foam	mat, and Steps 4 - 6 repeated?	
7.	Another piece of foam placed on top of	wrapped specimen?	<u> </u>
8.	Wrapped specimen pressed with another	er specimen of same size to eliminat	e air pockets from surfaces?
9.	Sharp knife used to trim excess film fro	om sides of sample?	
			<u> </u>
			each side of specimen?
10.	Backing peeled off remaining piece of	film and ends stretched to 400 mm ((16 in.)?
11.	One end of stretched film placed on sid	le of specimen and rolled over so file	m stretched tightly over surface?.
<i>12</i> .	Edges folded and pressed over edges of		
<i>13</i> .	Mass of covered specimen in air determ		
14.	Mass of covered specimen in water bat	h at 25 + 1 °C (77 + 1.8 °F)? (F)	
	(a) Is correction made if temperat	ture of water differs from 25 + 190	
	(b) If water differs by more than 2	ore of water at jets from 25 ± 1°C (to 15 minutes?
	(b) If water adjets by more than 2	C (3.0 1), specimen immersed 10	10 15 minutes:
<u>Calc</u> u	ulation <u>s</u>		
1.	Specific gravity of film determined by p	procedure in Sec. 8.3? (F)	
<i>2</i> .	If specimen contains moisture, correct		
<i>3</i> .	Bulk Specific Gravity calculated as followed		
	1 3	A	
	Bulk Specific Gravity =		
	z specific Gravity	D-E-((D-A)/F)	
		= = ((= 11), 1)	

Revised 2011-03-25

(D1188)

RESISTANCE OF COMPACTED BITUMINOUS MIXTURES TO MOISTURE INDUCED DAMAGE (TSR / Lottman)

(T283)	
(D4867)	

		APPARATUS Date:	
1.		Equipment for one of the following:	
1.		(a) Method T245/D6926 (Marshall), T247/D1561 (CA kneading compactor), T312/D4013	
		(Superpave Gyratory), or D3387 (US Corp of Engineers Gyratory Testing Machine)?	
0	r	(b) ASTM only: D1074 (Compressive strength) or D3496 (Dynamic Modulus)?	
2.		Vacuum Apparatus	
		(a) Vacuum container [ASTM only: preferably vacuum bowl from Rice test D2041]?	
		(b) Vacuum pump or water aspirator, conforms to (T209 / D2041)?	
		(c) Includes a manometer or vacuum gauge?	
3.		Balance, conforming to (T166 / D2726), reads to 0.1% of sample mass, G2/GP2?	
1.		Water bath(s):	
		(a) Conforming to (T166 / D2726), has balance suspension apparatus, etc?	
		(b) Capable of maintaining a temperature of 140.0 ± 1.8 °F (60 ± 1 °C) [ASTM: for 24 h]?	
		(c) Capable of maintaining a temperature of 77.0 \pm 1°F (25.0 \pm 0.5°C) [ASTM: \pm 1.8 °F (1 °C)]?	
5.		Freezer, maintained at 0 ± 5 °F (-18 ± 3 °C) [ASTM only: optional]?	
5.		AASHTO only: Oven, forced air draft, capable of maintaining any desired temperature setting from room	
		temperature to 176° C (350 °F) within $\pm 3^\circ$ C ($\pm 5^\circ$ F)?	
7.		Testing Apparatus	
		(a) Loading jack and ring dynamometer, conforms to (T245 / D6926)?	
0	r	(b) AASHTO: Mechanical or hydraulic testing machine (conforms to AASHTO T167) that provides a range of rates including 2 in. (50 mm) per minute?	
		ASTM: Mechanical or hydraulic testing machine capable of maintaining the required	
		strain rate and measuring load with equal or better precision?	
3.		Loading Strips [ASTM only: conforming to D4123 - Indirect Tension Test for Resilient Modulus]	
		(a) Concave surface with a radius of curvature equal to the nominal radius of the test specimen?	
		(b) The width is:	
		(1) 0.5 in. for a 4 in. diameter specimen?	
		(c) The length exceeds the thickness of the specimens?	
		(d) The edges are rounded by grinding?	
9.		Miscellaneous:	
•		(a) Plastic film or heavy-duty plastic bags, masking tape, 10 mL graduated cylinder?	
		(b) AASHTO only: Pan, bottom surface area of 75 - 200 sq. in. and depth of approximately 1 in.?	
		Note to Assessors – some helpful facts	

1 atmosphere (sea level) = 14.7 psi = 760 mm Hg (or torr) = 30 in Hg = 101.3 kPa

An <u>absolute</u> pressure gauge reads the pressure difference between a complete vacuum (0 pressure) and the sample. Examples include mercury manometers and some electronic gauges. Absolute gauges read atmosphere pressure (see above) when the vacuum system is off.

A <u>relative</u> pressure gauge reads the pressure difference between the normal atmosphere and the sample. Examples include most "vacuum gauges close to the source." Relative gauges read 0 when the vacuum system is off.

COMMENTS (T283 / D4867):

(T283 / D4867)

COMMENTS (T283):

RESISTANCE OF COMPACTED BITUMINOUS MIXTURES TO MOISTURE INDUCED DAMAGE (TSR / Lottman)

(T283)

		AASHTO PROCEDURE Date:
Sa	mnla	Preparation (laboratory mixed and compacted)
<u>sa</u>	пріє	Specimen size:
1.		(a) 4 in. diameter and 2.5 in. thick specimens used?
	or	(b) 6 in. diameter and 3.55 – 3.95 in. (90 – 100 mm) thick?
	01	Note: 6 in. specimens should be used if aggregate larger than 1 in. is presented.
2.		After mixing:
		(a) Mixture placed in a pan and cooled at room temperature for 2.0 ± 0.5 hours?
		(b) Mixture placed in a 140°F (60°C) oven for 16 ± 1 hour for curing?
		(c) Placed on spacers if shelf is not perforated?
3.		After curing:
		(a) Mixture placed in an oven at compaction temperature, $\pm 3^{\circ}C$ ($\pm 5^{\circ}F$), for
		2 hours \pm 10 min. prior to compaction?
		(b) Mixture compacted to 7.0 ± 0.5 percent air voids, or a void level expected in the field?
4.		After extraction from molds, test specimens are stored for 24 ± 3 hour at room temperature?
	mple	<u>Preparation</u> (field mixed and laboratory compacted)
1.		Specimen size:
		(a) 4 in. diameter and 2.5 in. thick specimens used?
	or	(b) 6 in. diameter and 3.55 – 3.95 in. (90 – 100 mm) thick?
_		Note: 6 in. specimens should be used if aggregate larger than 1 in. is presented.
2.		Field-mixed asphalt mixtures sampled in accordance with ASTM D979?
3.		No loose mix curing shall be performed?
4.		After sampling, mixture placed in oven until it reaches compaction temperature to within \pm 3°C (\pm 5°F)?
5.		After extraction from molds, test specimens are stored for 24 ± 3 hour at room temperature?
Sa	mnle	preparation (core test specimens)
1.	p.c	At least 6 cores for each set of mix conditions?
2.		
		Separate core layers as necessary by sawing or other suitable means, and layers to be stored at room temperature?
	aluat	ion of test specimens and grouping
1.		Theoretical maximum specific gravity of mixture determined by AASHTO T209?
2.		Specimen thickness determined by ASTM D3549 (average four thickness measurements at quarter points)?
3.		Bulk specific gravity determined by AASHTO T166?
4.		Volume of specimens expressed in cubic centimeters?
5.		Air voids calculated by AASHTO T269?
6.		Specimens sorted into two equal subsets of at least three specimens each so that average air voids of the
		two subsets are approximately equal?
	<u>y sub</u>	set - Preconditioning of test specimens
1.		Specimens stored at room temperature for 24 ± 3 hours?
2.		Specimens wrapped with plastic or placed in a heavy duty leak proof plastic bag?
3.		Specimens placed in a $77 \pm 1^{\circ}F$ ($25 \pm 0.5^{\circ}C$) water bath for at least 2 hours ± 10 min. and then tested?
4.		At least 1 in. of water above surface?

(T283)

RESISTANCE OF COMPACTED BITUMINOUS MIXTURES TO MOISTURE INDUCED DAMAGE (TSR / Lottman)

(T283)
(1200)

AASHTO PROCEDURE (C	Continued)
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	AASHTO PROCI	EDURE (Continued	<u>1)</u>	Date:	
	ioned subset – Vacuum saturation procedure				
	Specimens placed in the vacuum container supported				
	Container filled with potable water at room temperat				
	A partial vacuum <u>10-26 inches Hg</u> partial pressure (2				
	Note to Assessors: This is not the same amount of vacuum				
	27.5 mm Hg. A typical manometer that reads in mm Hg wi	ll not even move und	er the co	orrect vacuum for	T283.
		l man			-
	Correct TSR Pressure Absolute gauge (if vacuum is off reads 1 atm)	TSR pressure in.	TSR pi	ressure mm Hg	
	Relative gauge (if vacuum is off reads 1 aun)	10 - 26 in. $\sim 20 - 4 \text{ in.}$		254 – 660 mm Hg - 500 – 100 mm Hg	
	Relative gauge (ii vacuum is on reaus 0)	20 - 4 III.		- 300 – 100 mm 11g	_
	Vacuum applied for a short time, approximately 5-10	min as needed to a	chieve	correct saturation	on?
	Note to Assessors: The vacuum pressure and time is muc				
	Vacuum removed and specimens left submerged for				
	Mass of the SSD specimen after partial vacuum satur				
	SSD mass of conditioned samples compared with air				
	Degree of saturation determined by comparing volum				
	(a) If the volume of water is less than 70 percer				
	vacuum and/or more time?				
	(b) If the volume of water is more than 80 percentage (b)	ent, is the specimen	discar	ded?	
		· ·, · · · · · · · · · · · ·			
		percent, is the test	contini	ueu:	
	(c) If the volume of water is between 70 and 80	percent, is the test	contin	ueu?	
nditi	(c) If the volume of water is between 70 and 80	percent, is the test	contin	ued?	
nditi	(c) If the volume of water is between 70 and 80 ioned subset – Temperature conditioning procedure				
<u>nditi</u>	(c) If the volume of water is between 70 and 80 ioned subset – Temperature conditioning procedure Vacuum saturated specimens covered tightly with a p	plastic film and eac	h speci	men placed in a	plastic bag
<u>nditi</u>	(c) If the volume of water is between 70 and 80 ioned subset – Temperature conditioning procedure Vacuum saturated specimens covered tightly with a proceduring 10 ± 0.5 mL of water and placed in a free	plastic film and eac zer at $0 \pm 5^{\circ}$ F (-18	h speci ± 3°C)	men placed in a for a minimum of	plastic bag
<u>nditi</u>	ioned subset – Temperature conditioning procedure Vacuum saturated specimens covered tightly with a proceduring 10 ± 0.5 mL of water and placed in a free Specimens placed into a $140 \pm 2^{\circ}F$ ($60 \pm 1^{\circ}C$) water	plastic film and eac zer at $0 \pm 5^{\circ}$ F (-18 bath for 24 ± 1 hou	h specint 3°C) urs with	men placed in a for a minimum o a minimum	plastic bag of 16 hours?
<u>nditi</u>	ioned subset – Temperature conditioning procedure Vacuum saturated specimens covered tightly with a proceduring 10 ± 0.5 mL of water and placed in a free Specimens placed into a $140 \pm 2^{\circ}F$ ($60 \pm 1^{\circ}C$) water of 1 in. of water above specimen?	plastic film and eac zer at $0 \pm 5^{\circ}$ F (-18 bath for 24 \pm 1 hou	h specint 3°C) ars with	men placed in a for a minimum o a minimum	plastic bag of 16 hours?
<u>nditi</u>	ioned subset – Temperature conditioning procedure Vacuum saturated specimens covered tightly with a proceduring 10 ± 0.5 mL of water and placed in a free Specimens placed into a $140 \pm 2^{\circ}F$ ($60 \pm 1^{\circ}C$) water of 1 in. of water above specimen? Plastic bag and film removed from the specimens as	plastic film and eac zer at $0 \pm 5^{\circ}$ F (-18 bath for 24 ± 1 hou soon as possible af	h specing 3°C) ars with ter place	men placed in a for a minimum of a minimum of the ment in the wa	plastic bag of 16 hours?
<u>nditi</u>	ioned subset – Temperature conditioning procedure Vacuum saturated specimens covered tightly with a proceduring 10 ± 0.5 mL of water and placed in a free Specimens placed into a $140 \pm 2^{\circ}F$ ($60 \pm 1^{\circ}C$) water of 1 in. of water above specimen? Plastic bag and film removed from the specimens as	plastic film and eac zer at $0 \pm 5^{\circ}$ F (-18 bath for 24 ± 1 hou soon as possible af	h specing 3°C) ars with ter place	men placed in a for a minimum of a minimum of the ment in the wa	plastic bag of 16 hours?
<u>nditi</u>	ioned subset – Temperature conditioning procedure Vacuum saturated specimens covered tightly with a proceduring 10 ± 0.5 mL of water and placed in a free specimens placed into a $140 \pm 2^{\circ}F$ ($60 \pm 1^{\circ}C$) water of 1 in. of water above specimen? Plastic bag and film removed from the specimens as After 24 hours in the water bath, the specimens removed at $77 \pm 1^{\circ}F$ ($25.0 \pm 0.5^{\circ}C$), for 2 hours ± 10 min.?	plastic film and eac zer at $0 \pm 5^{\circ}$ F (-18) bath for 24 ± 1 hou soon as possible af eved and placed in a	h specing 3°C) ars with ter place	men placed in a for a minimum of a minimum ement in the wa bath, already	plastic bag of 16 hours? ter bath?
<u>aditi</u>	ioned subset – Temperature conditioning procedure Vacuum saturated specimens covered tightly with a proceduring 10 ± 0.5 mL of water and placed in a free Specimens placed into a 140 ± 2°F (60 ± 1°C) water of 1 in. of water above specimen? Plastic bag and film removed from the specimens as After 24 hours in the water bath, the specimens removed at 77 ± 1°F (25.0 ± 0.5°C), for 2 hours ± 10 min.? (a) If necessary, ice used to prevent water temp	plastic film and eac zer at $0 \pm 5^{\circ}$ F (-18) bath for 24 ± 1 hou soon as possible af wed and placed in a	h specing ± 3°C) arrs with ter place a water above	men placed in a for a minimum of a minimum ement in the wa bath, already	plastic bag of 16 hours?
<u>nditi</u>	ioned subset – Temperature conditioning procedure Vacuum saturated specimens covered tightly with a proceduring 10 ± 0.5 mL of water and placed in a free specimens placed into a $140 \pm 2^{\circ}F$ ($60 \pm 1^{\circ}C$) water of 1 in. of water above specimen? Plastic bag and film removed from the specimens as After 24 hours in the water bath, the specimens removed at $77 \pm 1^{\circ}F$ ($25.0 \pm 0.5^{\circ}C$), for 2 hours ± 10 min.?	plastic film and eac zer at $0 \pm 5^{\circ}$ F (-18) bath for 24 ± 1 hou soon as possible af wed and placed in a	h specing ± 3°C) arrs with ter place a water above	men placed in a for a minimum of a minimum ement in the wa bath, already	plastic bag of 16 hours?
	ioned subset – Temperature conditioning procedure Vacuum saturated specimens covered tightly with a proceduring 10 ± 0.5 mL of water and placed in a free Specimens placed into a 140 ± 2°F (60 ± 1°C) water of 1 in. of water above specimen? Plastic bag and film removed from the specimens as After 24 hours in the water bath, the specimens remo at 77 ± 1°F (25.0 ± 0.5°C), for 2 hours ± 10 min.? (a) If necessary, ice used to prevent water temp (b) The water bath should not require more than	plastic film and eac zer at $0 \pm 5^{\circ}$ F (-18) bath for 24 ± 1 hou soon as possible af wed and placed in a	h specing ± 3°C) arrs with ter place a water above	men placed in a for a minimum of a minimum ement in the wa bath, already	plastic bag of 16 hours?
	ioned subset – Temperature conditioning procedure Vacuum saturated specimens covered tightly with a proceduring 10 ± 0.5 mL of water and placed in a free Specimens placed into a 140 ± 2°F (60 ± 1°C) water of 1 in. of water above specimen? Plastic bag and film removed from the specimens as After 24 hours in the water bath, the specimens remo at 77 ± 1°F (25.0 ± 0.5°C), for 2 hours ± 10 min.? (a) If necessary, ice used to prevent water temp (b) The water bath should not require more than	plastic film and eac zer at $0 \pm 5^{\circ}$ F (-18) bath for 24 ± 1 hou soon as possible af eved and placed in a erature from rising	ter place the water above ch 77°F	men placed in a for a minimum of a minimum of a minimum of the walk bath, already of (25°C)?	plastic bag of 16 hours? ter bath?
	ioned subset – Temperature conditioning procedure Vacuum saturated specimens covered tightly with a proceduring 10 ± 0.5 mL of water and placed in a free Specimens placed into a 140 ± 2°F (60 ± 1°C) water of 1 in. of water above specimen? Plastic bag and film removed from the specimens as After 24 hours in the water bath, the specimens remo at 77 ± 1°F (25.0 ± 0.5°C), for 2 hours ± 10 min.? (a) If necessary, ice used to prevent water temp (b) The water bath should not require more than	plastic film and eac zer at $0 \pm 5^{\circ}$ F (-18 bath for 24 ± 1 hou soon as possible af wed and placed in a erature from rising in 15 minutes to reac pecimens determin	ter place above ch 77°F	men placed in a for a minimum of a minimum of a minimum of the walk bath, already of (25°C)?	plastic bag of 16 hours? ter bath?
	ioned subset – Temperature conditioning procedure Vacuum saturated specimens covered tightly with a proceduring 10 ± 0.5 mL of water and placed in a free specimens placed into a 140 ± 2°F (60 ± 1°C) water of 1 in. of water above specimen? Plastic bag and film removed from the specimens as After 24 hours in the water bath, the specimens remo at 77 ± 1°F (25.0 ± 0.5°C), for 2 hours ± 10 min.? (a) If necessary, ice used to prevent water temp (b) The water bath should not require more than the indirect tensile strength of dry and conditioned so The specimens in the 77°F water bath removed and prevent water tempore than the specimens in the 77°F water bath removed and prevent water tempore than the specimens in the 77°F water bath removed and prevent water tempore than the specimens in the 77°F water bath removed and prevent water tempore than the specimens in the 77°F water bath removed and prevent water tempore than the specimens in the 77°F water bath removed and prevent water tempore than the specimens in the 77°F water bath removed and prevent water tempore than the specimens in the 77°F water bath removed and prevent water tempore than the specimens water than t	polastic film and eac zer at $0 \pm 5^{\circ}$ F (-18 bath for 24 ± 1 hou soon as possible af eved and placed in a erature from rising in 15 minutes to reac pecimens determinal	th specifical has been a water above above at 77°F	men placed in a for a minimum of a minimum of a minimum of the walk bath, already of (25°C)?	plastic bag of 16 hours? ter bath?
	ioned subset – Temperature conditioning procedure Vacuum saturated specimens covered tightly with a proceduring 10 ± 0.5 mL of water and placed in a free Specimens placed into a 140 ± 2°F (60 ± 1°C) water of 1 in. of water above specimen? Plastic bag and film removed from the specimens as After 24 hours in the water bath, the specimens remo at 77 ± 1°F (25.0 ± 0.5°C), for 2 hours ± 10 min.? (a) If necessary, ice used to prevent water temp (b) The water bath should not require more than the indirect tensile strength of dry and conditioned so The specimens in the 77°F water bath removed and placed in the specimens in the bearing plates in the specimens in	blastic film and eac zer at $0 \pm 5^{\circ}$ F (-18) bath for 24 ± 1 hou soon as possible affived and placed in a erature from rising 15 minutes to reac pecimens determined blaced in the steel to the testing machine	h specing the specing with the specing water above the specing the	men placed in a for a minimum of a minimum o	plastic bag of 16 hours? ter bath?
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COMMENTS (T283): (T283)

EFFECT OF MOISTURE ON ASPHALT-CONCRETE PAVING MIXTURES ON TENSILE STRENGTH RATIO (TSR / Lottman)

(D4867)

		ASTM PROCEDURE Date:
Sa	mnle	Preparation (laboratory test specimens)
1.	<u>iipic</u>	Sample size:
		(a) 4 in. diameter and 2.5 in. thick specimens used?
	or	(b) Specimens of other dimensions used if aggregate larger than 1 in. is presented?
2.		Six specimens made for each test: three to be tested dry and three to be tested after partial
		saturation and moisture conditioning?
3.		Mixtures prepared in batches large enough to make at least 3 specimens or a batch large
		enough for just 1 specimen?
4.		Mixing temperatures and procedures followed for the method used?
5.		If an anti-stripping additive is used, are procedures in 6.4 and 6.5 used?
6.		After mixing:
		(a) Mixture placed in a closed container and placed in an oven for 1 to 2 hours to
		stabilize the specimen at the required compaction temperature?
		(b) If preparing multi-specimen batch, split into single specimens before placing into oven?
7.		After curing, mixture compacted to 7 ± 1 percent air voids, or a void level expected in the field?
8.		After compaction, test specimens are cooled as rapid as possible in a stream of moving air, extracted from
		molds, then procedure followed in Section 8 within 24 hours?
_		
	<u>mple</u>	Preparation (field specimens)
1.		Truck to be sampled selected in accordance with D3665 (Random Sampling of Construction Materials)?
2.		Sample taken from truck at plant in accordance with D979 (Sampling Bituminous Paving Mixtures)?
3.		Mixture temperature stabilized to approximately the temperature found in the field when
		rolling begins and temperature maintained in a closed container, in an oven for approximately
1		the time lapse between mixing and the start of actual rolling?
4. 5.		After curing, mixture compacted to 7 ± 1 percent air voids, or a void level expected in the field?
3.		After compaction, test specimens are cooled as rapid as possible in a stream of moving air, extracted from
(molds, then rest of procedure followed within 24 hours (saturation, testing, etc)?
6.		If specimens are not to be compacted in the field laboratory, place the samples in a sealed container, transported to the laboratory, and reheated to required temperature?
		container, transported to the laboratory, and reneated to required temperature?
Fv	กไมกร	ion of test specimens and grouping
1.		Theoretical maximum specific gravity of mixture determined by D2041?
2.		Specimen height determined by taking the average of four height measurements, ASTM D3549?
3.		Bulk specific gravity determined by ASTM D2726?
4.		Volume of specimens determined from Bulk specific gravity test (B - C expressed in cubic centimeters)?
5.		Air voids calculated D3203?
6.		Specimens sorted into two subsets of three specimens each so that average air voids of the

COMMENTS (D4867): (D4867)

two subsets are approximately equal?

8.

	EFFECT OF MOISTURE ON ASPHALT-CONCRETE PAVING MIXTURES (D4867) ON TENSILE STRENGTH RATIO (TSR / Lottman)
	ASTM PROCEDURE (Continued) Date:
Dr v sub	set - Preconditioning of test specimens
1.	Specimens stored at room temperature until test?
Conditi	oned subset – Preconditioning of test specimens
1.	Specimens placed in the vacuum chamber?
2.	Container filled with distilled water at room temperature?
	Note: The water used to saturate the specimens may be heated up to 140F (60 °C).
3.	A partial vacuum such as 20 in Hg applied for a short time (such as 5 minutes)?
	Note to Assessors: This is <u>not</u> the same amount of vacuum applied to Rice samples (T209 / D2041), which is 27.5 mm Hg.
4.	Volume of the partially saturated specimen determined in accordance D2726?
5.	Volume of the absorbed water determined by subtracting air dry mass of the specimen from the saturated surface-
J.	dry mass of the partially saturated specimen (Sec.8.6.2)?
6.	Degree of saturation determined by dividing the volume of the absorbed water by the volume of
	air voids and expressed as a percentage (Sec. 8.6.3)?
	(a) If the volume of water is less than 55 percent, is the procedure repeated using more vacuum?
	(b) If the volume of water is more than 80 percent, is the specimen discarded?
	(c) If the volume of water is between 55 and 80 percent, is the procedure continued?
7.	Specimens placed into a 140.0 ± 1.8 °F (60 ± 1 °C) water bath filled with distilled water for 24 hours?
8.	If a freeze-thaw cycle is desired, procedure in Note 6 used?
9.	After 24 hours in the water or air bath, specimens are removed and placed in a water bath already at
	77.0 ± 1.8°F (25 ± 1°C) for 1 hour?
10.	Height of the conditioned specimens determined by taking 4 measurements at quarter points (D3549)?
11.	Volume of conditioned subset specimens determined by D2726?
12.	Water absorption and degree of saturation determined in accordance with 8.6.2 and 8.6.3 (a degree of saturation
	exceeding 80% is acceptable after water bath soaking)?
13.	Swell calculated for partially saturated specimens (just after vacuuming procedure) and
	moisture-conditioned specimens (after additional water bath soaking time) (see Section 8.9.2)?
14.	Temperature of the dry subset adjusted by soaking in a water bath for 20 min. at 77.0 ± 1.8°F?
<u>Testing</u>	
1.	Height of dry specimens determined just before tensile testing by taking 4 measurements at quarter points?
2.	The tensile strength of dry and conditioned specimens determined at 77 ± 1.8 °F (25 ± 1 °C)?
3.	Specimens in the 77°F water bath removed and height measured?
4.	Specimens placed into the loading apparatus and the loading strips positioned so that they are parallel and centered
_	on the vertical diametral plane?
5.	Load applied to the specimen by means of the constant rate of movement of the testing
_	machine head of 2 in. per minute?
6.	Maximum compressive strength on the testing machine recorded? (P)
7.	Load continued until crack appears, specimen removed from the machine, pulled apart at the crack and inspected for degree of moisture damage?

Tensile Strength (kPa) = specimen thickness (mm) t =D = $\pi * t * D$ specimen diameter (mm)

P =

maximum load (N)

Calculations determined by as follows (see Section 9 for standard units calculation, also in IDT test)?..... 2000 * P

TSR = (average tensile strength of conditioned subset) / (average tensile strength of dry subset)

COMMENTS (D4867): (D4867)

(T287)	
(D4125)	

Date: _____

<u>APPARATUS</u>

1.		Nuclear gauge	
		(a) Neutron source - an encapsulated and sealed radioactive source (such as americium/beryllium)?	
		(b) Thermal neutron detector (such as helium-3 or boron tri-fluoride)?	
		(c) Read-out instrument displaying the percent asphalt binder to the nearest 0.1 percent?	
		(d) Daily standard log count?	
		(e) Factory or laboratory calibration data sheet?	
		(f) Leak test certificate?	
		(g) Shippers declaration for dangerous goods?	
		(h) Procedure memo for storing, transporting, and handling nuclear testing equipment?	
2.		Sample Pans, 3 or more made of stainless steel?	
3.		Balance, readable to 0.1 g with at least 20 kg capacity [ASTM: balance readable to 1 g]?	
4.		Heating Device	
		(a) Oven, capable of heating to $350 \pm 5^{\circ}F$ (177 $\pm 3^{\circ}C$)?	
	or	(b) AASHTO only: Microwave Oven, capable of maintaining a temp. of $177 \pm 3 ^{\circ}\text{C}$ (350 $\pm 5 ^{\circ}\text{F}$) and	
		determined to not be detrimental to aggregate?	
<i>5</i> .		ASTM only: straightedge, made of steel and approx. 18 inches long?	
6.		Leveling Plate, flat and rigid plate, made of:	
		(a) metal with a minimum thickness of 12.5 mm (0.5 in)?	
	or	(b) wood, with a minimum thickness of 19 mm (0.75 in) [ASTM: 20 mm]?	
	or	(c) AASHTO only: plexiglas, with a minimum thickness of 12.5 mm?	
7.		Thermometer, range of 50 to 500°F (10 to 260°C) [ASTM: range of 10 to 250°C (50 to 482°F)]?	
		Mechanical Mixer, with a 10 kg capacity?	
8.		Mechanical Mixer, with a 10 kg capacity?	
9.		Spoons and mixing bowls?	
10.		Splitting or quartering equipment?	
11.		AASHTO only: Sample containers, such as paint cans or unwaxed, non-absorbent cardboard boxes that can be closed to prevent contamination of the sample?	
<u> </u>	<u>M</u> (only: Additional apparatus for Test Method B (for compacted bituminous mixtures)	
1.		Apparatus necessary to prepare compacted specimens as specified in Test Methods D6926, D1561, D3387, or Practice D4013?	
2.		Molded laboratory specimen container provided by manufacturer (ref. Figure 1):	
-		(a) For 4 in. diameter specimens: container has 2 holes that can hold specimens (dia. 10.312 cm)?	
		(b) For 6 in. diameter specimens: container has 1 hole that can hold a specimen (dia. 15.392 cm)?	
		(c) Container fits inside nuclear gauge device (dimension are 24.689 x 18.161 x 6.985 cm)?	
		(d) Height from container bottom to specimen level is 5.715 cm?	

COMMENTS (T287 / D4125):

(T287 / D4125)

COMMENTS (T287):

ASPHALT CONTENT OF BITUMINOUS MIXTURES BY THE NUCLEAR METHOD

70	000	
(]	1287)

	ration and Verification (Assessor: Check Records)
1.	Sampling
	Approximately 50 kg (110 lbs) of aggregate obtained?
	Approximately 4 L (1 gal) of asphalt obtained?
3.	Aggregate Preparation
1.	If used, appropriate amount of lime hydrated onto aggregate?
<i>2</i> .	Aggregate dried to constant mass according to T255?
<i>3</i> .	Aggregate dry sieved?
	a) Sieve set includes 75-μm (No. 200) sieve?
	b) Cumulative mass required for each sieve size calculated as follows:
	X = T (100-P)/100 where:
	X = the required, cumulative batch mass for each specified sieve (g)
	T = the initial, total aggregate mass (g)
	P = the percent passing for each specified sieve according to the JMF*
4.	Aggregate dust correction performed?
7.	a) Prepare a washed-gradation sample from the masses calculated above?
	b) Perform a wash gradation according to T27 and T11?
	c) Calculate the corrected batch mass for each specified sieve for the calibration points as follows:
	c) Carculate the corrected batch mass for each specified sieve for the cultoration points as follows:
	$Z_n = X^2/Y$ where:
	$Z_n =$ the adjusted, cumulative batch mass for any sieve size, n (g)
	X = the pre-wash, cumulative batch mass for each specified sieve (g)
	Y = the post-wash, cumulative batch mass for each specified sieve (g)
	d) Blend the aggregate together at the proper proportion to match the JMF using the masses
	calculated in 4c?
	(Note to Assessors: JMF = Job Mix Formula. The job mix formula is the mix design specified for performing the calibrations.)

(T287)

COMMENTS (T287):

ASPHALT CONTENT OF BITUMINOUS MIXTURES BY THE NUCLEAR METHOD

(T287)

			AASHTO CALIBRATION (page 2 of 4) Date:
~			
		d Verification (Co	
<i>C</i> .		alt Binder Prepara	
1.			(0.5 gal) of asphalt binder to the mid-point of the mixing temperature
2			ainer? amount of liquid anti-stripping additive added to asphalt binder?
2. 3.			oon as it reaches the mid-point of the mixing temperature range (if this is not
J.			mperature for not more than 4 hours)?
4.			reheated?
7. 5.			r and aggregate required calculated by one of the following methods:
	11111011	ni oj uspitati otitue	and agging are required editionated by one of the following memous.
	Metho	od A – Asphalt bin	der percent by mass of the asphalt mixture
	1.		nass of asphalt binder for each calibration point as follows:
		$B = (E)(P_{bm})$	where:
			B = the mass of the asphalt binder to the nearest $0.1 g$
			$E = the \ mass \ of \ asphalt \ mixture \ (g)$
			P_{bm} = the percent of asphalt binder by total mass of the asphalt mixture,
			expressed as a decimal
	2.	1 minimum of t	our samples mixed, containing the following binder contents: 0.8% below
	2.		timum, 0.8% above optimum, and 1.6% above optimum?
	3.		nass of aggregate required for each calibration point (asphalt content) as follows?
	J.	Carculate the m	ass of aggregate required for each cultoration point (asphalt content) as follows:
		A = E - B	where:
			A = the mass of the aggregate to the nearest 0.1 g
			E = the mass of the asphalt mixture (g)
			B = the mass of asphalt (from above)
<u>OR</u>			
		A/	ASHTO Materials Reference Laboratory
	Metho	od B – Asphalt bin	der percent by mass of the aggregate
	1.	Calculate the m	nass of aggregate for each calibration point as follows:
		$A = E/(1+P_{ba})$	where:
			P_{ba} = the percent of asphalt binder by mass of the aggregate, expressed as a decimal
			E = the mass of the asphalt mixture (g)
	2	1	our samples mixed, containing the following binder contents: 0.8% below
	2.		timum, 0.8% above optimum, and 1.6% above optimum?
	<i>3</i> .		nass of asphalt binder required for each calibration point as follows:
	5.	Culculate the m	ass of aspiran officer required for each canoration point as follows
		$B = (A)(P_{ba})$	where:
		(-/ (- 04/	P_{ba} = the percent of asphalt binder by mass of the aggregate, expressed as a decimal
			$A = Mass \ of \ aggregate \ (from \ above)$

(T287)

9.

10.

11.

ASPHALT CONTENT OF BITUMINOUS MIXTURES BY THE NUCLEAR METHOD

(T287)

	AASHTO CALIBRATION (page 3 of 4) Date:
Calib	ration and Verification (Continued)
D.	Target Mass Determination
1.	Butter batch prepared to determine the mass to be used for the calibration samples?
2.	Mix the preheated aggregate and asphalt according to preparation of specimens section?
<i>3</i> .	Mass of a clean gage-sample pan determined and scale tared?
4.	Asphalt mixture placed into pan until pan is half full?
5.	Asphalt sample lightly tamped with preheated spoon or spatula?
6.	Remaining asphalt mixture placed in pan until it is mounded about 13 mm (0.5 in.) above the top of the pan?
7.	Leveling plate placed on top of mixture immediately after filling the pan?
8.	Sample compacted into the pan until it is level with the top of the pan by pressing down on the plate?
9.	Sight across the top of the pan to ensure that the asphalt mixture is not above the pan?
10.	Determine and record the mass (i.e. target mass) of the filled pan?
11.	Calibration and sample specimens within ± 5 g of the target mass?
E .	Preparation of Calibration Specimens (Mixing)
1.	Mass of the aggregate and asphalt binder determined for each sample according to Aggregate Preparation
	and Asphalt Binder Preparation Sections of the method?
2.	A minimum of four aggregate samples prepared, containing the following binder contents: 0.8% below
	optimum, at optimum, 0.8% above optimum, and 1.6% above optimum?
<i>3</i> .	Target mass used for each aggregate sample?
4.	Aggregate and asphalt binder materials heated to the mid-point of the mixing temperature range and
	allowed to stabilize at that temperature?
5.	Mass of heated mixing bowl determined to the nearest 0.1 g?
6.	Heated aggregate specimen placed in the mixing bowl?
7.	Crater formed in aggregate large enough to hold the required amount of asphalt binder?
8.	Mixing bowl placed on scale and required asphalt binder added into the crater to the nearest 0.1 g?
9.	Aggregate and asphalt mechanically mixed for a minimum of two minutes?
	Note: hand mixing is acceptable using large bowl and mixed for a minimum of five minutes. All material thoroughly coated after mixing procedure?
10.	All mater <mark>ial tho</mark> roughly coated after mixing procedure?
11.	If necessary, remix the sample by hand until it is thoroughly mixed?
<i>12</i> .	Mixture removed from mixing bowl, and the bowl weighed to ensure that all material is removed?
<i>13</i> .	Mass of bowl within ±5 g of its original mass?
14.	If not, bowl scraped with spatula and added to sample until sample mass is within tolerance?
F.	Calibration (Testing Calibration Specimens)
1.	Mass of a clean gage-sample pan determined and the pan tared on the scale?
2.	Asphalt mixture place in the pan until it is half full?
<i>3</i> .	Asphalt sample lightly tamped with preheated spoon or spatula?
4.	Remaining asphalt mixture placed in pan so that the mixture is mounded about 13 mm (0.5 in.) above
	the top of the pan?
<i>5</i> .	Leveling plate placed on top of the asphalt mixture immediately after filling the pan?
<i>6</i> .	Sample compacted into the pan until it is level with the top of the pan by pressing down on the leveling plate?.
<i>7</i> .	Sight across the top of the pan to ensure that the asphalt mixture does not protrude above the top of the pan?
8.	Mass of the compacted asphalt mixture in the pan determined and recorded?
	-,

COMMENTS (T287): (T287)

Pan placed into the gage, and manufacturer's instructions for operating the equipment and

Mass within ± 5 g of the target mass?.....

Repeat steps 1 through 11 for all calibration samples?.....

(12871

	AASHTO CALIBRATION (page 4 of 4) Date:
Calibi	ration and Verification (Continued)
<u>G.</u>	Calibration Curve
1.	Calibration curve prepared for each asphalt mixture type, aggregate blend, asphalt binder source, or addition of liquid anti-strip or hydrated lime?
2.	Calibration curve covers the range of expected values found in field samples?
3.	Do the limits for the calibration curve consider the allowable range of asphalt binder content plus the allowable aggregate moisture (which reads as asphalt binder in the gage)?
4.	At least four calibration-curve pans prepared at 0.8 below, optimum, 0.8 above, and 1.6 above the optimum asphalt binder content?
I.	Presentation of Calibration Data
For G	ages that generate the calibration internally to the gage:
1.	Formula coefficients, coefficient of fit, and the calculated percent difference for each calibration point printed or recorded?
2.	Calibration not acceptable if the coefficient of fit is less than 0.998 for a dense-graded asphalt mixture or 0.995 for an open-graded asphalt mixture, or any calibration point has a calculated difference greater than 0.09 percent?
3.	If calibrations are not acceptable are they performed again?
4.	Acceptable calibrations stored in the gages memory, using the JMF and an easily recognizable calibration number, according to the manufacturer's instructions?
For go	ages that do n <mark>ot generate the</mark> calibration internally to the gage:
1.	Calibr <mark>ati</mark> on curve pr <mark>e</mark> pared by plotting the gage readings for calibration samples versus asphalt binder content on linear graph paper, choosing convenient scale factors for the gage readings and asphalt binder content?
	Cross-Calibration (When Applicable) rocess creates a relationship between the field gage and the gage used in the JMF calibration (see Section A10 of the for details).
COM	MENTS (T287): (T287)

(T287)

	AASHTO PROCEDURE Date:	
Stando	ardization	
1. 2.	A background count performed in accordance with manufacturer's procedure each day prior to testing? Measurement times of the background count the same as testing time?	
3.	Background count does not change by more than 1 percent; if it does are more background counts performed?	
Proces	dure	
1.	Obtain a representative sample according to T168?	
<i>2</i> .	If required, sample reduced to appropriate size by splitting and quartering according to T248, Method B?	
<i>3</i> .	Test performed while mixture is hot, if sample cools it may be reheated to the mid-point of the compaction temperature range for the asphalt binder used?	
4.	Determine the mass of a clean gage-sample pan and tare the pan on the balance?	
<i>5</i> .	Pan filled half full with the asphalt mixture?	
6.	Asphalt mixture lightly tamped with a preheated spatula or spoon?	
7.	Additional asphalt added to the pan until the required mass, as found in the	
	Target Mass Determination, is reached within ± 5 g?	
8.	Leveling plate placed on top of the asphalt mixture immediately after filling the pan?	
9.	Sample compacted into the pan until it is level with the top of the pan by pressing down on the leveling plate?	
10.	Sight across the top of the pan to ensure that the asphalt mixture does not protrude above the top of the pan?	
11.	Mass of asphalt mixture compacted in the pan determined and recorded?	
12.	Mass within ± 5 g of the target mass?	
13.	If the gage has the ability to store multiple calibrations, activate the calibration for the mixture?	
14.	Pan containing mixture placed into the gage and a 4 minute count performed?	
<i>15</i> .	Uncorrected asphalt binder content determined by the direct read out on the gage, calibration	
16.	graph, or the formula supplied by the manufacturer, and recorded to the nearest 0.1 percent?	
10.	(a) Moisture correction determined by performing T110, T329, or by microwave oven at temp. of $177 \pm 3 \text{C}$ (350 $\pm 5 \text{F}$), using a representative portion of the sample?	
	(b) Moisture content recorded to the nearest 0.1 percent?	
	(c) Moisture content subtracted from the uncorrected asphalt binder content and reported to the nearest 0.1 percent?	
COM	MENTS (T287):	(T287)

(D4125)

		ASTM CALIBRATION (page 1 of 2) Date:
Calib	ration (g	eneral)
1.		ration curve developed according for each mix-type and aggregate blend?
2.		calibration curve developed whenever there is a change is asphalt/aggregate source,
		icant change in aggregate gradation, or new or repaired apparatus?
	υ	
Calib	ration –	Method A
A.	Blank	Sample
	1.	Blank sample (with no asphalt) prepared to determine mass used for calibration and
		plant mix samples?
	2.	Blank prepared by filling a sample pan with aggregate in 3 layers, avoiding segregation?
	3.	Each layer settled by raising pan 1 to 2 in. and tapping to work surface 2 or 3 times?
	4.	Third layer filled to a point slightly above top edge of pan?
	5.	Straightedge used to make sample flush with top of pan with no compaction to sample?
	6.	Mass of blank sample determined and recorded to nearest 1 g?
	7.	This mass used for all calibration and plant mix samples?
	8.	Sample pan placed in gauge and sample count recorded?
В.	Calib	ration Curve Samples
υ.	1.	Enough aggregate obtained for a minimum of 3 samples [approx. 30 kg (65 lb)]?
	2.	Approx. 2.5 kg (5.5 lb) of asphalt obtained?
	3.	Minimum of 3 samples prepared to establish calibration curve?
	4.	Range of asphalt contents vary by at least 2% from lowest to highest?
	5.	Range encompasses the design asphalt content?
	6.	Masses of the samples agree within 10 g of each other?
	7.	Calibration samples prepared and tested as close as possible to temperature of test samples
		of completed mixes?
	8.	Calibration samples tested within ±5°C (9°F) of each other?
	9.	Aggregates and asphalt for the three samples heated to approx. 150°C (300°F)?
	10.	All bowls, sample pans, and tools also heated to approx. 150°C (300°F)?
	11.	Prior to mixing first sample, mixture of asphalt and aggregate fines used to butter the bowl?
	12.	Aggregate placed in bowl and asphalt added to within 1 g of desired percent by mass?
	13.	Mixed thoroughly?
	14	Mixed sample placed in sample pan in 3 layers?
	15.	Each layer distributed evenly using scoop or spatula?
	16.	Each layer settled by raising pan 1 to 2 in. and tapping to work surface 2 or 3 times?
	17.	Third layer filled to a point slightly above top edge of pan?
	18.	Material added or removed from sample until mass is within 10 g of blank sample?
	19.	Mass recorded?
	20.	Sample compacted with flat metal or wood plate until level with top of pan?
	21.	Sample pan placed in gauge and sample count recorded?
	22.	Steps 1 through 21 repeated for each of the remaining mixes?
	23.	Bowl not completely cleaned between remaining mixes?

COMMENTS (D4125): (D4125)

COMMENTS (D4125):

ASPHALT CONTENT OF BITUMINOUS MIXTURES BY THE NUCLEAR METHOD

(D4125)

Calib	ration – Method B
1.	Aggregates sampled and prepared for blending according to method of compaction selected?
2.	Enough asphalt sampled for test?
3.	Four known asphalt contents selected?
4.	Range of asphalt contents vary by at least 2% from lowest to highest?
5.	Three specimens prepared for each of the selected asphalt contents?
6.	Specimens compacted by one of the following methods: D1561, D3387, D6926 or Practice D4013?
7.	Masses of each set of replicate specimens within 10 g (0.02 lb) of each other?
8.	Mass of each specimen determined to nearest 1 g (0.002 lb)?
9.	For 4-in. specimens: Two specimens whose masses are closest to each other selected for
<i>)</i> .	each of the asphalt contents?
10.	For 6-in. specimens: One specimen selected from each asphalt content such that masses
10.	of four specimens selected are as close as possible?
11.	Specimens not used retained for further testing?
12.	Specimens placed in molded specimen container and container placed in gauge?
13.	Sample count recorded?
14.	Process repeated for each set of calibration samples?
15.	For samples containing RAP: RAP material is of uniform gradation, asphalt content, and asphalt type?
16.	RAP calibration samples contain same percentage of RAP as samples to be tested?
10. 17.	Method not used for job control testing if RAP material is not uniform?
1/.	Method not used for job control testing if KAT material is not uniform?
10	Final galibration regroups is in form of ourse data table or equation for computer processing?
18.	Final calibration response is in form of curve, data table, or equation for computer processing?
18.	Final calibration response is in form of curve, data table, or equation for computer processing? Note to Assessors: Some gauges will do this on their own.
18.	Final calibration response is in form of curve, data table, or equation for computer processing?
	Final calibration response is in form of curve, data table, or equation for computer processing?
<u>Calib</u>	Final calibration response is in form of curve, data table, or equation for computer processing?
<u>Calib</u> 1.	Final calibration response is in form of curve, data table, or equation for computer processing?
<u>Calib</u>	Final calibration response is in form of curve, data table, or equation for computer processing?
<u>Calib</u> 1.	Final calibration response is in form of curve, data table, or equation for computer processing?
<u>Calib</u> 1. 2.	Final calibration response is in form of curve, data table, or equation for computer processing?
<u>Calib</u> 1. 2. Back ;	Final calibration response is in form of curve, data table, or equation for computer processing?
<i>Calib</i> 1. 2. <i>Back</i> 1.	Final calibration response is in form of curve, data table, or equation for computer processing?
<u>Calib</u> 1. 2. <u>Back</u> 1. 2.	Final calibration response is in form of curve, data table, or equation for computer processing? Note to Assessors: Some gauges will do this on their own. **ration Factor – Methods A and B* For each calibration, correlation factor calculated according to Equation (1) of test method? Correlation factor greater than or equal to 0.995? **ground and Stability Check** Background radiation count obtained each day before taking measurements? Measurement period of background count ≥ normal measurement period?
<u>Calib</u> 1. 2. <u>Back</u> 1. 2. 3.	Final calibration response is in form of curve, data table, or equation for computer processing?
Calib 1. 2. Backs 1. 2. 3.	Final calibration response is in form of curve, data table, or equation for computer processing? Note to Assessors: Some gauges will do this on their own. ration Factor – Methods A and B For each calibration, correlation factor calculated according to Equation (1) of test method? Correlation factor greater than or equal to 0.995? Background and Stability Check Background radiation count obtained each day before taking measurements? Measurement period of background count ≥ normal measurement period? New background count taken if gauge has been moved (even within same room)? New background count taken if environment around gauge has been changed significantly?
Calib 1. 2. Backs 1. 2. 3.	Final calibration response is in form of curve, data table, or equation for computer processing? Note to Assessors: Some gauges will do this on their own. For each calibration, correlation factor calculated according to Equation (1) of test method? Correlation factor greater than or equal to 0.995? Background and Stability Check Background radiation count obtained each day before taking measurements? Measurement period of background count ≥ normal measurement period? New background count taken if gauge has been moved (even within same room)? New background count taken if environment around gauge has been changed significantly? For gauges that have not been moved, background count within 1% of previous count
Calib 1. 2. Back; 1. 2. 3. 4.	Final calibration response is in form of curve, data table, or equation for computer processing? Note to Assessors: Some gauges will do this on their own. For each calibration, correlation factor calculated according to Equation (1) of test method? Correlation factor greater than or equal to 0.995? Background and Stability Check Background radiation count obtained each day before taking measurements? Measurement period of background count ≥ normal measurement period? New background count taken if gauge has been moved (even within same room)? New background count taken if environment around gauge has been changed significantly? For gauges that have not been moved, background count within 1% of previous count for gauge to be considered stable?
Calib 1. 2. Back; 1. 2. 3. 4.	Final calibration response is in form of curve, data table, or equation for computer processing? Note to Assessors: Some gauges will do this on their own. Pration Factor – Methods A and B For each calibration, correlation factor calculated according to Equation (1) of test method? Correlation factor greater than or equal to 0.995? Background radiation count obtained each day before taking measurements? Measurement period of background count ≥ normal measurement period? New background count taken if gauge has been moved (even within same room)? New background count taken if environment around gauge has been changed significantly? For gauges that have not been moved, background count within 1% of previous count for gauge to be considered stable? For gauges that have been moved, background count within 2 - 3% of previous count
Calib 1. 2. Backs 1. 2. 3. 4. 5.	Final calibration response is in form of curve, data table, or equation for computer processing? Note to Assessors: Some gauges will do this on their own. Pration Factor – Methods A and B For each calibration, correlation factor calculated according to Equation (1) of test method? Correlation factor greater than or equal to 0.995? Background and Stability Check Background radiation count obtained each day before taking measurements? Measurement period of background count ≥ normal measurement period? New background count taken if gauge has been moved (even within same room)? New background count taken if environment around gauge has been changed significantly? For gauges that have not been moved, background count within 1% of previous count for gauge to be considered stable? For gauges that have been moved, background count within 2 - 3% of previous count for gauge to be considered stable? ———————————————————————————————————
Calib 1. 2. Backs 1. 2. 3. 4. 5.	Final calibration response is in form of curve, data table, or equation for computer processing? Note to Assessors: Some gauges will do this on their own. Factor – Methods A and B For each calibration, correlation factor calculated according to Equation (1) of test method? Correlation factor greater than or equal to 0.995? Background and Stability Check Background radiation count obtained each day before taking measurements? Measurement period of background count ≥ normal measurement period? New background count taken if gauge has been moved (even within same room)? New background count taken if environment around gauge has been changed significantly? For gauges that have not been moved, background count within 1% of previous count for gauge to be considered stable? For gauges that have been moved, background count within 2 - 3% of previous count for gauge to be considered stable? If background counts are not within limits or if gauge stability is suspect, statistical stability test
Calib 1. 2. Backs 1. 2. 3. 4. 5. 6.	Final calibration response is in form of curve, data table, or equation for computer processing? Note to Assessors: Some gauges will do this on their own. ration Factor – Methods A and B For each calibration, correlation factor calculated according to Equation (1) of test method? Correlation factor greater than or equal to 0.995? Background and Stability Check Background radiation count obtained each day before taking measurements? Measurement period of background count ≥ normal measurement period? New background count taken if gauge has been moved (even within same room)? New background count taken if environment around gauge has been changed significantly? For gauges that have not been moved, background count within 1% of previous count for gauge to be considered stable? For gauges that have been moved, background count within 2 - 3% of previous count for gauge to be considered stable? If background counts are not within limits or if gauge stability is suspect, statistical stability test performed according to manufacturer's instructions?
Calib 1. 2. Backs 1. 2. 3. 4. 5. 6.	Final calibration response is in form of curve, data table, or equation for computer processing?
Calib 1. 2. Backs 1. 2. 3. 4. 5. 6. 7.	Final calibration response is in form of curve, data table, or equation for computer processing? Note to Assessors: Some gauges will do this on their own. Pration Factor – Methods A and B For each calibration, correlation factor calculated according to Equation (1) of test method? Correlation factor greater than or equal to 0.995? Depound and Stability Check Background radiation count obtained each day before taking measurements? Measurement period of background count ≥ normal measurement period? New background count taken if gauge has been moved (even within same room)? New background count taken if environment around gauge has been changed significantly? For gauges that have not been moved, background count within 1% of previous count for gauge to be considered stable? For gauges that have been moved, background count within 2 - 3% of previous count for gauge to be considered stable? If background counts are not within limits or if gauge stability is suspect, statistical stability test performed according to manufacturer's instructions? Statistical stability test also performed when gauge is new or repaired? Statistical stability test performed at least once a month otherwise?
Calib 1. 2. Backs 1. 2. 3. 4. 5. 6. 7. 8. 9. 10.	Final calibration response is in form of curve, data table, or equation for computer processing? Note to Assessors: Some gauges will do this on their own. Factor – Methods A and B For each calibration, correlation factor calculated according to Equation (1) of test method? Correlation factor greater than or equal to 0.995? Measurement period of background count ≥ normal measurements? Measurement period of background count ≥ normal measurement period? New background count taken if gauge has been moved (even within same room)? New background count taken if environment around gauge has been changed significantly? For gauges that have not been moved, background count within 1% of previous count for gauge to be considered stable? For gauges that have been moved, background count within 2 - 3% of previous count for gauge to be considered stable? If background counts are not within limits or if gauge stability is suspect, statistical stability test performed according to manufacturer's instructions? Statistical stability test also performed when gauge is new or repaired? Statistical stability test performed at least once a month otherwise? Failure of statistical stability test prompts a check for hydrogen bodies in or around gauge?
Calib 1. 2. Backs 1. 2. 3. 4. 5. 6. 7.	Final calibration response is in form of curve, data table, or equation for computer processing? Note to Assessors: Some gauges will do this on their own. Pration Factor – Methods A and B For each calibration, correlation factor calculated according to Equation (1) of test method? Correlation factor greater than or equal to 0.995? Depound and Stability Check Background radiation count obtained each day before taking measurements? Measurement period of background count ≥ normal measurement period? New background count taken if gauge has been moved (even within same room)? New background count taken if environment around gauge has been changed significantly? For gauges that have not been moved, background count within 1% of previous count for gauge to be considered stable? For gauges that have been moved, background count within 2 - 3% of previous count for gauge to be considered stable? If background counts are not within limits or if gauge stability is suspect, statistical stability test performed according to manufacturer's instructions? Statistical stability test also performed when gauge is new or repaired? Statistical stability test performed at least once a month otherwise?

Revised 2011-03-25

(D4125)

COMMENTS (D4125):

ASPHALT CONTENT OF BITUMINOUS MIXTURES BY THE NUCLEAR METHOD

(D4125)

		ASTM PROCEDURE Date:
	mplir	
1.		Test sample obtained according to D979 (Standard Practice for Sampling HMA)?
2.		Moisture content of the test sample determined:
		(a) By ASTM D1461 (Moisture or Volatile Distillates Content of HMA)?
	or	(b) By drying test sample to constant mass in an oven at 110 ± 5 °C (230 ± 9 °F)?
_	ethod	
1.		Sample pan filled in three layers?
2.		Each layer distributed evenly with scoop or spatula?
3.		After each layer, pan lifted approx. 20 to 50 mm (1 to 2 in.) and tapped on working surface two
		or three times to settle contents?
4.		Last layer fills pan slightly above top edge?
5.		Material added or removed until mass of mix in pan is within 10 g (0.02 lb) of the calibration sample?
6.		Mass of mix in pan recorded?
7.		Sample compressed with flat plate until level with top edge of pan?
8.		Temperature of sample recorded?
9.		Temperature within 5°C (9°F) of calibration temperature unless the apparatus makes provision
		for temperature correction?
10.		Sample placed in chamber?
11.		Manufacturer's instructions followed to obtain sample counts?
12.		Asphalt content of mixture determined?
13.		Corrected for moisture content, if necessary?
	•	
Me	ethod	B
1.		Specimens prepared using Test Method, D1561, D3387, D6926, or Practice D4013?
2.		Two specimens used for 10-cm (4 in.) diameter specimens?
3.		Two specimens used for 10-cm (4 in.) diameter specimens? One specimen used for 15-cm (6 in.) diameter specimens?
4.		One of the following:
••		(a) For 10-cm specimens: the mass of the two test specimens are within 10 g (0.02 lb) of each other and
		the average of the two test specimens are within 10 g (0.02 lb) of the average of the calib. samples?
	or	(b) For 15-cm specimens: the mass of the test specimen is within 10 g (0.02 lb) of the calibration sample?
5.	OI	Sample(s) placed in the molded specimen container and then placed in the testing chamber?
6.		Manufacturer's instructions followed to obtain sample counts?
7.		Asphalt content of mixture determined?
8.		Corrected for moisture content, if necessary?
ο.		Corrected for moisture content, it necessary?

Revised 2011-03-25

(D4125)

DETERMINING THE ASPHALT CONTENT OF HMA BY THE IGNITION METHOD

(T308)	
(D6307)	

Date:

<u>APPARATUS</u>

1.	Ignition	Furnace						
	(a)	One of the following:						
		(1) Convection-type furnace that maintains 578°C (1072°F) [ASTM: 580°C]?						
		(2) Direct Irradiation-type furnace with an internal balance?						
	(b)	Dimensions adequate to accommodate a 3500 g sample [ASTM: 2500 g sample]?						
	(c)	Door cannot be opened during test (do not attempt to open it during the test!)?						
	(d)	Method for reducing furnace emissions?						
	(e)	Vented into a hood or to the outside?						
	(f)	A fan to pull air through the furnace?						
	(g)	Sample Baskets:						
	(Ο)	(1) Allows sample to be thinly spread?						
		(2) Allows air to flow through and around sample?						
		(3) Sets of two or more baskets nested?						
		(4) Sample completely enclosed with a mesh screen or perforated stainless steel plate?						
	(h)	Catch pan of sufficient size to hold baskets?						
2.	Internal	balance (Method A only)						
	(a)	AASHTO only: Thermally isolated from furnace chamber?						
	(b)	Accurate to 0.1 g?						
	(c)	Capable of weighing a 3500 g [ASTM: 2500 g] sample in addition to the baskets?						
3.	Data co	llection system (Method A only)						
<i>J</i> .	(a)	Mass can be determined and displayed during the test?						
	(b)	Built-in computer program?						
	(c)	Calculates the change in mass?						
	(d)	AASHTO only: Provides for the input of a correction factor?						
	(e)	AASHTO only: Audible alarm and indicator light?						
	(f)	Capable of changing the ending mass loss percentage to 0.02 percent [ASTM: 0.01%]?						
4.	Printed	ticket (Method A only)						
••	(a)	Records initial specimen mass?						
	(b)	Records specimen mass loss?						
	(c)	Records temperature compensation?						
	(d)	Records correction factor?						
	(e)	Records corrected asphalt content (%)?						
	(f)	Records test time?						
	(g)	Records test temperature?						
	(8)	Note to assessors: NCATs should be set to print the long ticket.						
5.	Miscella	aneous						
	(a)	Oven capable of maintaining a temperature of $110 \pm 5^{\circ}\text{C}$ ($230 \pm 9^{\circ}\text{F}$)?						
	(b)	Balance readable to 0.1 g, conforming to M231, Class G2 [ASTM: D4753, class GP2]?						
	(c)	AASHTO only: Protective cage capable of surrounding baskets?						

COMMENTS (T308 / D6307):

(1)

(2)

(3)

DETERMINING THE ASPHALT CONTENT OF HMA BY THE IGNITION METHOD

(T308)

		AASHTO PROCEDURE Date:
1.	Correc	ction Factor (AASHTO only)
	(a)	Determined for each job mix formula (JMF)?
	(b)	Determined before any acceptance testing is completed and if any change in ingredients or design occurs?
	(c)	A new correction factor is determined if any changes greater than 5% in stockpiled aggregate proportions occur?
	(d)	"Butter mix" prepared and discarded to condition bowl?
	(e)	"Blank" aggregate specimen batched and tested for aggregate gradation?
	(f)	Two correction factor samples mixed at the design asphalt content?
	(g)	Freshly mixed samples placed directly into basket assembly or
	(8)	if allowed to cool, dried to constant mass at $110 \pm 5^{\circ}$ C?
	(h)	Baskets are not preheated?
	(i)	Specimens tested according to method?
	(j)	Gradation analysis performed on residual aggregate and compared to blank [1(c) above]?
	(k)	Asphalt contents determined?
	(1)	If the asphalt contents differ by more than 0.15 percent:
		(1) Test repeated with two more samples?
		(2) Highest and lowest result discarded from the four tests?
		(3) Correction factor determined from the two remaining results?
	(m)	Convection-type furnace - if correction factor exceeds 1.0 percent, test repeated at
	()	482 ± 5 °C (900 ± 8°F), and the resulting correction factor used for further testing?
	(n)	One of the following
	` '	(1) Test temperature is the same as that of correction factor?
		(2) <u>Direct irradiation-type furnace</u> - DEFAULT burn profile used for most materials?
		Note: Burn profile OPTION 1 or OPTION 2 may be selected to optimize burn cycle?
		OPTION 1 is designed for samples with correction factor greater than 1.0 percent.
		OPTION 2 is designed for samples that may not burn completely using DEFAULT burn profile.
Note. 2 testing	AASHTO or gprocedure	nly <mark>: Histor</mark> ical data or scientific studies may be used to determine the correction factor(s) in lieu of using this if th <mark>e te</mark> sting agency provides reference to the studies/data.
2.	<u>Aggreg</u>	gate Correction Factor (AASHTO Only)
	(a)	Aggregate correction factor determined for aggregates known for excessive
		breakdown or from an unknown source?
	(b)	Gradation analysis performed on residual aggregate for each correction factor sample?
		Note: residual aggregate is the aggregate removed from the ignition oven when
		determining the asphalt content correction factor.
	(c)	Difference between percent passing a given sieve in the correction factor samples and in the blank sample determined and the average difference calculated?
	(d)	If the average difference is greater than allowable, correction factor average for any sieve (equal to

COMMENTS (T308): (T308)

resultant average difference) for all sieves applied to all test results?

Sieves larger than or equal to 2.36-mm (No. 8) allowed 5% difference?

Sieves larger than or equal to 75-µm (No. 200) allowed 3% difference?.....

Sieves smaller than 75- μm (No. 200) (or bottom pan) allowed 0.5% difference?.....

DETERMINING THE ASPHALT CONTENT OF HMA BY THE IGNITION METHOD

$(T308)_{-}$	
(D6307)	

Calibi	ration Facto	r (ASTM only)					
(a)				n mix design o	· ingredient	s?		
(b)						sed for the HMA test sample?		
(c)	Aggregate oven dried to constant mass (no temperature specified)?							
(d)	Aggregate, asphalt cement, and all mixing bowls and tools heated to approx. 150 °C?							
(e)	Butter mix prepared to condition bowl?							
Ó	Three calibration samples mixed at the design asphalt content?							
(g)								
(h)						•••••		
(i)		ontents detern						
	Convecti	on-type furnac	ce					
				that of calibrat	ion (540 ± 3	5°C)?		
						perature to 482 \pm 5 $^{\circ}$ C and		
						0 percent?		
		-		,		· I		
		adiation –type						
	(3)	Burn profile so	et to DEFAULT f	for most materia	ıls?			
						pptimize burn cycle?		
		OPTION 1 is de	signed for aggrega	tes with correction	n factor gree	iter than 1.0 percent.		
<i>(j)</i>	C <mark>ali</mark> brati	OPTION 2 is de on factor deter	rsigned for samples	that may not bur	n completely	using DEFAULT burn profile.		
(j)		OPTION 2 is de on factor deter	rsigned for samples	that may not bur	n completely	using DEFAULT burn profile. total mass <u>B</u> efore ignition total mass <u>A</u> fter ignition % of asphalt cement by		
<i>(j)</i>	C <mark>ali</mark> brati	OPTION 2 is deconfactor determined (B – A) *100	esigned for samples rmined by calcula 0 P	that may not bur tion below? Where:	B = A = P =	using DEFAULT burn profile. total mass <u>B</u> efore ignition total mass <u>A</u> fter ignition % of asphalt cement by		
	Calibrati C _F =	OPTION 2 is deconfactor determined (B – A) *100 B	esigned for samples rmined by calcula 0 P	that may not burtion below? Where:	B = A = P =	using DEFAULT burn profile. total mass <u>B</u> efore ignition total mass <u>A</u> fter ignition % of asphalt cement by		
	Calibrati C _F =	OPTION 2 is deconfactor determined (B – A) *100 B	esigned for samples rmined by calcula 0 P	that may not burtion below? Where:	B = A = P =	total mass <u>B</u> efore ignition total mass <u>A</u> fter ignition % of asphalt cement by mass of the total mix		
<i>(k)</i> Sampl	Calibrati C _F = Average	OPTION 2 is decon factor deter (B - A) *100 B AAS H of the three tai	esigned for samples rmined by calcula 0 P ken and used as to	that may not bur tion below? Where: he calibration f	B = A = P = feren	total mass <u>B</u> efore ignition total mass <u>A</u> fter ignition % of asphalt cement by mass of the total mix		
(k) Sample (a)	Calibration CF = Average de Preparation Mixture	OPTION 2 is deformable on factor determined in an open constant $\frac{(B-A)*100}{B}$.	rmined by calcula P ken and used as to	that may not bur tion below? Where: the calibration for	B = A = P = feren	total mass <u>Before</u> ignition total mass <u>After</u> ignition % of asphalt cement by mass of the total mix		
(k) Sampl (a) (b)	Calibration CF = Average de Preparation Mixture of Sample in	OPTION 2 is deform factor determined in an expectation of warmed in an expectation of warmed in $\frac{1}{2}$	rmined by calcula P ken and used as to oven at 110 ± 5°C oven for extended	that may not bur tion below? Where: the calibration f (230 \pm 9°F) unt period of time?	B = A = P = A il it can be h	total mass <u>Before</u> ignition total mass <u>After</u> ignition % of asphalt cement by mass of the total mix		
(k) Sample (a) (b) (c)	Calibration CF = Average de Preparation Mixture of Sample in Particles	on factor determined in an expectation of mixture separate determined in an expectatio	esigned for samples rmined by calcula 0	that may not bur tion below? Where: the calibration f (230 \pm 9°F) unt period of time? a or trowel?	B = A = P = A il it can be h	total mass <u>Before</u> ignition total mass <u>After</u> ignition % of asphalt cement by mass of the total mix		
(k) Sampl (a) (b) (c) (d)	Calibration CF = Average Mixture of Sample in Particles Sample of	on factor determined in an of of mixture september of the three tandard in the continuous of mixture september of the three by red	esigned for samples rmined by calcula 0	that may not bur tion below? Where: (230 ± 9°F) unt period of time? a or trowel? ple [ASTM: by	B = A = P = il it can be be splitting or	total mass Before ignition total mass After ignition % of asphalt cement by mass of the total mix andled if necessary?		
(k) Sample (a) (b) (c)	Calibration CF = Average Mixture of Sample in Particles Sample of Sample in Sample	on factor determined in an of the three tandard of mixture septembers at least as	princed for samples rmined by calcula P ken and used as to even for extended arated with spatul ucing a larger sam much as indicated	that may not bur tion below? Where: (230 ± 9°F) unt period of time? a or trowel? pple [ASTM: by d on table below	B = A = P = il it can be be splitting or	total mass <u>Before</u> ignition total mass <u>After</u> ignition % of asphalt cement by mass of the total mix		
(k) Sampl (a) (b) (c) (d)	Calibration CF = Average Mixture of Sample in Particles Sample in 1200 g for sample in 120	on factor deter (B-A) *100 B of the three tan of warmed in an of of mixture septiatined by red nass at least as or No. 4 [ASTA)	exigned for samples rmined by calcula 0	that may not bur tion below? Where: (230 ± 9°F) unt period of time? a or trowel? pple [ASTM: by d on table below	B = A = P = il it can be be splitting or	total mass Before ignition total mass After ignition % of asphalt cement by mass of the total mix andled if necessary?		
(k) Sampl (a) (b) (c) (d)	Calibration CF = Average Mixture of Sample of Sample of 1200 g for 1200 g	on factor determined in an object of mixture septential by red mass at least as or No. 4 [ASTM or 3/8 in. [ASTM]	princed for samples rmined by calcula P ken and used as to even for extended arated with spatul ucing a larger sam much as indicated	that may not bur tion below? Where: (230 ± 9°F) unt period of time? a or trowel? pple [ASTM: by d on table below	B = A = P = il it can be be splitting or	total mass Before ignition total mass After ignition % of asphalt cement by mass of the total mix andled if necessary?		
(k) Sampl (a) (b) (c) (d)	Calibration CF = Average Mixture of Sample of Sample of 1200 g for 1500 g	on factor determined in an experiment of the three tables at least as as in No. 4 [ASTM or 3/8 in. [ASTM or 1/2 in.?]	exigned for samples rmined by calcula 0	that may not bur tion below? Where: (230 ± 9°F) unt period of time? a or trowel? pple [ASTM: by d on table below	B = A = P = il it can be be splitting or	total mass Before ignition total mass After ignition % of asphalt cement by mass of the total mix andled if necessary?		
(k) Sampl (a) (b) (c) (d)	Calibration CF = Average Le Preparation Mixture of Sample of Particles Sample of 1200 g for 1500 g for 2000 g for 20	on factor determined in an experiment of the three tandard warmed in an experiment of mixture septiatined by rednass at least as an No. 4 [ASTM or 1/2 in.? or 1/2 in.?	exigned for samples rmined by calcula 0	that may not bur tion below? Where: (230 ± 9°F) unt period of time? a or trowel? pple [ASTM: by d on table below	B = A = P = il it can be be splitting or	total mass Before ignition total mass After ignition % of asphalt cement by mass of the total mix andled if necessary?		
(k) Sampl (a) (b) (c) (d)	Calibration CF = Average Le Preparation Mixture of Sample of Particles Sample of 1200 g for 1200 g for 2000 g for 3000 g for 30	on factor determined in an order of the three tands of mixture septembers at least as as To No. 4 [ASTM or 3/8 in. [ASTM or 3/4 in.?]	exigned for samples rmined by calcula 0	that may not bur tion below? Where: (230 ± 9°F) unt period of time? a or trowel? pple [ASTM: by d on table below	B = A = P = il it can be be splitting or	total mass Before ignition total mass After ignition % of asphalt cement by mass of the total mix andled if necessary?		
(k) (a) (b) (c) (d) (e)	Calibration CF = Average Mixture of Sample in Particles Sample in 1200 g for 1500 g for 2000 g for 3000 g for 4000 g f	on factor determined in an experiment of the three tandard warmed in an experiment of mixture septiatined by reduces at least as for No. 4 [ASTM or 3/8 in. [ASTM or 3/4 in.?] or 1 in.? or 1 1/2 in.?	exigned for samples rmined by calcula 0	that may not bur tion below? Where: (230 ± 9°F) unt period of time? a or trowel? ple [ASTM: by d on table below []? [] in.]?	B = A = P = ference actor?	total mass <u>Before</u> ignition total mass <u>After</u> ignition % of asphalt cement by mass of the total mix nandled if necessary?		
(k) Sample (a) (b) (c) (d)	Calibration CF = Average Mixture of Sample in Particles Sample in 1200 g for 1500 g for 2000 g for 3000 g for 4000 g for AASHTO	on factor determined in an order of the three tandom warmed in an order of mixture septiatined by reduces at least as or No. 4 [ASTM or 3/8 in. [ASTM or 1/2 in.? or 1/2 in.? only: Specime only: Specime of the three tandom is a septiment of the three tandom is a	exigned for samples rmined by calcula 0	that may not bur tion below? Where: (230 ± 9°F) unt period of time? a or trowel? ple [ASTM: by I on table below I? B in.]? than 500 g great	B = A = P = ference actor?	total mass Before ignition total mass After ignition % of asphalt cement by mass of the total mix andled if necessary?		

COMMENTS (T308 / D6307):

DETERMINING THE ASPHALT CONTENT OF HMA BY THE IGNITION METHOD

(T308)	
(D6307)	

Date:

PROCEDURE	(Continued))
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(a)		Convection	on-type furnace preheated	d to 538°C (1000°F) o	r the correc	etion		
` ′								
	or							
<i>(b)</i>		<i>AASHTO</i>	only: Convection-type f	urnace, temperature r	ecorded pri	ior to test (can be automatic)?		
(c)		Sample d	ried to constant mass at 1	105 ± 5 °C (221 ± 9°F)	ASTM: 1	$10 \pm 5^{\circ}C$]?		
	or		imen for moisture determ					
		determine	ed according to (T110 / D	01461)?				
(d)		<i>AASHTO</i>	only: Correction factor	entered for the mix or	manually i	recorded?		
(e)		Basket(s)	placed in catch pan and	weighed with guards i	n place?			
<i>(f)</i>		<i>AASHTO</i>	only: Sample evenly dis	tributed in the basket,	material ke	ept away from edges and leveled?		
(g)						rded?		
(h)		Initial ma	iss of the specimen calcul	lated?				
(i)		<i>AASHTO</i>	only: Initial mass entere	ed into the furnace cor	troller and	l verified?		
(j)		Baskets p	placed in the furnace and	chamber door closed?				
(k)		<i>AASHTO</i>	only: Furnace scale agr	rees within 5 g of the to	otal mass?			
(l)		Pressing t	the start button locks the	door and starts the blo	wer?			
(m)						or three consecutive minutes?		
		Note, AAS	SHTO only: Ending mass los	ss percentage of 0.02 per	cent may be	used for excessive aggregate loss.)		
	ar ræ	00116						
	SHI		ethod A (continued)	1 1 1 1 1 1 1	1.	2		
(a)		Pressing	the stop button unlocks th	ne aoor ana prints the	iesi resuiis	;?		
(b)	/	Corrected	a aspnait content (%) jroi	m tne printea ticket re _l	00rtea?			
	or	Ij aspnan	content on ticket is not c	the printed ticket and	nnaer corr	rection factor subtracted?		
(0)	or	Percent n	noisiure subtractea from	ine primiea lickei ana	ine resuitat	nt value reported?roximately 30 minutes?		
(c)		Duskeis r	emovea ana allowea lo c	ooi io room iemperatu	re jor appr			
107	TM I	Only Moth	had A (continued)	laterials Re	eterer	nce Laboratory		
(a)	I IVI	Final ma	<u>10u A (comunueu)</u> 155 obtain <i>a</i> d by subtractiv	ng the wass loss by th	o furnaco f	from the initial mass of the mix?.		
(u) (b)		Corrected	as oouuneu oy suonucui d asnhalt contont calcula	ig the muss toss by the ited by the formula be	i juinuce ji Iow?			
(0)			e furnace may measure and					
		%AC =	(B-A) *100	Where:	B =	total mass <u>B</u> efore ignition		
			C_F		A =	total mass <u>A</u> fter ignition		
			\boldsymbol{B}		$C_F =$	calibration factor		

COMMENTS (T308 / D6307):

COMMENTS (T308 / D6307):

DETERMINING THE ASPHALT CONTENT OF HMA BY THE IGNITION METHOD

(T308)	
(D6307)	

(a)	<u>Procedure, Method B</u> Furnace preheated to 538°C (1000°F) or the correction factor temperature [$ASTM: 540 \pm 5$ ° C]?
(b)	Sample dried to constant mass at 105 ± 5 °C (221 ± 9°F) [ASTM: 110 ± 5 °C]?
or	AASHTO only: Test specimen for moisture determination obtained if necessary and moisture
	content determined according to (T110 / D1461) or by other suitable method?
(c)	AASHTO only: Correction factor recorded for the mix?
(d)	Basket(s) placed in catch pan and weighed with guards in place?
(e)	AASHTO only: Sample evenly distributed in the basket, material kept away from edges and leveled?
(f)	Total mass of the sample, basket, catch pan and basket guards recorded?
(g)	Initial mass of the specimen calculated?
(h)	Sample burned in the furnace for at least 45 minutes?
(i)	Sample removed and allowed to cool to room temperature at least 30 minutes [ASTM: 10 minutes]?
(j)	Sample weighed after ignition to the nearest 0.1 g?
(k)	Sample placed back in the furnace?
(1)	Sample burned for at least 15 minutes after reaching set temperature?
(m)	Steps (i) through (l) repeated until change in mass does not exceed 0.01
	percent of the initial sample mass?
	Note, AASHTO only: Ending mass loss percentage of 0.02 percent may be used for excessive aggregate loss.)
(n)	Corrected asphalt content (%) determined by calculation listed as ASTM above?
or	AASHTO only: If a moisture content has been determined, subtract the percent moisture from
	the AC percent, and report the resultant value as the corrected asphalt binder content?
Guadat	ion (AASHTO only)
(a)	Contents emptied into a flat pan, including any residual fines?
(b)	Gradation analysis performed according to T30?
(c)	Sample allowed to cool to room temperature in sample baskets?
(0)	
	AASHTO Materials Reference Laboratory
Report	(ASTM only)
(a)	Mass of HMA sample before and after ignition (to nearest 0.1 g) included on report?

PREPARING HMA SPECIMENS BY MEANS OF THE SUPERPAVE GYRATORY COMPACTOR

(T312)	
(D6925)	

			<u>APPARATUS</u>	Date:
Gyrator	v Com	pactor		
(a)		facturer:		
(b)	AASH	TO: Capable of applying of	a pressure of 600 ± 18 kPa?	
· /	ASTM	I: Capable of applying a c	constant vertical pressure of 600.	
(c)				of the compactor?
(d)				
(e)	Recor	ds height of specimen to 0.	1 mm during compaction once pe	r gyration [AASHTO: if density is
(f)	Applie			_
` /	(1)		gle of 20.2 ± 0.35 mrad (1.16 ± 0.0))2 degrees)?
or	(2)		•)2 degrees)?
Balance		TO. C5 halamas (usadah)		
	ASTN.			
		1: Minimum capacity of 1	0,000 grams with a sensitivity of	0.1 grams?
Ovens -	forced	d: Minimum capacity of I draft oven capable of bein vens recommended:	$0,000$ grams with a sensitivity of ag thermostatically controlled to \pm	3°C?
Ovens - ASTM: One for	– forced Two o	d: Minimum capacity of I draft oven capable of bein vens recommended: ift oven capable of mainta	$0,000$ grams with a sensitivity of \pm ining the temperature required, μ	3°C?
Ovens - ASTM: One for	– forced Two o	d: Minimum capacity of I draft oven capable of bein vens recommended: ift oven capable of mainta	$0,000$ grams with a sensitivity of \pm ining the temperature required, μ	3°C?
Ovens - ASTM: One for One oven	- forced Two o rced dra en shall	draft oven capable of being vens recommended: aft oven capable of maintal thave a range to a minimum. armored, glass or dial-types.	ining the temperature required, not of 204°C for heating aggregative with metal stems, range of at le	3°C?
Ovens - ASTM: One for One oven	- forced Two o rced dra en shall	draft oven capable of being vens recommended: aft oven capable of maintal thave a range to a minimum. armored, glass or dial-types.	ining the temperature required, not of 204°C for heating aggregative with metal stems, range of at le	3°C?
Ovens - ASTM: One for One oven	forced Two orced drawn shall ometers only: n	draft oven capable of being draft oven capable of being wens recommended: aft oven capable of maintal have a range to a minimular armored, glass or dial-typinimum sensitivity of 3 °C armored in accordance.	ining the temperature required, many of 204°C for heating aggregative with metal stems, range of at least ever with AASHTO TP 71 (at least ever)	3°C?
Ovens - ASTM: One for One oven	- forced Two o rced dra en shall ometers only: n	draft oven capable of being vens recommended: If oven capable of maintal have a range to a minimum sensitivity of 3 °C ASTHO: in accordance ASTM: in accordance	ining the temperature required, in most a sensitivity of the temperature required, in most 204°C for heating aggregation with metal stems, range of at least even with manufacturer's instructions.	0.1 grams?
Ovens - ASTM: One for One oven	- forced Two o reed dra en shall ometers only: n	draft oven capable of being vens recommended: If oven capable of maintage to a minimum sensitivity of 3 °C ASTM: in accordance the following been verified.	ining the temperature required, in most a sensitivity of the temperature required, in most 204°C for heating aggregation with metal stems, range of at least even with manufacturer's instructions.	3°C?

Gyration frequency?

LVDT (or other device to continuously record height)? Mold dimensions?

Plate faces?

Oven temperature?

Note, ASTM only: If manufacturer's instructions are not available these pieces of equipment should be checked at the following intervals: angle of gyration, vertical pressure, and height measurement system (monthly);

frequency of gyration (quarterly); mold and platen dimensions (annually).

COMMENTS (T312 / D6925):

(c) (d)

(e)

(f)

(g)

(T312 / D6925)

PREPARING HMA SPECIMENS BY MEANS OF THE SUPERPAVE GYRATORY COMPACTOR

(T312)	
(D6925)	

	PROCEDURE Date:
Prepara	tion of Apparatus
1.	Main power for compactor turned on for manufacturer's required warm-up period?
2.	Angle, pressure and number of gyrations set?
3.	Bearing surfaces lubricated as needed per manufacturer's instructions?
٥.	bearing surfaces tubricated as needed per manufacturer's instructions?
Dranara	tion of Mixtures (Laboratory mixed)
1 10para 1.	Aggregate fractions weighed into separate pan and combined to desired batch weight?
2.	If specimens are used for determination of volumetric properties, are the batch weights adjusted to result in a
۷.	compacted specimen 150 mm in diameter and 115 mm in height?
3.	Aggregate and binder placed in oven and heated to required mixing temperature?
3. 4.	ASTM only: Mixing implements heated to mixing temperature?
	Mixing and compacting temps. determined from a temperature-viscosity chart (T316/D4402)?
5.	
6.	Mixing temperature based on viscosity of 0.17 ± 0.02 Pa·s and compaction temperature based on viscosity of
-	0.28 ± 0.03 Pa·s [ASTM: temperatures based on viscosity of 0.17 ± 0.02 Pa·s]?
7.	Mixing bowl charged with heated aggregate and thoroughly dry-mixed?
8.	Crater formed in aggregate and binder added?
9.	Aggregate and binder mixed quickly and thoroughly?
10.	Mix placed in pan and aged [AASHTO only: in accordance with R30]?
11.	Mix kept in oven 2 hours ± 5 minutes at compaction temperature ± 3°C (Volumetric Design)?
12.	Mixture stirred every 60 ± 5 minutes to maintain uniform aging?
13.	AASHTO only: Mold and base plate preheated to compaction temperature for at least 30 minutes?
	ASTM only: Compaction mold assembly preheated to compaction temp. $\pm 5 \%$ for at least 45 minutes?
14.	Mix removed from oven and immediately compacted?
15.	AASHTO only: If the compaction temperature differs from the conditioning temperature used in accordance
	with R30, mix placed in oven at compaction temperature, maximum 30 min., to achieve the required temp.?
	AASHTO Materials Reference Laboratory
Prepara	tion of Mixtures (Plant sample)
1.	AASHTO: Loose mix brought to compaction temp. by uniform heating in an oven prior to molding?
	ASTM: For samples of as-produced mixture, follow one of the following short-term aging procedures:
	(a) No conditioning- compacted immediately as produced?
or	(b) Condition for $2h \pm 5$ min. at the compaction temperature ± 3 °C, stirring after 60 ± 5 min?
or	(c) Any conditioning method which has been demonstrated to replicate design conditioning?

COMMENTS (T312 / D6925):

(T312 / D6925)

PREPARING HMA SPECIMENS BY MEANS OF THE SUPERPAVE GYRATORY COMPACTOR

(T312)	
(D6925)	

	PROCEDURE (Continued) Date:
~	
1. 2. 3.	Mold, base plate, and upper plate (if required) removed from oven and paper disk placed on bottom of mold? Mixture placed into mold in one lift, mix leveled, and paper disk and upper plate (if required) placed on top of material [ASTM only: Quickly place mix into mold using a transfer bowl or other suitable device]? Mold loaded into compactor and a pressure of 600 ± 18 kPa applied?
4. or	Internal angle of 20.2 ± 0.35 mrad (1.16 ± 0.02 degrees) applied to the mold and compaction started?
5.	AASHTO: Compactor shuts off when desired number of gyrations are reached?
	ASTM: Compaction shall proceed until the desired endpoint – either a required number of gyrations (volumetric properties), or a specified height (physical property testing)?
6.	AASHTO only: No leveling off load applied (dwell gyrations, reverse gyrations, or square load), unless specified in another standard referencing T312?
	Note to Assessors: The following is guidance for certain models of gyratory compactors. If a delay is being applied a counter will often be displayed counting down [Reference: LAP technical bulletin 1-08].
	Pine Model AFGB Compactor - requires a dwell setting of "2" for no delay. This does not mean a 2-second delay. Pine Models AFGI & AFGC125X – may indicate "Compaction Complete, Squaring specimen, please wait" but this is ok. The ram will retract immediately after the angle is removed, which is correct.
7. 8. 9.	Mold removed and specimen extruded (may require cooling time before extruding)?
Density 1. 2. 3.	Procedure Maximum specific gravity (T209/D2041) determined on companion sample aged to same extent? Bulk specific gravity (T166/D2726 or T275/D1188) of specimen determined? Height recorded to nearest 0.1 mm after each revolution (when monitored)?
$G_{mbx} = G$	ssessors: the relative density at any given gyration of interest can be determined as follows $_{\text{mbfinal}}$ (h_{final} / h_{x}) (1) (G_{mbx} / G_{mm}) * 100 (2)
$G_{mm} = m$ $h_{final} = he$ $h_{x} = heig$ $G_{mbfinal} = % G_{mm} = % G_{$	alk specific gravity of the extruded specimen, at any gyration, x aximum theoretical specific gravity of the mixture (companion sample) ight of the specimen recorded at the final gyration, mm ht of the specimen recorded at any gyration, x, during the compaction process, mm bulk specific gravity of the extruded specimen at the final gyration relative density at any gyration, x, expressed as a percentage of the maximum theoretical specific gravity, arest 0.1 % at selected number of gyrations.

COMMENTS (T312 / D6925):

(T312 / D6925)

HAMBURG WHEEL-TRACK TESTING OF COMPACTED HMA

(T324)

		APPARATUS Date:					
1.	Uaml	burg Wheel-Tracking Machine					
1.	(a)	Electrically powered machine capable of moving a 203.2-mm (8-in.) diameter, 47-mm (1.85-in.)					
	. ,	wide steel wheel over a test specimen?					
	(b)	The load on the wheel is $705 \pm 4.5 \text{ N}$ (158 lb \pm 1.0 lb) while traveling back and forth across the					
	()	specimen?					
	(c) (d)	Wheel makes approximately 50 passes across the specimen per minute?					
	(u)	of the specimen?					
2.	Temp	perature Control System					
	(a)	Water bath controlled to within ± 1.0°C (1.8°F) over a range of 25 to 70°C (77 to 158°F)?					
	(b)	Water bath has a mechanical circulating system to stabilize the temperature in the specimen tank?					
3.	<u>Impre</u>	ession Measurement System					
	(a)	An LVDT device capable of measuring the depth of the impression of the wheel within 0.01 mm (0.0004 in.), over a minimum range of 0 to 20 mm (0.8 in.)?					
	(b)	System mounted to measure the depth of the impression at the midpoint of the wheel's path on the slab specimen?					
	(c)	Impression measured at least every 400 passes of the wheel?					
	(d)	System capable of measuring rut depth without stopping the wheel?					
4.	Whee	Wheel Pass Counter					
	(a)	Non-contacting solenoid that counts each wheel pass over the specimen?					
	(b)	Data from signal of counter coupled to wheel impression measurement, allowing for rut depth to b expressed as a function of wheel passes?	e 				
5.	Speci	imen Mounting System					
	(a)	Stainless steel tray that can be mounted rigidly to the machine?					
	(b)	Mounting restricts shifting of the specimen to within 0.5 mm (0.02 in.) during testing?					
	(c)	System suspends the specimen, allowing for free circulation of the water bath on all sides?					
	(d)	Minimum of 20 mm (0.8 in.) of free circulating water on all sides of the specimen?					
6.	<u>Balar</u>	nce, capacity of 12,000 g, accurate to 0.1 g?					
7.	Oven	1 - for heating aggregate and asphalt binders?					
8.		rpave Gyratory Compactor and molds, conforming to AASHTO Test Method T312?					
	Note i	to Assessors: this is only needed if the laboratory is testing gyratory specimens in the wheel tester.					
9.		er of Paris - mixed at approximately a 1:1 ratio of plaster to water?					
0	r <u>Plasti</u>	ic mounting sheets?					
COM	MENTS	(T324):	(T324)				

HAMBURG WHEEL-TRACK TESTING OF COMPACTED HMA

(T324)

			<u>PROCEDURE</u>	Date:
O-134	:/E	Varionalian (Ontional)		
		Verification (Optional)		
1.				ure readout every 6 months?
2.			mm (0.002 in.) between the three (
3.	Wheel force ve	rified, mid-stroke, at the	e correct level elevation, to be 705	± 4.5 N (158 ± 1.0 lb)?
			accurate to $0.4N(0.1lb)$ is sufficient t	
4.	Steel wheel rec	iprocating back and forth	h on the test sample at 50 ± 5 pass	ses per minute verified?
Specime	en Preparation			
1.	At least two spe	ecimens prepared for each	ch test?	
2.	Slab specimens	or cylinders?		<u>—</u>
	-	•		
	Circle One:	Slab Specimen	Core or Gyratory Specimen	1
3.	Either laborator	y-produced HMA or fie	eld-produced HMA?	
Laborat	ory-Produced HI	<u>MA:</u>		
	mixin	g temp. range:t	to compaction temp. rai	nge: to ula?
1.	Mixture propor	tions batched in accorda	ince with the desired job-mix form	ula?
2.	Mixing temp. is	the temperature the asp	phalt binder must be heated to achi	leve a viscosity of 170 ± 20cSt?
3.	Mixing tempera	nture recommended by the	he manufacturer used for modified	l binders?
4.	Aggregates and	mineral admixture (if u	sed) dry-mixed?	
5.				e thoroughly coated?
			vet-mixed if a lime slurry or other wet	
6.	Test sample con	nditioned at the appropri	iate compaction temperature in acc	cordance with the short-term
	conditioning pr	oce <mark>du</mark> re in R 30?		
7.	Compaction ter	np. is the temperature th	e asphalt binder must be heated to	achieve a viscosity of 280 ± 30cSt?.
8.				y the manufacturer used?
Laborat	ory Compaction	of Specimens – Slah Sp	ecimens erials Refero	ence Laboratory
1.	Compacted into	of Specificis — Siab Sp odaho ucing a Linear Kr	neading Compactor (or equivalent)?
2.	Specimens are	320 mm (12.5 in) long a	and 260 mm (10.25 in) wide?	
3.	Slab thickness i	s between 38 mm (1.5 ii	n) and 100 mm (4 in)?	·····
<i>3</i> . 4.	Slab thickness i	is at least twice the nomi	inal maximum aggregate size?	·····
4. 5.	Compacted spe	oiman goolad at normal	room temperature on a clean, flat	surface until the specimen
3.				
	is cool to the to	ucii?		
Laborat	ory Compaction	of Specimens – Gyrator	ry Specimens	
1.	Material compa	cted into specimens usir	ng a Gyratory compacter according	g to AASHTO T312?
2.	Specimen thick	ness is between 38 mm ((1.5 in.) and 100 mm (4 in.)?	
3.	Specimen thick	ness is at least twice the	nominal maximum aggregate size	e?
4.	Two specimens	prepared?		<u> </u>
5.	Compacted spe	cimen cooled at normal	room temperature on a clean, flat	surface until the specimen
COMM	ENTS (T324):			(T324)

COMMENTS (T324):

HAMBURG WHEEL-TRACK TESTING OF COMPACTED HMA

(T324)

		PROCEDURE (Continued) Date:	
	re / S	Slab Specimens	
1.		Specimens are wet saw-cut compacted specimens taken from HMA pavements?	
2.		Specimen size:	
		(a) Field core is 250 mm (10 in.) in diameter and may be 38 mm (1.5 in.) high?	
	or	(b) Cut slab specimen is approximately 260 mm (10.25 in.) wide and 320 mm (12.5 in.) long and	
		38 mm (1.5 in. high)?	
_			
	termi	ining air void content	
1.		Bulk specific gravity determined in accordance with AASHTO T166?	
2.		Maximum specific gravity of the mixture determined in accordance with AASHTO T209?	
3.		Air void of the specimens determined in accordance with AASHTO T269?	
4.		Air voids:	
		(a) Air void of laboratory-compacted specimens is 7.0 ± 2.0 percent?	
	or	(b) Field specimens (cores / cut slabs) tested at the air void content at which they were obtained?	
_			
	cedu		
1.		Plaster of Paris used to mount the specimens in the mounting trays?	
2.		Plaster poured to a height equal to that of the specimen so that the air space between the specimen and	
_		the tray is filled?	
3.		Plaster layer underneath the specimen does not exceed 2 mm (0.08 in.)?	
4.		Plaster set for at least 1 hour?	
_		Note: If other mounting material is used, it should be able to withstand 890 N (200 lb) of load without cracking.	
5.		Test temperature selected based upon the applicable specifications?	
6.		Drain valves closed and wheel-tracking device filled with hot water until the float device floats	
7		to a horizontal position?	
7.		After the water has reached the test temperature for 30 minutes, wheels lowered onto the specimens?	
8.		Micro-control unit's LVDT reads between 10 mm (0.4 in.) and 18 mm (0.7 in.) and test started?	
9.		Wheel-tracking device shut off when (a) 20,000 passes have occurred or (b) if the average LVDT	
1.0		displacement is 40.90 mm (1.6 in.) or greater for a specimen?	
10.		Screen readout subtracts the initial LVDT reading from the total displacement?	
11.		Machine and the main power supply turned off?	
12.		Valves opened to drain the bath, and the wheels raised and the rutted specimens and spacers removed?	
13.		Water baths, heating coils, wheels, and temperature probe cleaned with water and scouring pads?	
14.		Wet-dry vacuum used to remove particles that have settled to the bottom of the baths?	
15.		Filter element and spacers cleaned after every test?	
16.		Calculations performed according to the test method?	
D ~-	aart		
<u>Ke</u> j 1.	<u>oort</u>	UMA production (field or lab) and composition method wood?	
		HMA production (field or lab) and compaction method used?	
2.			
3. 1		Test temperature?	
4. 5		Specimen air voids? Type and amount of anti-stripping additive?	
5. 6.		Type and amount of anti-stripping additive?	
υ.		Creep stope, surp stope, and surpping infrection point!	

(T324)

MOISTURE CONTENT OF HOT-MIX ASPHALT (HMA) BY OVEN METHOD (T329)

	APPARATUS Date:
1.	Oven, maintains 163 ± 14°C (325 ± 25°F)?
2.	Sample container, of sufficient size to contain the sample without danger of spilling?
3.	Balance, 2-kg (4.4-lb) capacity, readable to at least 0.1 g?
	<u>PROCEDURE</u>
1. 2.	Test sample obtained by AASHTO T248, Method B (quartering)?
3.	Mass of the sample container determined to the nearest 0.1 g?
4.	Sample placed into the container, distributed evenly, and the temperature of the test sample determined?
5.	Mass of the sample container and moist test sample determined to the nearest 0.1 g?
6.	Mass of the moist test sample determined by subtracting the mass of the sample container from the
7.	total mass of the sample container and moist test sample? (M_i)
8.	If a mixing temperature range is not supplied, dried at $163 \pm 14^{\circ}$ C $(325 \pm 25^{\circ}F)$?
o. 9.	Sample initially dried for 90 ± 5 minutes and mass determined?
10.	Sample then dried at 30 ± 5 minutes and mass determined? Sample then dried at 30 ± 5 minute intervals until a constant mass is reached?
11.	Test sample removed from oven and cooled to approximately the same temperature determined in Step 4?
12.	Mass of the sample container and dry test sample determined to the nearest 0.1 g?
13.	Mass of the final dry test sample determined by subtracting the mass of the sample container from the
	total mass of the sample container and dry test sample? (M_f)
14.	Moisture content calculated according to Section 7.1.1 or 7.1.2 (see below)?
For AC	content reported as % of HMA: % moisture = $M_i - M_f$ Where: $M_i = \underline{\underline{B}}$ Efore drying (moist mass) $M_f = \underline{\underline{M}}$ fter drying (dry mass)

COMMENTS (T329): (T329)

BULK SPECIFIC GRAVITY OF COMPACTED HMA USING AUTOMATIC VACUUM-SEALING METHOD

(T331)	
(D6752)	

Date: _____

<u>APPARATUS</u>

1.	Balance	
	(a)	AASHTO: Readable to 0.1 percent of the sample mass or better, conforming to M231?
		figures (0.1 g for 130.0 to 999.9 g) (meets D4753, GP2)?
	(b)	Equipped with an apparatus for weighing specimen while suspended in water?
2.	Water E	Bath
	(a)	Minimum dimensions of 610 x 460 x 460 mm (24 x 18 x 18 in.) or a large cylindrical
	()	container capable of completely submerging the specimen while suspended?
	(b)	Equipped with an overflow outlet to maintain constant water level?
	(c)	AASHTO: Suspension wire of smallest practical size?
	()	ASTM: Has a cushioned specimen holder, without sharp edges, for submerging samples?
3.	Vacuum	n Chamber
	(a)	Equipped with a pump capable of evacuating a sealed and enclosed chamber to a minimum
	()	pressure of 5 mm Hg absolute [ASTM: 10 mm Hg] in less than 60 seconds at sea level?
	(b)	Chamber large enough to seal samples with dimensions of 150 x 350 x 150mm (6 x 14 x 6 in.)?
	(c)	Automatically seals bag?
	(d)	Exhausts air back into chamber in a controlled manner to ensure plastic conforms to specimen?
	(e)	Air exhaust and vacuum operation time calibrated at factory prior to initial use?
	(f)	Air exhaust system calibrated to bring chamber to atmospheric pressure in 80 to 120 seconds,
	(1)	after the completion of the vacuum operation?
	(g)	Vacuum system provided with a latch to control the chamber door opening?
4.	Vacuun	n Measurement Gauge
т.	(a)	Independent from the vacuum sealing device?
	(b)	AASHTO: Calibrated gage capable of reading 1 mm Hg (1 torr) pressure with a minimum
	(0)	range of 10 to 0 mm Ha (10 to 0 torr)?
		range of 10 to 0 mm Hg (10 to 0 torr)?
5.	Plastic 1	
٥.		One of the two following sizes:
	(a)	(1) Smaller bags: Openings from 235 to 260 mm (9.25 to 10.25 in.)
		[ASTM only: 241 mm to 260 mm (9.5 to 10.25 in.)]?
	(b)	(2) Larger bags: Openings from 375 to 394 mm (14.75 to 15.5 in.)?
	(b)	Does not adhere to asphalt film? Capable of withstanding temperatures of up to 70°C (158°F)?
	(c)	Capable of withstanding temperatures of up to 70 C (138 r)?
	(d)	Impermeable to water and is puncture resistant?
	(e)	Contains no air channels for evacuation of air from bag?
	(f)	Thickness of bags 0.100 to 0.152 mm (0.004 to 0.006 in.)?
	(a)	Specific gravity known [AASHTO: provided by manufacturer]?
	(g)	Specific gravity known [AASITIO. provided by manujacturer]:
6.		nal Apparatus
	(a)	Specimen sliding plate with a smooth, flat surface?
	(b)	Bag cutting knife, scissors, or other type of clipping device?
	(c)	ASTM only: For bag density verification, sufficient aggregate and AC to prepare a lab-compacted sample of 4.75 mm design mixture with min. dimensions of 100 mm diameter by 60 mm thick?
	(d)	AASHTO only: Drying oven?
	(e)	AASHTO only: Thermometer, ASTM 17C (range 19 to 27 °C, subdivisions of 0.1 °C)?
	or	AASHTO only: Thermometer, ASTM 17F (range 66 to 80 °F, subdivisions of 0.2 °F)?
		AASHTO only: An electronic temperature measuring device?

COMMENTS (T331 / D6752):

(T331 / D6752)

BULK SPECIFIC GRAVITY OF COMPACTED HMA USING AUTOMATIC VACUUM-SEALING METHOD

(T331)	
(D6752)	

			PROCEDURE	Date:
Sai	mnlin	g and Te	est Specimens	
1.	шртт		f specimens:	
		(a)	Laboratory prepared specimens?	
	or	(b)	Field samples [ASTM only: obtained in accordance with Practice L	
2.			ter of cylindrically molded or cored specimens, or the length of the side our times the maximum aggregate size?	s of sawed specimens at
3.		Thickne	less of specimens at least 1 ½ times the maximum aggregate size?	
4.			ken to avoid distortion, bending, or cracking of specimen and stored in	
5.			TO: Sample conforms to the requirements of T166?	
<u>Ve</u>	rifica			
1.		-	Verification	
		(a)	Vacuum settings of the device verified every three months, after repa each shipment or relocation?	
		(b)	Verification performed with an absolute vacuum gage capable of beir	
			chamber and reading the vacuum setting of the sealing device?	
		(c)	Vacuum gage indicates a reading of 10 mm Hg (10 torr) or less?	
		(d)	ASTM only: Vacuum gage used for verification calibrated every 12	? months?
2.		Plastic	Bag Verification	
		(a)	Plastic bag apparent specific gravity provided by manufacturer verific	ed periodically?
		(b)	Laboratory compacted sample used to verify bags?	
			(1) 4.75 mm mixture compacted by Marshall compactor or Gyra	
			(2) Minimum sample diameter of 100 mm by 60 mm thick?	
			(3) AASHTO: Sample compacted to produce air voids in the rar	
			ASTM: Sample compacted to air voids in the range of 4%	
		(c)	Average of results from three bags (per size) used to measure the bull	
		(d)	Bulk specific gravity of the same sample determined using (T166 / D	
		(e)	AASHTO: Average bulk spec. gravity calculated for the lab-compact	
			$(\pm 0.020 \text{ g/cm}^3)$ of the bulk spec. gravity determined by T166 for the s	
			ASTM: Average bulk spec. gravity calculated for the lab-compacted	
			$(\pm 0.010 \text{ g/cm}^3)$ of the bulk spec. gravity determined by D2726 for the	
		(f)	AASHTO only: If difference between T166 and T331 bulk spec. gravi	
			dried and verification repeated?	
		(g)	AASHTO only: Manufacturer contacted if second test fails?	

COMMENTS (T331 / D6752):

(T331 / D6752)

BULK SPECIFIC GRAVITY OF COMPACTED HMA USING AUTOMATIC VACUUM-SEALING METHOD

(T331)	
(D6752)	

Date: _____

PROCEDURE (Continued)

Tes	ting			
1.		Mass in air determination:		
		(a) Laboratory-prepared, dry specimens mass determined at room temperature? (A)		
	or	(b) Cores and specimens containing moisture mass determined after drying to constant mass,		
	01	[AASHTO: at 52 ± 3 °C (125 ± 5 °F)] or dried by D7227 (vacuum drying) less than		
		0.05% change between consecutive 15 minute drying intervals? (A)		
2.		ASTM only: Anything that would prevent the bag from complying with the sample removed by sawing?		
3.		Appropriate size bag selected and mass determined [AASHTO only: and inspected for holes or damage?]?		
٥.				
		(b) Small bag used for 150 mm (6 in.) diameter specimens with a thickness less than 50 mm (2 in.)		
		[ASTM only: thickness less than 75 mm (3 in.)]?		
		(c) Large bag used for 150 mm (6 in.) diameter specimens with a thickness greater than 50 mm (2 in.)		
		[ASTM only: thickness greater than 75 mm (3 in.)]?		
,		Note: Use manufacturer's recommendation for specimens weighing more than 5500 g or abnormally shaped.		
<i>4</i> .		AASHTO only: If needed, filler plates added or removed before inserting specimen?		
5.		Bag placed on top of specimen sliding plate inside vacuum chamber?		
6.		AASHTO: Specimen placed in bag, smoothest side down?		
7		ASTM: Specimen placed in bag without puncturing, dropping, or impacting the bag?		
7.		AASHTO only: End of bag pulled over the sample, centered over the sealing bar with at least 1 in. overlap?		
8.		Any wrinkles in the bag straightened just prior to closing the lid and latching the bar?		
9.		Chamber door latched to avoid automatic opening of door after the completion of the test?		
10.		Vacuum chamber allowed to remove air from chamber and bag and automatically seal bag?		
11.		Exhaust air into chamber until chamber door opens indicating atmospheric pressure within the chamber?		
12.		Sealed sample removed from the vacuum chamber without puncturing bag?		
13.		AASHTO: Mass of sealed specimen in air determined? (B)		
		ASTM: Sample immediately placed in water bath at $25 \pm 1^{\circ}C$ (77 ± 1.8°F) and mass determined?		
14.		AASHTO only: Specimen fully submerged in bath and no air bubbles entrapped under specimen?		
<i>15</i> .		AASHTO only: Mass of sealed specimen in a water bath at 25 ± 1 °C (77 ± 1.8 °F) determined? (E)		
		Note: The time between the lid opening and putting the specimen in the water bath should not exceed 1 min.		
<i>16</i> .	ASTM only: If temperature differs from specified range, is a correction to the bulk specific gravity to			
		25°C made in accordance with Section 8.3?		
17.		Sample removed from bag and mass determined? (C)		
18.		Specimen's new mass in air (C) checked against the initial mass (A) and if the check fails is test restarted?		
		(a) AASHTO: check passes if less than 0.08 % is lost or no more than 0.04 % is gained?		
		(b) ASTM: check passes if dry mass after test < (the initial dry mass + 5 g)?		
19.		AASHTO only: Specimen oven or vacuum dried to constant mass prior to recording specimen dry mass?		
<i>20</i> .		AASHTO only: Specimen must be vacuum dried for referee testing?		
<i>21</i> .		ASTM only: Specimen dry mass after procedure recorded?		
22.		Bulk specific gravity calculated according to the method to four significant figures (see formula below)?		
		Note to Assessors: The formula given below is the ASTM version. The AASHTO version is slightly different and uses		
		different letters to designate the values. A = initial dry mass of specimen, g		
		A B = mass of sealed specimen in air, g		
		$G_{mb} =$ $G_{mb} =$ $[C+(B-A)] - E - [(B-A)/F]$ $G_{mb} =$ $[C+(B-A)] - E - [(B-A)/F]$ $G_{mb} =$		
		[C+(B-A)] - E - [(B-A)/F] E = mass of sealed specimen in water at 25°C, g		
		$F = $ spec. gravity of bag to nearest 0.001 $G_{mb} = $ specimen bulk specific gravity		
23.		Density of the specimen calculated and reported to nearest 0.001 (see formula below)?		
<i>23</i> .		$G_{mb} = $ specimen bulk specific gravity		
		$\tilde{n} = G_{mb} * \tilde{a}$ specified bulk specific gravity $\tilde{a} = density \text{ of water at } 25^{\circ}\text{C } (77^{\circ}\text{F}), 0.999 \text{ g/cm}^{3}$		
		$\tilde{n} = \frac{\text{density of water at 25 C (77 T), 6.599 g/cm}}{\text{density of specimen kg/m}^3 (lb/ft^3)}$		
CO	MM	ENTS (T331 / D6752): (T331 / D6752)		

EFFECT OF WATER ON COHESION OF COMPACTED HMA

((D1075)	

			<u>APPARATUS</u>	Date:
1.		Water B (a) (b)	Sufficient size to permit total immersion of 3 specimens? Temperature: 25 ± 1°C (77.0 ± 1.8°F)?	
2.		Water B (a) (b) (c) (d) (e)	Automatic temperature control, capable of controlling temperature to ± 1°C (± Lined with copper, stainless steel, or other non-reactive material?	
3.		Air Bath	$\underline{\text{n}}$ capable of being maintained at $25 \pm 1^{\circ}\text{C}$ (77.0 $\pm 1.8^{\circ}\text{F}$) for 4 hours?	
4.	or	Transfer (a) (b)	Plates Flat; glass, metal, or other non-reactive material; at least 3 available? One large transfer plate sufficient to hold all three specimens?	
5.		Compre	ssive Strength Testing Machine available [required for test method T167 (D1074	4)]?
6.		Apparat	us for T166, Method A / D2726, Method A?	
СО	MM	ENTS (D	01075):	(D1075)

EFFECT OF WATER ON COHESION OF COMPACTED HMA

(T	107	<i>E</i> \	
	107	つ 1	

		PROCE	<u>EDURE</u>	Date:	
Toot Sn	ecimens				
1.		x 102 mm diameter specimens prepare	ed in accordance with (T167/D10	74)?	
Determi	nation of Bulk Sr	pecific Gravity (sequence of steps opt	ional)		
1.		ved to cool at least 2 hrs. after remova			
2.	Oven-dry mass of	of each specimen determined?			
3.	Each specimen i	mmersed for 3 to 5 minutes in water a	at 25 ± 1°C?		
4.	Immersed mass	of specimen recorded?			
5.	Blotted quickly	with damp towel?			
6.		s of each specimen determined?			
7.	Bulk specific gra	avity of each specimen calculated?		······	
Procedu	ıre				
1.		pecimens separated into groups of thr	ee specimens each so that the aver	rage	
		wity of Group 1 is essentially the sam			
2.		ept under each specimen during imme			
		sting?			
3.	Test specimens	n each group tested as follows:			
	Group 1				
	(a) Specim	ens stored in air bath at 77.0 ± 1.8 °F ($(25 \pm 1^{\circ}\text{C})$ for at least 4 hours?		
	(b) Compre	essive strength of each Group 1 specir	nen by (T167 / D1074)?		
	Group 2		,		
		ens on transfer plates immersed in wa	ter at 140.0 ± 1.8 °F (60 ± 1 °C) fo	r 24 hrs?	
	(b) Specim	ens transferred to water bath at 77.0 ±	1.8° F (25 ± 1°C) for 2 hrs?		
	(c) Compre	essive strength of each Group 2 specir	men by (T167 / D1074)?		
	Group 2, Alterna				
		ens on transfer plates immersed in wa	ter at 120.0 ± 1.8 °F (49 ± 1 °C) fo	r 4 days?	
		ens transferred to water bath at 77.0 ±			
	(°)	essive strength of each Group 2 specin	als Reference La	aboratory —	
Calcula					
Numeri	cal index of resist	ance to water calculated according to	equation below?	·······························	
		Index of retained streng	$\text{tth, } \% = (S_2 / S_1) \times 100$		
	Where:	S_1 = Compressive strength of dry sp	ecimens (Group 1)		
		S_2 = Compressive strength of immer			
COMM	ENTS (D1075):			(D1075)	

DENSITY OF BITUMINOUS CONCRETE IN PLACE BY NUCLEAR METHODS (D2950)

	<u>APPARATUS</u>	Date:
Nuclear 1. 2. 3.	Device: An electronic counting instrument capable of being seated on the surface of the mate A sealed high energy gamma source such as cesium or radium? Equipped with a gamma detector such as a Geiger-Mueller tube?	
Referen	ce Standard Block: A block of dense material able to produce reference count rates?	
1. 2.	A metal plate, straightedge, or suitable leveling tool capable of conditioning the test smoothness? Drive Pin: (a) A steel rod of slightly larger diameter than the rod in the direct transmission (b) A drill capable of creating a perpendicular hole?	instrument?
	CALIBRATION	
1. 2.	New gages initially calibrated? Existing gages calibrated to re-establish calibration curves, tables, or equivalent coeff once each year and after all major repairs? Calibrations completed in accordance with manufacturer's recommended procedures Calibration produces calibration response within ± 16 kg/m³ (± 1.0 lb/ft³) on standard established and constant densities (can be done by manufacturer, user, or independen Note: one-point calibration is not acceptable. The densities of the materials used to establish or verify the calibration extend through include the types and densities on the in-place materials to be tested? Blocks used to establish calibration identified on the calibration data sheets? STANDARDIZATION	ficients at least ?
Records 1. 2.	Standardization performed at the start of each day's work? Permanent records of this data retained?	
Location 1. 2.	n: Standardization performed with apparatus located at least 8 m (25 ft) away from other Area clear of large mass or other items that could affect the reference count?	
 2. 3. 	Device turned on prior to standardization and allowed to stabilize?	esults.

DENSITY OF BITUMINOUS CONCRETE IN PLACE BY NUCLEAR METHODS (D2950)

	STANDARDIZATION (Continued) Date:
4.	Are current day count ratios determined using the following equation?
	$\begin{aligned} \left N_s - N_o \right &\leq 2.0 \sqrt{N_o} F \\ N_s &= & \text{value of current standard count} \\ N_o &= & \text{average of past four values of N}_s \text{ taken} \\ &= & \text{previously} \\ F &= & \text{value of any pre-scale (a divisor supplied by the} \\ &= & \text{manufacturer)} \end{aligned}$
5. 6. 7.	If the value is outside of limits, is the device allowed additional stabilization time before repeating procedure? If the device fails a second time, is the device adjusted or repaired as recommended by the manufacturer? If measured densities become suspect, is another standardization count performed?
	<u>PROCEDURE</u>
Nuclear 1. 2.	Device: Instrument turned on prior to use and allowed to stabilize? Power left on during the entire day's testing?
Site Pres 1. 2. 3. 4.	If the instrument will be closer than 250 mm (10 in.) to any vertical mass, manufacturer's correction procedure followed? Test site leveled with the guide / scraper plate?
Backsca 1. 2. 3.	Device placed on prepared test site?
Direct T 1. 2. 3. 4. 5. 6.	Guide / scraper plate placed on test site?
<u>Calculat</u> 1. 2. 3.	In-place density (bulk or wet density) determined from the ratio and the calibration and adjustment data? Note: Some instruments may have built-in provisions to compute the ratio and density. Otherwise, calibration charts, tables, equations, or coefficients may be used. Adjustment bias may be calculated by comparing the results from instrument measurements to results D2726 (Bulk Specific Gravity of Bituminous Mixtures)?
COMM	ENTS (D2950): (D2950)

COMMENTS (D4013):

PREPARATION OF TEST SPECIMENS OF BITUMINOUS MIXTURES BY MEANS OF GYRATORY SHEAR COMPACTOR

(D4013)

	<u>APPARATUS</u>				Date:						
Curo	tory Shoor Compactor										
(a)	Gyratory Shear Compactor (a) Platen and ram face ground flat?										
(b)	Low pressure gauge with resolution of ± 0.3 psi (± 2.0 kPa) ca					•••••	•••••				
(0)	(1) Pregyration stress of 31.8 psi (219 kPa) (400 lbf tot	al on 4	-in. sr	ecime	ь. n)?						
	(2) End point stress of 95.3 psi (657 kPa) (1200 lbf total on 4-in. specimen)?										
(c)	High pressure gauge with resolution of ± 16.0 psi (± 110 kPa) capable of displaying:										
	(1) Consolidation stress of 1590 psi (11.0 MPa) (20000 lbf total on 4-in. specimen)?										
(d)	(d) Tilt mechanism capable of cocking the mechanism 6 degrees while under pre-gyration stress?										
(e)	Gyration mechanism driving at a rate of 1 revolution per sec										
(f)	Metered hydraulic pump (applies 0.020-in. (0.508-mm) of ra	m mov	emen	t per a	pplicat	ion)?					
Chec	k the following items at room temperature:										
Mol	ds, Ram Faces, and Base Plates		1	2	3	4	5	6			
	e diameter of ram: 3.995 - 4.000 in. (101.47 - 101.60 mr	n.) A									
Face	e diameter of base plate: 3.995 - 4.000 in. (101.47 - 101.60 mr										
	ekness of base plate rim: $0.052 - 0.072$ in. $(1.32 - 1.82 \text{ m})$										
Thic	ekness of base plate: $0.55 - 0.58$ in. $(13.89 - 14.65 \text{ m})$	m) E									
Insid	de diameter of mold: 3.96 - 4.06 in. (100.58 – 103.12 m										
Heig	ght of mold: 3.87 – 4.00 in. (98.30 mm - 101.60 m										
Balar	Balance: Minimum capacity of 4500g with a sensitivity of 0.1g?										
Sieve	e or screen: A 1-in. (25-mm) screen (round openings) or a 7/8 in.	(22.4-	·mm)	sieve (square	openin	gs)?				
<u>Spatı</u>	Spatula: Blade about 4 in. (100 mm) long and 3/4 in. (20 mm) wide?										
Spoo	Spoon: Bent to a right angle between bowl and handle?										
<u>5000</u>	m. Dent to a right angle between bowr and handle:	•••••	•••••	•••••	•••••	•••••	•••••				
Meas	suring device: Caliper or micrometer for measuring the height of	the sa	mple?	?	•••••		•••••				
Spec	Specimen Extrusion Device: (Any means which does not distort or damage the specimen is acceptable)?										
Over	Oven: Range of 100 to 300°F (37.8 to 148.9°C) thermostatically controlled to \pm 5°F (\pm 3°C)?										
Misc	Miscellaneous: Gloves, trowels, mixing pans, and thermometers?										
Kero	sene?										
	Paper Disks, appropriate in diameter to the mold being used?										

(D4013)

PREPARATION OF TEST SPECIMENS OF BITUMINOUS MIXTURES BY MEANS OF GYRATORY SHEAR COMPACTOR

(D4013)

	PROCEDURE Date:	
C-13.	and an and Marie Conditions	
	ration and Verification:	
1.	Low- and high-pressure gauges calibrated to indicate distinct points for:	
	(a) Pre-gyration stress? (low pressure gauge)	
	(b) End point stress? (low pressure gauge)	
	(c) Consolidation stress? (high pressure gauge)	
	pressure applied to the ram. In any of these cases, verify that the appropriate pressure is applied	
	to the specimen at each of the calibration points.	
2.	Low- and high-pressure gauges verified on the gyratory shear molding press at each of the calibrated poi	nts?
Sampl	ole and Apparatus Preparation:	
1.	Sample mass required to produce a specimen conforming to requirements calculated by Section 7?	
2.	If mixing, HMA samples mixed at $250 \pm 5^{\circ}F$ ($121 \pm 3^{\circ}C$)?	
3.	Samples cured to constant mass in an oven at $140 \pm 10^{\circ}$ F ($60 \pm 6^{\circ}$ C)?	
4.	Samples stirred throughout curing?	
5.	If the sample contains material larger than 7/8 in., material separated over a 7/8-in. (22.4-mm) sieve, or	
	through a 1-in. (25.0-mm) round opening screen, scraping material to ensure recovery of fines clinging	
	to the surface of larger particles?	
6.	Molds and base plates preheated to approximately 100°F (38°C)?	
7.	Gyratory apparatus in loading position?	
8.	Apparatus run through at least one gyration prior to testing?	
9.	Platen, lower bearing, and top of the mold ring lubricated?	
10.	Mold and base plate removed from oven and insides wiped down with either kerosene or light lubricating	g oil?
11.	Paper disk placed in the bottom of the mold?	
Comp	paction:	
1.	Sample heated to $250 \pm 5^{\circ}F$ (121 ± 3°C) prior to compaction?	
2.	Mixtures funneled into the mold in two layers? After each lift, top leveled by pressing down with the spoon?	
3.	After each lift, top leveled by pressing down with the spoon?	::y
4.	Any large aggregates moved away from the walls?	<u></u>
5.	Paper disk placed on top of the material?	
6.	Mold placed onto the lower platen and centered beneath the press?	
7.	Ram pumped down until pre-gyration stress is reached?	
8.	Mold immediately tilted to the specified angle of gyration?	
9.	Handle of the pump lifted up to fill the metering pump?	
10.	Gyration mechanism switched on and mold gyrated three times?	
11.	Mold immediately squared (angle removed) and one full stroke of the metering pump performed?	
	Note: One stroke of the pump should take approximately one second.	
12.	Low pressure gauge reaches the end-point stress, otherwise pressure reduced below pre-gyration stress at steps 7 to 11 repeated until end-point stress is reached on one application of the metering pump?	
13.	Pump slowly applied until the low-pressure gauge is cut out of the system, then pressure pumped up to	
13.	consolidation stress (on the high-pressure gauge)?	
14.	Pumping stopped once consolidation stress is reached?	
15.	Pressure released slowly so as to prevent damage to the gauge?	
16.	Ram retracted and mold removed from the press?	
17.	Any method of removal that does not damage the specimen used to extract the sample from the mold?	
18.	Sample measured for height to confirm conformance to specification?	
19.	Mold cleaned with a kerosene rag if more specimens are to be molded?	
17.	The decided with a reference rag it more specimens are to be moracu.	
COM	IMENTS (D4013):	(D4013)

Revised 2011-03-25

RECOVERY OF ASPHALT FROM SOLUTION USING THE ROTAVAPOR

(D5404)

APPARATUS Date:

1.		Rotavapor apparatus						
		(a)	Distillation flask, depth of approximately 40 mm (1.5 in) when fully immersed?					
		(b)	Variable speed motor, capable of rotating the distillation flask at least 50 rpm?					
		(c)	Condenser?					
		(d)	Solvent recovery flask?					
		(e)	Heated oil bath?					
		(f)	Angle of distillation flask from horizontal to bath is approx. 15 degrees?					
2.			ge apparatus (either of the following):					
		(a)	Batch unit capable of 770g?					
		(b)	Centrifuge tubes (either of the following types):					
			(1) Wide-mouth bottles, 250 to 500 mL capacity?					
			(2) Cylindrical tubes, 6 or 8 in. long, with conical ends; capacity 100 mL?					
	or	(c)	Continuous unit capable of 3000 gravity?					
3.		Manome	eter or vacuum gage, suitable for measuring the specified vacuum?					
4.		Gas flov	v meter, capable of indicating a gas flow of up to 1000 mL/min.?					
5.		Sample container, having an adequate volume to hold the sample and added solvent?						
6.		<u>Vacuum system</u> , capable of maintaining a vacuum to within ± 0.7 kPa (± 5 mm Hg) of the desired level up to and including 80 kPa (600 mm Hg)?						
7.		Nitrogen gas or carbon dioxide gas?						
8.		Bath liqu	uid: USP White Oil? SHTO Materials Reference Laboratory					
	or	(a) (b)	Silicone Fluid SWS-101, with flash point above 215°C (420°F)?					
	or or	(b) (c)	Other?					
		(-)						
9.		Solvent						
		(a)	Reagent grade trichloroethylene or methylene chloride?					
			Note: Trichloroethylene, Technical Grade, Type I may be used, but it is recommended that for					
		(L)	each new supply of the solvent, a blank should be run. Normal Propyl Bromide (nPB)?					
	or	(b)	Normal Propyl Bromide (nPB)?					
			each new supply of the solvent, a blank should be run.					
10.		Solvents	s listed used only under a hood or with an effective exhaust system in a well ventilated area?					
СО	MM	ENTS (D	(D5404):					

RECOVERY OF ASPHALT FROM SOLUTION USING THE ROTAVAPOR

(D5404)

		PROCEDURE Date:						
Sama	le Preparati	tion:						
1.								
2.	Solution	Solution obtained by D2172, extraction method A?						
۷.	(a)	At least 30 min at 770 times gravity in either wide-mouth bottles or centrifuge	• ——					
	(a)	tubes in batch apparatus?						
	(b)	At least 3000x gravity in a continuous centrifuge charged at a rate not to exceed 150 mL/min?	• ——					
	(0)	At least 3000x gravity in a continuous centifuge charged at a fate not to exceed 130 mil/min?	•					
Proce	dure							
1.		th heated to 140 ± 3 °C $(285 \pm 5$ °F)?						
2.		vater circulated through condenser?						
3.	Vacuum	m of 40 mm below atmospheric pressure applied?						
4.	Approx.	x. 600 mL of AC solution drawn from sample container into flask through the sample line?						
5.	Approx.	x. 500 mL/min [AMRL: 50 ml/ min] (check records) flow of nitrogen or CO ₂ begun?						
6.	Distillat	ation flask rotated at approx. 40 rpm?						
7.	Flask lo	owered into the oil bath?						
8.	Steady,	controlled stream of condensed solvent maintained?						
9.	Gas flov	ow discontinued when the amount of solution in the flask is low and more is to be added?						
10.	Remain	Remaining asphalt solution drawn into flask and gas flow readjusted?						
	Note to A	Assessors: The equipment may be modified to allow a continuous flow of solution into the distillation flask -						
		time in the flask should be maintained at approx. 600 mL [AMRL: ± 50 mL/min]. The gas flow should not be						
		until all the solution has entered the flask.						
11.		mmersed to depth of 40 mm (1.5 in.) when most of the solvent has been distilled and no						
10	condens	nsation is occurring on the condenser?	·					
12.	Vacuum	m of 600 mm below atmospheric pressure applied slowly?	•					
13.	Gas flov	ow increased to approx. 600 mL/min [AMRL: ± 5 rpm]?	•					
14.	Spin rat	Spin rate of flask increased to approx. 45 rpm (see Note 5)? Note to Assessors: A 1 or 2 min delay before applying the vacuum is recommended. Hold or reduce vacuum						
	if foamin	ing occurs. Apply maximum vacuum when foaming stops.						
15.	This cor	ondition maintained for 15 ± 1 min?						
16.	Dietillat	ation flask removed and wiped clean of oil?	• ——					
17.	A enhalt	It poured (or drained) into proper size container?	•					
1/.	лэрнан	t pource (or dramed) into proper size container:	•					
COM	MENTS (D	D5404)·	D5404)					
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Revised 2011-03-25

COMMENTS (D6931):

INDIRECT TENSILE (IDT) STRENGTH OF BITUMINOUS MIXTURES

(D6931)

		<u>APPARATUS</u>	Date:			
	1. D.					
	ding Device	C 'IN (ID N				
(a)	Maker:	Serial No. (or I.D. N	0.)?			
(b)		$f 50 \pm 5$ mm/min $(2.00 \pm 0.15 \text{ in. /r})$	min)?			
(c)	Load measuring device:					
	(3) Sensitivity: minimum	50 N (10 lb)?				
		al indicator graduated in increments	s of 0.0025 mm			
Load	ding Strips					
(a)	Concave surface with a radius	of curvature equal to the nominal ra	adius of the test specimen?			
(b)	Widths:					
	(1) 4 in. diameter specime	ens, 12.70±0.3 mm (0.50±0.01 in.)?	?			
	(2) 6 in. diameter specime	ens, 19.05±0.3 mm (0.75±0.01 in.)?	?			
(c)	The length exceeds the thickne	ss of the specimens?	······			
(d)	The outer edges of the loading	strips beveled slightly to remove sl	harp edges?			
(e)	Upper loading strip clean and s	lides freely on the posts?	·····			
(f)	Two guide posts perpendicular	to base?				
(g)	Guide rods free of appreciable	binding or loose motion?				
(8)	Tr					
Tem	perature Control System					
(a)	An air or water bath capable of	maintaining the specimens at the s	specified test			
Ther	rm <mark>om</mark> eter					
(a)		f suitable range or any other thermo				
(b)	Subdivisions readable to 0.1°C	(0.2°F)?				
(c)	Calibrated (verify records)?	Materials Refere	nce Laboratory			
A ta _l	pe, ruler, or set of calipers for measure	uring specimens?				
Spec	cimens prepared according to one of	the following methods?				
	M D1074 (Compressive Strength of					
	M D1561 (California Kneading Co.					
	TM D3387 (USCE Gyratory Testing Machine [GTE])? TM D3496 (Dynamic Modulus Testing Preparation)?					
	M D4013 (Gyratory Shear Compac					
	M D6925 / AASHTO T312 (Super					
	M D6923 / AASHTO 1312 (Super) M D6926 / AASHTO T245 (Marsh					
ASI	M D0920 / AASHTO 1243 (Marsh					
Spec	eimens					
(a)	Specimen size					
(u)		ens - minimum height of 50 8 mm ((2 in.)?			
	(2) 6 in. diameter specime	one - minimum height of 75 mm (2	.95 in.)?			
(b)						
(b)	Note: A in specimens are suitable	e for mixtures with a nominal maximum	m particle size of 10 mm (2/4 in) or			
			particle size of 37.5 mm (1.5 in) or less.			
	1000. U-III. Specificity are sulfable I	oi matures with a nonlinal maximum '	particle Size of 37.3 Hill (1.3 HI) of ICSS.			

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(D6931)

COMMENTS (D6931):

INDIRECT TENSILE (IDT) STRENGTH OF BITUMINOUS MIXTURES

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				<u>PROCEDUR</u>	<u>E</u>	Date:				
Procedu	ıre									
1.		determinatio	n:							
	(a)			to ASTM D3549?		<u> </u>				
	(b)					sing a tape, rule, or calipers?				
(c) If using measurement jig, results consistently within ±0.05 in. (±0.13 cm) of those obtained using tape, rule, or calipers?										
(d) Measured to the nearest 1 mm (0.05 in)?										
	(e)			g volume by cross-se ith dense paving mixtur		area? than 10% air voids)				
2.	Diamet	er determina	tion:							
	(a)	Measured t	to the nearest 1 mn	n (0.05 in)?						
	(b)	Average of	four measuremen	ts taken at 90° increm	nents?					
	(c)	Measureme	ents taken at mid-h	eight?						
3.	Conditi	oning:								
	(a)				°F) by	one of the following methods				
			ided temperature is			f four house?				
	or	(1) Pr	rocedure A. Placed	in heavy duty leak-n	mum o	of four hours?				
	OI	or (2) Procedure B: Placed in heavy duty leak-proof plastic bag and paced in water bath for a minimum of 2 hours?								
	or	(3) Pr	ocedure C: Place	the specimens in a wa	ater bat	th for 30 to 120 minutes?				
4.	Load Determination: (a) Specimen removed from bath or oven and placed in lower segment of loading strip?									
5.	Calcula (a)		th calculated as fo	llower						
	(a)	IDT Such	gii caiculated as 10		C.					
			2000 * P		$S_t = P =$	IDT strength, kPa (psi) maximum load, N (lbf)				
In kPa		$S_t =$			t =	specimen height immed. before test mm (in.)				
•		-1	Pi * t * D		D=	specimen diameter, mm (in.)				
			2 * P		$S_t =$	IDT strength, kPa (psi)				
In psi	S	$S_t =$	2 * P		P = t =	maximum load, N (lbf) specimen height immed. before test mm (in.)				
тт ры		Pi * t * D D= specimen diameter, mm (in.)								

(D6931)