ASPHALT BINDER WORKSHEET INDEX

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^{**} NP for "Not Presented"

^{❖ -} Indicates the line has been modified since the previous version of the worksheets, 2013-09-20.

PRESSURIZED AGING VESSEL (PAV)

(R28)
(D6521)
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te
en?
± 0.02 MPa]?
est?

			<u> </u>	<u>APPARATUS</u>	Date:			
Draccuri	zed Agin	g Vaccal	(DAV).					
1.	Manufac	-						
2.								
3.					older?			
4.					······			
5.	Pan holder capable of holding pans in a level position?							
6.	Vessel							
	(a)	Separat						
		(1)	Contains a stand or shelf which the power surface of the oven		a level position above			
		(2)	Sufficiently large interior dim	ensions to allow forced	air to freely circulate			
					el is placed in the oven?			
or	(b)	Indeper	ndent temperature-controlled sy	stem (Prentex and ATS	s models)?			
Pressure	e Controll	ing Syst	em:					
1.	Pressure	gauge c	capable of measuring and control	olling the pressure to wi	ithin ± 1% [ASTM: ± 0.02 MPa]?			
2.	Capable	of main	taining the pressure within the	loaded vessel at 2.1 ± 0	.1 MPa during the test?			
	ature Con							
1.			ging the un-pressurized vessel to					
_		_	• /					
2.	Maintaii	ns the te	mperature within the vessel to 0	.5°C during the test?	······			
Tampar	ature Mea	curing I	Davica					
1.	Platinun	DTD o	r aquivalent, accurate and reads	able to the nearest 0.1%	C?			
2.								
	1							
Tempera	ature Rec	ording D	Device: SHTO Mar	terials Refe	erence Laboratory ng temperature throughout			
1.	Strip cha	art recor	der or other data acquisition de	vice capable of recordir	ng temperature throughout			
or								
2.	Electron	ic devic	e capable of reporting maximur	n and minimum require	ements (to 0.1°C)?			
COMM	ENTS (R	28/D652	21):		(R28/D6521)			

PRESSUR	IZED A	GING	VESSEL	(PAV)

(R28)	
(D6521)	

Date: _____

<u>APPARATUS (</u>	(Continued)	

Stainless Steel Pans	1	2	3	4	5	6	7	8	9	10
Diameter 140 mm (AMRL: 140 ± 5 mm)?										
Depth 9.5 mm (AMRL: 9.5 ± 5 mm)?										
Approximate thickness 0.64 mm [AMRL: at least 0.38 mm?]										
Bottom not badly warped?	·									

Note to assessors: It is recommended that the laboratory has at least ten pans.

	, p
Balanc	e:
1.	Class G2/ <i>GP2</i> (readable to 0.1 g)?
Vacuur	n Oven:
1.	Capable of maintaining temperatures within \pm 5.0°C [ASTM: an accuracy of \pm 5°C], for temperatures up to 180°C?
2.	Capable of a vacuum of 1.0 kPa (7.5 mm Hg) absolute [ASTM: $15 \pm 1 \text{ kPa}$]?
	(a) Temperature measuring device or temperature sensor capable of measuring the vacuum oven chamber temperature to within $\pm 5C$?
	(b) Vacuum measuring device or a vacuum gauge or digital vacuum measuring system capable of measuring the absolute pressure in the chamber to within ± 0.5 kPa (± 1.0 in. Hg)?
3.	AASHTO: If the vacuum oven is equipped with a relative pressure gauge, has it been adjusted for elevation according to the requirements of the test method?
	Note: weather data should not be used for this adjustment, as most weather data is pre-corrected for elevation.
Oven:	
1.	ASTM: Capable of maintaining a temperature of 168 ± 5 C?
<u>Air Su</u> 1.	pply: Commercial bottled air or equivalent?
COMM	MENTS (R28/D6521): (R28/D6521)

COMMENTS (R28/D6521):

	PRESSURIZED AGING VESSEL (PAV) (R28) (D6521)
	CALIBRATION AND STANDARDIZATION Date:
PAV Tei	mperature Detector [ASTM: Sensor]:
1.	AASHTO: Has the PAV temperature detector been verified to 0.1°C at least every 6 months using a calibrated thermometer?
2.	ASTM: Has the PAV temperature sensor calibration been verified at its respective meter or electronic circuitry to within \pm 0.1 $^{\circ}$ C at least every 6 months using a calibrated
3.	temperature measuring device traceable to a national standard?
PAV Pre	essure Gauge or Digital Pressure Measurement System:
1.	AASHTO: Has the pressure gauge been calibrated to an accuracy of ± 1% (± 0.02 MPa) at least every 6 months?
2.	ASTM: Has the pressure gauge or digital pressure measurement system calibration been verified to within ± 0.02 MPa at least every 6 months using a calibrated pressure indicator traceable to a national standard?
3.	ASTM: Is the verification performed near the pressure of use within a range of 2.00 to 2.10 MPa?
Vacuum	Oven Temperature Verifications:
1.	Is the vacuum oven temperature sensor calibration verified to within ±1 °C at least every 6 months using a calibrated temperature measuring device traceable to a national standard?
2.	Is the verification performed near the temperature of use within a range of 150 to 190 C?
Vacuum	Oven Pressure Verifications:
1.	Is the Vacuum oven vacuum gauge or digital vacuum measurement system - calibration verified to within ± 0.5 kPa (± 1.0 in. Hg) absolute pressure at least every 6 months using a calibrated vacuum or pressure indicator traceable to a national standard?
2.	Is the verification performed near the absolute pressure of use within a range of 12.5 to 17.5 kPa?
	Note: ASTM standardization for operator controlled pressure application is listed in Section 9.4. This gives guidance on how to determine the best temperature at which to apply the pressure during the aging procedure.
	SAMPLE PREPARATION
1.	Asphalt binder conditioned in accordance with RTFO, T240/D2872?
2.	Hot residue from RTFO combined into single container and stirred?
3.	If conditioned binder has cooled to room temperature, is it heated until sufficiently fluid to pour and stirred before it is poured into pans?
4.	Pan holder (without pans) placed inside the pressure vessel (if oven is used, vessel placed inside oven)?
5.	Aging temperature selected and vessel preheated (preheating the vessel 10 to 15°C above the aging temperature is acceptable)?

Revised 2014-06-02

(R28/D6521)

PRESSURIZED AGING VESSEL (PAV)

(R28)	
(D6521)	

	PROCEDURE Date:
Loading	$\mathbf{p}_{\mathbf{A}\mathbf{V}}$
1. 2. 3.	50 ± 0.5 g added to each stainless steel pan (not all pans need to be filled)?
4. 5. 6.	Pan holder with filled pans placed in PAV and PAV closed?
PAV Tes	
1.	If temperature inside vessel does not reach the desired temperature for applying pressure within 2 hours of loading pans into vessel, is procedure discontinued and samples discarded?
2.	Air pressure of 2.1 ± 0.1 MPa applied?
3.	Γiming of test started when pressure is 2.1 ± 0.1 MPa?
4.	Γemperature and pressure maintained for 20 hours ± 10 minutes?
5.	At the end of 20 hour test period, air pressure reduced over a 9 ± 1 minute interval
	[ASTM: 8 to 15 min.] (at an approximately linear rate of pressure decrease)?
	Note to Assessors: PAV depressurization timing should be stopped at 0.02 MPa Note ASTM: The bleed valve should be preset to equalize internal and external pressures on the PAV?
6.	AASHTO: If temperature indicated by recording device falls 0.5 $^{\circ}$ C above or below the target
0.	aging temperature for a total of more than 60 minutes during the 20 hours aging period,
	est declared invalid?
7.	ASTM: Test declared invalid if any of the following occur?
	Temperature deviates from the conditioning temperature by more than $\pm 0.5^{\circ}C$ for more than
	60 minutes total during the conditioning period?
	Temperature deviates from the conditioning temperature by more than $\pm 5^{\circ}C$ for more than 10
	minutes total during the conditioning period?
	c) If the device is capable of indicating only the minimum and maximum temperatures during the test, any temperature outside of ±0.5°C from the conditioning temperature?
Conditio	ning PAV Aged Material
1.	Pan holder and pans removed from PAV and pans placed in an oven at an oven set at a minimum
2.	emperature for a minimum time until sufficiently fluid to pour [ASTM: 168±5 °C for 15 ±1 minutes]?
3.	AASHTO: Any temperatures used beyond 175°C noted in the report:
<i>3</i> . 4.	Vacuum oven preheated until it stabilizes at 170 ± 5 °C prior to use?
5.	Pans removed from oven and residue poured into a single container?
6.	Container allows residue depth to be between 15 and 40 mm?
7.	Container transferred to the vacuum oven within one minute after last pan is scraped?
0	being performed on the collected residue.
8. 9.	Vacuum oven maintained at 170 ± 5°C for 15 ± 1 minutes without vacuum?
	(93.5 to 131.5 mm Hg or 3.7 to 5.2 in. Hg) absolute?
	AASHTO note: If the material foams over the lip of the container during degassing, slowly and
	temporarily reduce the vacuum until foaming ceases.
10.	Vacuum maintained for 30 ± 1 minutes?
11.	Vacuum released and container removed?
12.	If bubbles are visible on the surface, removed by flashing surface with a torch or hot knife?
13.	ASTM: If further testing is not performed immediately, container covered and stored at
	oom temperature?

7.

	GRADING OR VERIFYING PGB (R29)	
	APPARATUS Date:	
1.	Does the laboratory have equipment available to run the following tests or practices?*	
	(a) Pressure Aging Vessel (PAV) (R28)?	
	(b) Flash point (T48)?	
	(c) Rolling Thin-Film Oven (RTFO) (T240)?	
	(d) Bending Beam Rheometer (BBR) (AASHTO T313)?	
	(e) Direct Tension (DT) (T314) (Optional)?	
	(f) Dynamic Shear Rheometer (DSR) (T315)?	
	(g) Rotational Viscosity (T316)?	
	SAMPLE PREPARATION	
For Gr	ading an Unknown Asphalt Binder	
1.	Approximately 400 g of unaged asphalt binder obtained?	
E M-	wife in a ska Naminal Coada of an Asakak Dindon	
1.	rifying the Nominal Grade of an Asphalt Binder Approximately 250 g of unaged asphalt binder obtained?	
1.	Note to Assessor: If the laboratory does not do both grading and verification, an informational note will be written.	
	There to insense in any order to the decision of the configuration of th	
	<u>PROCEDURE</u>	
For G	rading an Unknown Asphalt Binder: (Assessor: Ask to see records):	
1.	DSR test (AASHTO T315) performed on the original asphalt binder?	
	(a) Sample tested beginning at 58°C?	
	(b) Test temperature increased or decreased in 6° increments until a value for	
	$G^*/\sin \delta \le 1.00 \text{ kPa is obtained?}$	
	(c) The highest test temperature where the value for G*/sin δ is ≥ 1.00 kPa is	
	used as the starting grade?	
2.	The flash point (AASHTO T48) determined on original binder is greater than 230°C*?	
3.	The viscosity (AASHTO T316) determined on original binder is less than 3 Pa·s*?	
4.	Approximately 200 g of asphalt binder is aged in the RTFO (AASHTO T240)?	
	(a) Change in mass determined is less than or equal to 1.00 percent*?	
5.	DSR test (AASHTO T315) performed on RTFO residue?	
	(a) The test temperature determined as the starting grade (from DSR original binder) is used?	
	(b) If $G^*/\sin \delta$ is ≥ 2.20 kPa, this test temperature used as the high temperature grade?	
	(c) If the value of $G^*/\sin \delta$ is less than 2.20 kPa, the material is tested again at a temperature 6° lower?	
_	(d) The lowest test temperature where $G^*/\sin \delta$ is ≥ 2.20 kPa is used as the high temperature PG grade?	
6.	RTFO residue aged in the PAV (AASHTO T28)?	
	(a) Aged at a temperature of 90°C for material with a high temperature grade of 46 or 52°C?	
	(b) Aged at a temperature of 100°C for material with a high temperature grade of 58°C or higher?	
	(c) Aged at a temperature of 110°C to simulate a desert environment?	

*If these requirements cannot be met (according to AASHTO M320), then no further testing is required.

COMMENTS (R29): (R29)

PAV residue combined into a single container?....

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(R29)	
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<u>PROCED</u>	<u>URE (</u>	Continued)	

8.	DSR tes	t (AASHTO T315) performed on PAV residue?
	(a)	Initial test temperature conforms to the table below unless there is other information to suggest a
		temperature at which G*sin δ exceeds 5,000 kPa?

Starting Grade*	Initial DSR Test Temperature- PAV Residue (°C)
PG 52	16
PG 58	19
PG 64	22
PG 70	28

*Starting grade determined from DSR test on original and RTFO aged residue.

	(b)	Test temperature increased or decreased at 3° C increments until the value for
		G*sin δ exceeds 5,000 kPa?
9.	BBR 1	test (AASHTO T313) performed on PAV residue?
	(a)	Two specimens prepared and tested (results averaged)?
	(b)	The initial test temperature determined using the high temperature grade,
		the temperature determined for the DSR test on PAV residue, and Table of M320?
	(c)	Test temperature increased in 6°C increments until the stiffness is less than
		or equal to 300.0 MPa and the slope m is greater than or equal to 0.300?
	(d)	New BBR test specimens prepared for each change in test temperature?
10.	DT te	st performed on PAV residue (Optional)?
	(a)	Creep stiffness from BBR test on PAV residue is between 300 and 600 MPa?
	(b)	Slope from BBR test on PAV residue is greater than or equal to 0.300?
	(c)	Results of four specimens reported and averaged?
	(d)	Initial test temperature used is the first temperature from BBR test on PAV aged
		residue at which the slope is greater than or equal to 0.300?
	(e)	Test temperature increased in 6°C increments until a failure strain greater than or
		equal to 1.0 percent is obtained?
	(f)	equal to 1.0 percent is obtained?
11.	Final	grade of the asphalt binder determined using the data collected and Table 1 of M320?

COMMENTS (R29): (R29)

		GRADING OR VERIFYING PGB	(R29)
		PROCEDURE (Continued) Date: _	
For V	erifying t	the Nominal Grade of an Asphalt Binder: (Assessor: Ask to see records)	
1.	DSR	test (AASHTO T315) performed on the original asphalt binder?	
	(a)	The test temperature used is the high temperature grading designation?	
	(b)	If the value of G*/sin δ is ≤1.00 kPa, testing continued?	
	(c)	If the value of G*/sin δ in not \leq 1.00 kPa, the specimen is treated like a	
		binder of unknown grade (see above)?	
2.	The f	lash point (AASHTO T48) determined on original binder is greater than 230°C*?	
3.	The v	viscosity (AASHTO T316) determined on original binder is less than 3 Pa·s*?	
4.	Appro	oximately 100 g of asphalt binder aged in the RTFO (AASHTO T240)?	
	(a)	Mass loss determined is less than or equal to 1.00 percent*?	
5.	DSR	test (AASHTO T315) performed on RTFO residue?	
	(a)	The test temperature used is the high temperature grading designation?	
	(b)	If the value of G*/sin δ is \geq 2.20 kPa, testing continued?	
	(c)	If the value of G*/sin δ in not \geq 2.20 kPa, the specimen is treated like a binder	
		of unknown grade (see above)?	
6.		residue combined into a single container?	
7.		test (AASHTO T315) performed on PAV residue?	
	(a)	The test temperature determined from the temperature grading designation and Table 1	
	(b)	If the value of G*sin δ is less than 5,000 kPa, testing continued?	
	(c)	If the value of $G*\sin\delta$ in not less than 5,000 kPa, the specimen is treated like a	
_		binder of unknown grade (see above)?	·····
8.		test (AASHTO T313) performed on PAV residue?	
	(a)	Two specimens prepared and tested (results averaged)?	
	(b)	The test temperature determined from the temperature grading designation and Table 1	
	(c)	If the slope is ≥0.300, testing continued?	
•	(d)	If the slope is not ≥ 0.300 , the specimen is treated like a binder of unknown grade (see a	
9.		est performed on PAV residue (Optional)?	
	(a)	Creep stiffness from BBR test on PAV residue is between 300 and 600 MPa?	
	(b)	Slope from BBR test on PAV residue is greater than or equal to 0.300?	oratory—
	(c)		
	(d)	If the failure strain is greater than or equal to 1.0 percent, the grade is considered verifi	ea!
	(e)	If the failure strain is not greater than or equal to 1.0 percent, the specimen is	
		treated like a binder of unknown grade (see above)?	

*If these requirements cannot be met (according to AASHTO M320), then no further testing is required.

COMMENTS (R29): (R29)

COMMENTS (R29):

(R29)

	GRADING OR VERIFYING PGB		(R29)
	<u>REPORTING</u>	Date:	
For Gra	ding an Unknown Asphalt Binder:		
1.	Results of all tests performed reported?		
2.	PG grade designation reported (example PG 52-34)?		
For Ver	ifying the Nominal Grade of an Asphalt Binder:		
1.	Results of all tests performed reported?		
2.	Whether or not binder meets the requirements of M320 reported?		



SOLUBILITY OF ASPHALT MATERIALS IN TRICHLOROETHYLENE

(144)	
(D2042)	

		<u>APPARATUS</u>	Date:
1.	Good	h Crucibles	
	(a)	Glazed surface throughout except bottom exterior unfinished?	
	(b)	Approximately 44 mm at the top tapering to 36 mm at the bottom [Al	
	(c)	Approximate depth of 24 to 28 mm [ASTM: 20 to 30 mm]?	
2.		ation Assembly	
	(a)	Heavy walled filter flask with side tube, capacity 250 mL or larger [A Note to assessors: any other assemblies permitting vacuum filtration with a Gooch cru	
	(b)	Glass fiber pads with a diameter of 32, 35, or 37 mm [ASTM: 32 to 3	
	(0)	fast flow rate, 1.5 µm particle retention?	
	Sucti	on Assembly	
•	(a)	Satisfactory assembly?	
١.	\ /	loroethylene Solvent	
	(a)	Technical Grade Type 1 (Reagent Grade) Trichloroethylene,	
	()	or Technical Grade 1,1,1 Trichloroethane	
	(b)	ASTM: Trichloroethylene, Technical (Reagent) grade?	
i.	Desic	•	
	(a)	Satisfactory design and charged with effective desiccant?	
		ng Oven	
	(a)	Maintains temperature at 110 ± 5 °C (230 ± 9 °F)?	
·.		ellaneous Items	
	(a)	Suitable container for weighing and dissolving sample?	
	(b)	Class A balance (readable to 0.0001 g) available?	
	(c)	Policeman (optional)?	
		· / AIVIEI	
		PREPARATION OF THE GOOCH CRUCIBLE	
	AASI	HTO Procedure	
•	(a)	Filtering annaratus assembled?	T. I
	(b)	Filtering apparatus assembled?	nce Laboratory
	(c)	New glass fiber pad placed in crucible?	
	(d)	Pad wetted with solvent and seated firmly with light suction?	
	(e)	Crucible and contents dried at 110 ± 5 °C (230 ± 9 °F) for at least 20	
	(f)	Crucible and contents cooled in a desiccator for at least 20 min.	<i></i>
	(1)	and then weighed to nearest .0001g?	·····
	(g)	Drying and cooling procedure repeated until constant mass (± 0.000	3 g) is obtained?
	(h)	Crucible stored in a desiccator until used?	·····
	ASTA	<u> 1 Procedure</u>	
	(a)	New filter pad placed in crucible and dried in oven at $110 \pm 5^{\circ}C$ for (no wetting and seating required)?	
	(b)	Cooled in a desiccator for 30 ± 5 and weighed to the nearest 0.0001	
	(c)	Stored in a desiccator until ready to use?	
	\-/	,	
COM	MENTS	(T44/D2042):	(T44/D2042

SOLUBILITY OF ASPHALT MATERIALS IN TRICHLOROETHYLENE

(144)	
(D2042)	

	PROCEDURE Date:		
1.	If the sample is not fluid is it heated with care to prevent local overheating?		
2.	Sample stirred occasionally and the entrapment of air avoided?		
3.	ASTM: Sample heated at any temperature not more than 100 °C above softening point?		
4.	Approximately 2 g [AMRL: ± 0.5 g] of sample placed in tared (nearest 0.001 g) container?		
5.	Container with sample allowed to cool and then weighed to nearest 1 mg (0.001 g)?		
6.	100 mL of solvent added to container, flask stoppered, and then container agitated		
	as necessary until the sample is dissolved?		
7.	ASTM: Solvent added in small portions with constant agitation?		
8.	Lumps completely digested and container sides free of undissolved sample?		
9.	ASTM Only: No undissolved material visible after 15 minutes in the solvent?		
	Note to Assessors: for referee testing, flask and solution shall be placed in a water bath at $38.0\pm0.3\%$ for		
10	1 hour before filtering. Please discuss with laboratory if necessary.		
10.	Crucible placed in filter tube and wetted?		
11. 12.	Asphalt solution decanted through filter with light suction?		
12.	If insoluble matter is visible:		
	(a) Retained in container until solution has drained through filter?		
	(c) Container and policeman (if used) rinsed?		
	(d) Insoluble matter washed until the filtrate is substantially colorless?		
	(e) Strong suction applied to remove remaining solvent?		
13.	Crucible removed and bottom washed free of dissolved matter?		
14.	Placed in oven at 110 ± 5°C for at least 20 min?		
15.	Cooled in desiccator for at least 20 min. [ASTM: $30 \pm 5 \text{ min}$] and then weighed to nearest .0001g?		
16.	Steps (14) and (15) repeated until constant mass of \pm 0.0003 g obtained?		
10. 17.	Percent insoluble reported to nearest 0.1%?		
18.	ASTM: If percent insoluble is less than 1%, reported to nearest 0.01%?		
19.	ASTM: If the crucible in desiccator must be left overnight, is it placed		
1).	in oven for 30 min, and then cooled again in desiccator?		
	in oven for 30 min. and then cooled again in desiccator?		

COMMENTS (T44/D2042):

(T44/D2042)

COMMENTS (T48/D92):

APPARATUS FOR FLASH AND FIRE POINT BY CLEVELAND OPEN-CUP

(T48)	
(D92)	

		<u>APPARATUS</u>	Date:
1	Elech T	Poston.	
1.			
		Automatic or Manual? (Please Record)	······
(b) Supported on 1 (c) Draft-free local 2. Test Cup (a) Outer diamete (b) Outer diamete (c) Inner diameter (d) Rim to the fill (e) Depth of the c (f) Bottom thickn 3. Heating Plate (a) Plate of accept (b) Hard board of (c) Support holds 4. Test Flame Applicator (a) Mounted on at (b) Swing radius r (c) Orifice diamet (d) Orifice not mot (e) Diameter of th (f) Bead 3.8 to 5. (g) Bead mounted 5. Heater (a) Centered unde (b) No local super (c) No gases or flat 6. Thermometer (a) ASTM 11C/11 or Electronic (b) Support can be (c) Thermometer (d) Positioned on 7. Filling Level Gage (AA (a) Height of projects	Draft-free location?		
2	· /		······································
۷.		Outer diameter of the flange 97 to 101 mm [ASTM: 97 to 100 mm]?	
	` /	Outer diameter of the flange 67.5 to 69 mm?	
	` '	Inner diameter of the cup 62.5 to 64 mm [ASTM: 63 to 64 mm]?	
		Rim to the fill mark 9 to 10 mm in length?	
	` /	Depth of the cup 32.5 to 34 mm?	
	` '	Bottom thickness 2.8 to 3.6 mm [ASTM: 2.8 to 3.5]?	
3.	` /		
		Plate of acceptable design covering heater top?	
		Hard board of suitable design?	
	(c)	Support holds plate level and steady?	
4.	Test Fl	ame Applicator	
	(a)	Mounted on apparatus?	······
	(b)	Swing radius not less than 150 mm (6 in.)?	
	(c)	Orifice diameter approximately 0.8 mm (0.031 in.)?	
	(d)	Orifice not more than 2.5 mm [ASTM: 2 mm] above the cup?	<u></u>
	(e)	Diameter of the tip 1.6 to 5.0 mm [ASTM: approximately 1.6 mm]?	
	(f)	Bead 3.8 to 5.4 mm [ASTM: 3.2 to 4.8 mm]?	
	(g)	Bead mounted on apparatus [ASTM: not required to be mounted]?	<u></u>
5.			
	(a)	Centered under plate opening?	
	(b)	No local superheating?	······ <u> </u>
		No gases or flames up around the cup?	······· <u> </u>
6.		ometer ASTM 11C/11F or IP 28C/28F?	Laboratory
	(a)		
		or Electronic device such as a resistance thermometer or thermocouple?	
	. ,	Support can hold thermometer vertically 6.4 ± 0.1 mm from the cup bottom	
		Thermometer positioned halfway between the center and side of cup?	
7	` /	Positioned on diameter perpendicular to arc of sweep of test flame?	
/.		Level Gage (AASHTO Only – Optional)	
	` /	Height of projections 9 to 10 mm?	
		Flame orifice guide-hole diameter 0.8 mm?	
	(<i>C</i>)	Guide hole not more than 2.5 mm above bottom edge of gage?	······

Revised 2014-06-02

(T48/D92)

	FLASH AND FIRE POINTS BY CLEVELAND OPEN-CUP (148) (D92)
	CALIBRATION AND STANDARDIZATION Date:
1.	Temperature measuring device calibrated according to manufacturer's instructions?
2.	Performance of manual or automated apparatus verified (annually) by testing a certified reference material (CRM), or equivalent (not AMRL Proficiency Sample), which is reasonably close to the expected temperature range of the samples to be tested? (CRM must have certificate from manufacturer)?
3.	Using a secondary working standard (SWS), a mean flash point and statistical control limits (3σ) for the SWS established? (Optional)
	(a) When an observed flash point is outside of these control limits, is a close check of the apparatus made and the test repeated with a fresh specimen while paying close attention to the procedural

Additional information taken from Annex A2 (included only for guidance)

<u>CRM</u> - A certified reference material is a stable, pure (99 + mole % purity) hydrocarbon or other stable petroleum product with a method-specific flash point established by a method-specific interlaboratory study following RR: D02-1007* or ISO Guide 34 and 35.

Typical Flash Point Values and Typical Limits for CRM						
Hydrocarbon Purity Flash Limits						
	(mole%) Point (°C) (
n-tetradecane	99+	115.5	±8.0			
n-hexadecane	99+	138.8	±8.0			

Calculation of the limits for other CRM's can be determined from the reproducibility values of this test method, reduced by interlaboratory effect and then multiplied by 0.7 (see Research Report RR:S15- 1008^*).

SWS – A secondary working standard is a stable, pure (99 + mole % purity) hydrocarbon, or other petroleum product whose composition is known to remain appreciably stable.

Research Reports and supporting data are filed at ASTM International Headquarters.

PREPARATION OF TEST CUP

1.	Cup washed with solvent to remove any test specimen or traces of gum or residue?
2.	If deposits of carbon are present, removed with steel wool?
3.	Cup rinsed with cool water and dried on hot plate or over open flame?
4.	Cup cooled to at least 56°C (100°F) below the flash point before using?
COM	MENTS (T48/D92): (T48/D92)

FLASH AND FIRE POINTS BY CLEVELAND OPEN-CUP

(148)	
(D92)	

		<u>PROCEDURE</u>	Date:
Antic	cipated Flash Point:°C°F		
1.			pable flash point?
2.			
3.			and refilled? Air bubbles destroyed?
4.			
5.			F)] per min?
		nknown, test flame applied beginn	
			to 6 °C (9 to 11 °F)] per min?
		ry test flame applications during	
		ng whether unexpected volatile m	aterial is present in
	the sample (recommend 10°C)		
		point material or with highly vise	cous material it is
	advised to use the 5 to 6 °C / m		
6.	When sample temp is approximately 56° point is heat decreased?		F)] below anticipated flash
7.	Rate adjusted to 4 to 7°C (7 to 13°F) [AS	STM: 5 to 6 ℃ (9 to 11 ℉)] per r	ninute?
8.	Starting at least 28°C (50°F) below flash	n point, rate maintained and test fla	ame applied?
9.	Test flame applied for each 2°C (5°F) m	nark with smooth continuous motion	on?
10.	Direction reversed with each pass of test	t flame?	<u></u>
		testers that only pass the flame in	
11.			
12.	Care taken to avoid disturbing vapors by		
13.			irring rod?
14.			ture?
15.	Temperature read when flash appears at	any point on surface of sample?	ith a fresh sample?
16.	If flash is detected on first application, is	s the test stopped then restarted w	ith a fresh sample?
17.			
	Note to Assessors: The sample is deeme	v v	* *
	and instantaneously propagates itself ov	ver the entire surface. A blue hald	or an enlarged
	flame is not a flash.		
18.			······
	Note to Assessors: Many aneroid barom		
	are pre-corrected to give sea level reading		
19.		<u> </u>	using formula in book?
20.	Flash point reported to nearest 1°C (2°F)	7)?	······
COM	IMENTS (T48/D92):		(T48/D92)

PENETRATION OF BITUMINOUS MATERIALS

(T49)	
(D5)	

			<u>APPARATUS</u>		Date:			
1.	Penet	rometer						
1.	(a)	Dial accurate to 0.1 mm?						
	(b)	Spindle readily detaches?						
	(c)							
	(d)	Mass of 50 g weight 49.95 to 50.05 g	?					
	(e)	Mass of 100 g weight 99.95 to 100.05	g?					
	(f)	Needle moves vertically and perpendicular						
	(g)	Device equipped with a leveling indic						
2	(h)	Level indicator verified annually with	a hand held level	?				
2.		PENETROMETER NEEDLES	1	2	3			
			1	2	3			
		Needle Number (not required, please reference for reporting)?						
		Mass of needle 2.45 to 2.55 g?						
		Needle Diameter of 1.00 to 1.02 mm?						
		Ferrule Diameter of 3.15 to 3.25 mm?						
		Ferrule length of 37 to 39 mm?						
		Needle straight (roll on flat surface)?						
		Surface finish smooth?						
		End symmetrically tapered and straight?						
	(a) (b) (c) (d)	For penetrations less than 40, container For penetrations less than 200, container For penetrations between 200 and 350. For penetrations greater than 350, con <i>Note to Assessors:</i> For referee testing below 40 shall be 55 mm in diameter.	ner 55 mm diameto, container 55 to 7 tainer 55 mm diar 4, the container fo	er and 35 mm dee 75 mm diameter a meter and 70 mm or r testing materials	p?d 45 to 70 mm deep?deep?			
4.	Wate	r Bath	by 33 mm in aepir	<i>i</i> .				
	(a)	Capable of maintaining temperature w	rithin 0.1°C (0.2°F	E) (e.g. 77.0 ± 0.29	°F)?			
	(b)	At least 10 liters of water in bath?			- / · · · · · · · · · · · · · · · · · ·			
	(c)	Perforated shelf at least 100 mm below						
	(d)	Water in bath clean?						
5.	Ther	<u>mometer</u>						
	(a)	Any thermometer or thermometric dev						
	(b)	Thermometer used calibrated?						
_	(c)	AASHTO only: Thermometer immers						
6.		Transfer Dish for Container (for penetr						
	(a) (b)	Capacity of at least 350 mL? Sufficient depth for water to cover san						
	(c)	Means of preventing rocking of sampl						
7.		ng Device	e container provie	icu (usuany magn	Ct):			
•	(a)	Electric timer, stopwatch, or other dev	vice graduated to 0	0.1 s or less				
	V-7	and accurate to ± 0.1 s for 60-s interva						
or	(b)	Audible seconds counter providing on						
or		Automatic timing device on penetrom						
8.		Source for illuminating surface of specir						
COMN	MENTS	(T49/D5):				(T49/D5)		

PENETRATION OF BITUMINOUS MATERIALS

(T49)	
(D5)	

			SAMPLI	E PREPARATI	<u>ON</u>	Date:		
1.	Sample heated	to loss than 00°C ab	ova avnaatad (oftoning point	for acphalt?			
2.								
2. 3.	Sample heated for the minimum time necessary to make sample sufficient fluid?							
3.	Sample surred	١٢٠			•••••	•••••		
			<u>PR</u>	OCEDURE				
1.	Expected pene	etration value?						
2.								
3.								
4.								
5.		s less than 65 mm in						
6.	•	ed to cool at 15 to 30						
				mm or smaller	containers?			
7.)	
8.							a time of 5 s?	
							at the bottom of page	
9.								
10.	If transfer dish is used, water covers entire sample?							
11.	Levelness of apparatus ensured using the level indicator?							
12.								
13.								
14.								
15.								
16.	If transfer dish	is used, dish with sa	mple_returned	to bath after ea	ach penetration	?		
17.	When one nee	dle is used, cleaned v	with solvent me	oistened cloth a	after each pene	tration	oratory	
	and then wiped	d with a clean, dry cl	oth?					
18.	Test run with a	a long needle for sam	ples with a per	netration greate	er than 350, rur	with a short n	eedle otherwise?	
19.	If penetration	value is over 200, are	e needles left in	n sample until o	completion of t	est?		
20.								
21.	Penetrations at	t least 1 cm (10 mm)	from side and	bottom of cont	ainer and each	other?		
22.		least three penetratio						
	amount shown	in the table below?						
							_	
		Penetration	0 to 49	50 to 149	150 to 249	250 to 500		
	Max Difference 2 4 12 20							

COMMENTS (T49/D5): (T49/D5)

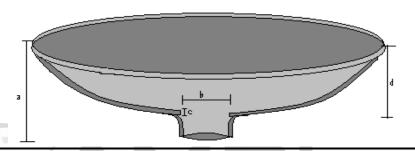
FLOAT TEST FOR BITUMINOUS MATERIALS

(T50) _____ (**D139**) _____

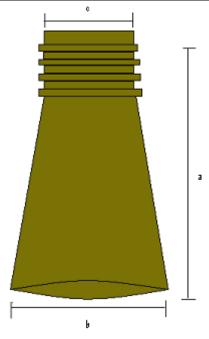
Date: ____

<u>APPARATUS</u>

Float	1	2	3	4
Made of Aluminum alloy?				
Weight: 37.90 ± 0.20g				
a =Total height: 35.0 ± 1.0 mm				
b =Diameter of opening: 11.1 ± 0.1mm				
c =Thickness of shoulder: 1.4 ± 0.1mm				
d =Height of rim above lower side of				



Collar	1	2	3	4	
Made of Brass?					
Weight: 9.8 ± 0.20g					
$a = \text{Height: } 22.5 \pm 0.2 \text{mm}$					atory
b = Inner diameter at bottom: 12.82 ± 0.10mm					
c = Inner diameter at top: 9.70 ± 0.05mm					



FLOAT TEST FOR BITUMINOUS MATERIALS

(T50) _ (D139) _	

				APPARATUS (Contin	nued)	Date:
1.	Have f	oat and c	collar assemblies been	standardized for depth o	of immersion?	
2.						
3.	Water			1		
	(a)	Testing	Bath			
		(1)	Circular bath with in			of at least 185 mm?
	or	(2)	Rectangular bath at 1	east 150 by 300 mm and	d water depth of at least	110 mm?
		(3)				
		(4)				<u></u>
		(5)		rted in bath to a depth of	f 40 ± 2 mm below the w	vater surface?
	(b)	Cold B				
		(1)				
		(2)				
4.						
5.					ge enough for the sample	
6.			_			
7.	Stop watch?					
				PROCEDURE		
				FROCEDURE		
1.	Collar	placed wi	th smaller end down o	n plate prepared with re	lease agent?	
2.					to a fluid condition (If e	
						······ <u> </u>
3.	Sample	stirred w	vithout incorporating a	ir bubbles?		
4.						
5.	Sample	cooled a	it room temperature fo	r 15 to 60 min. (not nece	essary for tar products)?	·······
6.	Sample	immerse	ed in 5°C (41°F) water	bath for 5 minutes?	Reterence	Laboratory
7.	Excess	material	trimmed with slightly	heated spatula or steel k	nife?	
8.	Sample	flush wi	th top of collar?			
9.	Collar	and plate	put back in 5°C (41°F) water bath for 15 to 30) minutes?	············ <u> </u>
10.	Testing	bath ma	intained at ± 0.5 °C (0.	9°F) of test temperature	throughout the test?	
11.						nter surface?
12.	Collar	and conte	ents removed from plat	e and screwed into float	?	
13.	Float a	nd collar	assembly completely i	mmersed in 5°C (41°F)	water for 1 minute?	············ <u> </u>
14.	Water	removed	from inside of float?			
15.						
16.						<u></u>
17.						
18.						
19.	Time re	ecorded?				

COMMENTS (T50/D139):

(T50/D139)

COMMENTS (T51/D113):

DUCTILITY OF BITUMINOUS MATERIALS

(T51)	
(D113)	

Date: ____

APPARATUS

Molds	1	2	3	4	5	6
Design conforms to Fig. 1?						
Thickness: 9.9 to 10.1 mm?						
Width at midpoint: 9.9 to 10.1 mm?						
Made of brass?						
Mold Plates	1	1	2	2	3	
Non-absorbent?						
Made of brass?						
Flat and level?						

1.	Wate	<u>r Bath</u>
	(a)	Maker:
	(b)	AASHTO: Depth of water shall be not less than 50 mm
	(c)	AASHTO: Specimens can be immersed to a depth of at least 25 mm?
	(d)	AASHTO: Water free from oil and slime?
	(e)	AASHTO: Volume of water not less than 10 L?
	(f)	ASTM: Specimens immersed and supported such that they are surrounded by water?
	(g)	Bath maintains temperature within 0.5°C (0.9°F)?
2.	Ducti	lity Machine
	(a)	Maker:
	(b)	Serial No. (or I.D. No.)?
	(c)	Space for at least 25 mm of water above and below sample at start of test?
	(d)	Machine maintains specified speed within 5 percent (e.g. 5.00 ± 0.25 cm/min.)?
	(e)	Machine functions without undue vibration?
	(f)	ASTM: Means of measuring elongation of the specimen in centimeters?
3.	Agen	t for Adjusting Specific Gravity (information only)
	(a)	Specimen sinks to the bottom:
	(b)	Specimen floats to the top:
4.	Ther	mometer
	(a)	ASTM 63C or 63F thermometer or any other liquid in glass equivalent?
	` ′	or
	(b)	An equivalent thermometric device that has been calibrated in accordance with Test Method E220
		or E644 (Assessors: Records should indicate use of one of these methods)?
	(c)	AASHTO: An electronic temperature device may be used if it exhibits the same temperature
		response as the mercury thermometer?
5.	Relea	ise Agent
	(a)	One of the following:
	. ,	Glycerin and Dextrin, Talc, or Kaolin (China Clay)
		Versamid Resin and Mineral Oil
		AASHTO: Dow-Corning stopcock grease
		Other materials that have been shown not to affect physical properties of the test specimen:
		
6.	•	ellaneous Equipment
	(a)	Straight-edged trimmer at least 1 ½ in. wide [ASTM: Wider than the specimen]?
	(b)	ASTM: 300-µm (No. 50) sieve?

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(T51/D113)

DUCTILITY OF BITUMINOUS MATERIALS

(151)	
(D113)	

	<u>PROCEDURE</u>	Date:
1.	Mold assembled on plate prepared with release agent?	
2.	Interior surface of mold sides treated with release agent along center pieces only?	
3.	Sample carefully heated to prevent local overheating and then stirred?*	
4.	Mold filled by pouring a thin stream back and forth from end to end?	
5.	Mold filled until more than level full?	
<i>5</i> . 6.	Disarrangement of mold parts avoided during filling?	
7.	Sample, mold, and plate cooled at room temperature? Start Time:	
8.	Cooling time 30 - 40 min. at room temperature? End Time:	······
o. 9.	Placed in water bath for 30 min. [ASTM 30 to 40 min]? Start Time:	······
9. 10.		
	Bath water within 0.5°C (0.9°F) of test temperature?	
11.	Excess material cut off with hot straight-edged putty knife or spatula? End Time:	
12.	Mold level full?	
13.	Specimen not pulled away from mold?	
14.	Sample, mold and plate placed again in water bath? Start Time:	
15.	Bath water within 0.5°C (0.9°F) of test temperature?	
16.	Conditioned in water bath for 85 - 95 min.? End Time:	
17.	Mold taken off plate and side pieces of mold detached?	
18.	Bending of the specimen, distortion, or fracture avoided?	
19.	Briquette placed in testing machine and tested immediately?	
20.	Water in testing machine within 0.5°C (0.9°F) of test temperature?	
21.	Specimen pulled to a point or thread and does not touch bottom or surface?	
22.	Is gravity adjusted and test repeated if thread does contact top or bottom?	<u></u>
23.	Distance clips pulled to produce rupture measured in cm?	
24.	If specimen does not rupture is it noted on the report (ASTM: Noted as "length+" - i.e.	
25.	For referee testing, three specimens tested and results averaged?	

*Note: ASTM: If emulsion or cutback residue is used, it is recommended that the material be sieved through at 300 mm (No. 50) sieve prior to pouring. If the material is sieved prior to testing, it should be noted on the report. For referee testing residue material must be poured through a 300 mm sieve.

COMMENTS (T51/D113): (T51/D113)

SOFTENING POINT OF BITUMINOUS MATERIALS

(T53)	
(D36)	

Date: _____

<u> AP</u>	<u>PA</u>	\mathbf{R}	\TI	<u>US</u>

RINGS (at least two)	1	2	3	4	5	6
Outer Diameter at top of shoulder: 22.7 to 23.3 mm?						
Inner Diameter at top of shoulder: 19.5 to 20.1 mm?						
Outer Diameter at bottom of ring: 18.5 to 19.1 mm?						
Inner diameter at bottom of ring: 15.6 to 16.2 mm?						
Total Height of rings: 6.0 to 6.8 mm?						

STEEL BALLS (at least two)	1	2	3	4	5	6
Diameter approximately 9.5 mm [AMRL: 9 to 10 mm]						
Weight: 3.45 to 3.55 g?						

(a)	Two brass guides resembling guides in Fig. 1 (c) available?
Bath	
(a)	Glass vessel with a minimum inside diameter of 85 mm and not less than 120 mm in depth?
	Note to Assessors: An 800 mL beaker, Griffin Low-Form, heat resistant glass, meets this requirement
Thern	nometers .
(a)	ASTM 15C or 15F for distilled water?
(b)	ASTM 16C or 16F for USP Glycerin?
(c)	ASTM 113C or 113F [ASTM: ASTM 16C or 16F] for Ethylene Glycol?
(d)	Thermometer positioned so bottom of bulb is level with the bottom of rings and within
	13 mm (0.5 in.) of the rings but not touching them or the ring holder?
(e)	Thermometer can be read after 3 minutes with bulb at bottom of rings?
(f)	AASHTO: Another thermometric device meeting the following requirements
	(1) Maximum scale error no greater than that of the thermometer specified in ASTM E1?
	 (1) Maximum scale error no greater than that of the thermometer specified in ASTM E1? (2) Capable of indicating the temperature within 0.2°C (0.5°F)?
(g)	ASTM: Any thermometric device can be used as long as meets the following requirements:
	(1) Equal accuracy or better?
	(2) Capable of indicating temperature to within 1° C $(2^{\circ}F)$?
	(3) Stable to within 1° C $(2^{\circ}$ F) for the duration of exposure?
Ring 1	<u>Holder</u>
(a)	Holder accommodates two rings only in a horizontal position?
(b)	Bottom of rings 25 mm above upper surface of bottom plate?
(c)	Lower surface of bottom plate 16 ± 3 mm above bottom of the bath?
Bath l	Liquids
(a)	Freshly boiled distilled water for tests between 30 and 80°C (86 and 176°F)?
(b)	USP Glycerin for tests between 80 and 157°C (176 and 315°F)?
(c)	Ethylene Glycol for test between 30 and 110°C (86 and 230°F)?
	(1) Boiling point between 193 and 204°C (379 and 399°F)?
	(2) ASTM: Boiling point between 195 and 197°C (383 and 387°F)?
Misce	llaneous
(a)	Release agent available (standard or comparable)?
(b)	Base plate, brass, and approximately 50 by 75 mm [AMRL: Large enough for the samples]?
(c)	Forceps?
(d)	Knife or spatula?
(a)	Gos humar or alactric hantar?

COMMENTS (T53/D36):

(T53/D36)

SOFTENING POINT OF BITUMINOUS MATERIALS

(T53)	
(D36)	

	PROCEDURE Date:
1.	Sample heated less than 2 hours and to temperature less than 110°C (200°F) above the softening point?
2.	Pouring plate coated with release agent?
3.	Brass rings heated to approximate pouring temperature?
4.	Enough sample poured into two rings to provide excess when cool?
5.	Samples cooled on flat surface at room temperature? Start Time:
6.	If samples are soft at room temperature, cooling done at minimum of 10°C (18°F)
	below expected softening point?
7.	Elapsed cooling time at least 30 minutes? End Time:
8.	Excess material cut off level with warmed knife?
9.	Apparatus assembled with rings, correct thermometer, and ball centering guides in position?
10.	Bath filled to a depth of 102 to 108 mm (4 to 4.25 in.) with appropriate bath liquid?
11.	Balls adjusted to bath temperature before use?
12.	Bath maintained at the proper starting temperature for 15 minutes:
	(a) Ethylene glycol and freshly boiled, distilled water: 5 ± 1 °C (41 ± 2 °F)?
	(b) USP Glycerin: 30 ± 1 °C (86 ± 2 °F)?
13.	Care taken to avoid contamination of bath liquid?
14.	Ball placed in each ball centering guide with forceps?
15.	Heat applied from below avoiding drafts?
16.	After 3 minutes, is rate of temperature rise controlled to 5.0 ± 0.5 °C (9.0 ± 1.0 °F) per minute?
17.	Temperature shown by the thermometer at the instant the sample surrounding the
	ball touches the bottom plate recorded for each ring and ball?
18.	Total elapsed time from <u>preparation</u> to completion of testing less than 6 hours?
19.	Total elapsed time from <u>pouring</u> to completion of testing less than 4 hours?
20.	If softening point temperatures differ by more than 1°C (2°F), is test repeated?
21.	Results obtained in an ethylene glycol bath corrected to water or glycerin using appropriate formula?
22.	Mean temperature of duplicate determinations reported to:
	(a) Nearest 0.5°C or 1.0°F when using an ASTM 16C/16F thermometer?
	(b) Nearest 0.2°C or 0.5°F when using an ASTM 15C/15F thermometer?
	(c) Nearest 0.5°C or 1.0°F when using an ASTM 113C/113F thermometer?
COMM	ENTS (T53/D36): (T53/D36)

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(1)

(1)

(b)

(c)

For trap styles E verification performed as follows:

For trap styles F, verification performed as follows:

WATER IN BITUMINOUS MATERIALS

(T55)	
(D95)	

		APPARATUS Date:
1.		Solvents
		(a) Xylene (Industrial grade) (for asphaltic materials)?
	or	(b) 20% toluene and 80% Xylene?
	or	(c) Petroleum [ASTM: petroleum naphtha] or coal-tar naphtha?
		(d) Petroleum distillate
2.		Water content of solvent determined by running a blank using an equal amount of solvent
2		as used for the test sample?
3.		Still made of metal or glass?
4.		Gas burner or electric heater? (Ring burner used with metal stills)?
5.		Reflux condenser?
6.		Graduated glass trap?
		VERIFICATION
		VENITCATION
1.		Does the trap come with a manufacturer-supplied certificate stating accuracy and NIST traceability
		requirements have been met?
		or
2.		Have the accuracy of the graduation marks on the trap been verified by use of a NIST-traceable 5 mL
		Micro Burrette or Micro Pipette readable to the nearest 0.01 mL?
		(a) For trap styles A, B, C, and D verification performed as follows:
		(1) Verified at 0.1 mL intervals up to the 1.0 mL line?
		(2) Verified at 1.0 mL intervals up to the total volume of the trap?

Table 1: Trap Types			
	Bottom	Size of	
Trap Style	Shape	Trap	Divisions (mL)
Α	Conical	10	0.1 up to 1.0,
			0.2 up to 10.0
В	Conical	25	0.1
C	Conical	25	0.2
D	Conical	25	0.2
E (i)	Round	5	0.1
E (ii)	Round	5	0.05
E (iii)	Round	10	0.1
F	Round	2	0.05

Verified at 0.05, 0.5, 1.0, 1.5, and 2.0 mL lines?....

Verified at the 0.1, 1.0, 2.0, 4.0, and 5.0 mL lines?....

3.	The enti	re glassware assembly verified prior to first use and at regular frequency thereafter as follows:	
	(a)	400 mL of dry (0.02% water maximum) solvent, to be used for testing, tested in accordance with	the
		procedure below?	
		Contents of the trap discarded?	
COMME	ENTS (T	55/D95):	(T55/D95)

WATER IN BITUMINOUS MATERIALS

(T55)	
(D95)	

Date: _____

VERIFICATION (Continued)

(c)	Volume of water specified as the first test in Table 2 added to the distillation flask and the sample retested?
(d)	Contents of the trap again discarded?
(e)	Volume of water specified as the second test in Table 2 added to the distillation flask and the sample retested?
(f)	Trap readings are within the tolerances specified in Table 2?
(g)	If trap readings are not within tolerance, the device is checked for vapor leaks, too rapid boiling, or inaccuracies in verification of the trap and the steps above repeated?
	Note to Assessors: View at least one past record of verification of the glassware assembly.

Table 2: Trap Verification				
Trap Style	Test 1 Volume of Water (mL)	Permissable Limits for Test 1	Test 2 Volume of Water (mL)	Permissible Limits for Test 2
A	1	1 ± 0.1	9	9 ± 0.21
В	12	12 ± 0.2	24	24 ± 0.2
С	12	12 ± 0.2	24	24 ± 0.2
D	12	12 ± 0.2	24	24 ± 0.2
E (i)	1	1 ± 0.1	4.5	4.5 ± 0.1
E (ii)	₀ 1	2 ± 0.05	4.5	4.5 ± 0.05
E (iii)	5	5 ± 0.1	9	9 ± 0.1
F	1	1 ± 0.05	1.9	1.9 ± 0.05

AASHTO Materials Reference Laboratory

1.	Sample either measured in cylinder or weighed in still to accuracy of 1 percent?
2.	If cylinder is used, was it rinsed clean with total of 100 mL solvent?

3. If weighed into still, one 50-mL and two 25-mL solvent portions added to still?.....

	<u>PROCEDURE</u>
1.	Satisfactory assembly of components?
	Note to Assessors: Glass beads may be added to reduce bumping.
2.	Trap selected according to expected water content?
3.	Loose cotton (or similar) plug inserted in top of condenser?
4.	All connections vapor or liquid tight (caution if the apparatus leaks)?
5.	Cold water circulated in jacket of condenser?
6.	Heat applied and rate adjusted to have 2 to 5 drops distillate per second from the condenser?
7.	For metal stills, ring burner about 3 in. above bottom and lowered as test proceeds?
8.	Distillation continued until no water visible except in trap?
9.	No increase in water in trap for 5 minutes?
10.	Ring of water dislodged from condenser?
11.	Drops of water in trap dislodged?
12.	When cool, volume of water in trap read to nearest division?
13.	Water in the sample calculated?

COMMENTS (T55/D95):

(T55/D95)

DISTILLATION OF CUT-BACK ASPHALTIC PRODUCTS

(T'/8)	_
(D402)	

		APPARATUS Date:	
1	1711		
1.	Flasks	Flasks have a side arm?	
	(a)		
2.	(b)	500 mL capacity?	·····
۷.	Conder (a)	Jacket length of 200 to 300 mm?	
	(a) (b)	Tube length of 440 to 460 mm?	
	\ /	Glass jacket?	
3.	(c)	· ·	
3.	Adapte (a)		
	(a) (b)	Reinforced top?	
	` '	Inside outlet line vertical?	
	(c)		
	(d)	End out or ground?	
4	(e)	End 40 to 50° to inside line?	·····
4.	Shield		
	(a)	Made of 22 gauge steel?	
	(b)		
	(c)	Transparent mica windows?	
5.	(d)	Two piece top consisting of 6.4 mm mill board?	
<i>5</i> . 6.		nated Receiver 100 mL with 1 mL graduations or smaller receiver with 0.1 mL graduations	5 ?
0.	Heating	ng assembly (1) Coa human?	
		(1) Gas burner?	
		Two pieces of 1.18-mm gauze? Tripod or ring stand?	
		3. Shield for burner? (2) Electric heating mantle?	
		1. Connected to variable transformer?	
7.	Missoll	1. Connected to variable transformer?	
7.	(a)	Thermometers: Either ASTM 8C / 8F or IP 6C?	ratory
	(a) (b)	Flame snuffer for residue?	
	(c)	Suitable cork for flask?	
	(d)	Weighted receiver cover?	
	(e)	Tight joint between flask & condenser?	
	` '	Flask neck to adapter end: length adjustable to 650 ± 50 mm?	
	(f)	Class G2 balance?	
	(g) (h)	Residue container with slip-on cover, 75 ± 5 mm diameter and 55 ± 5 mm in height?	
	(11)	Residue container with sup-on cover, 13 ± 3 min diameter and 33 ± 3 min in height?	
COMN	MENTS (7	T78/D402):	(T78/D402)

DISTILLATION OF CUT-BACK ASPHALTIC PRODUCTS

(T'/8)	_
(D402)	

	PROCEDURE Date:
1.	Apparatus clean, dry, and assembled correctly?
2.	Thermometer properly aligned 6.4 mm above bottom of flask?
3.	Correction of nominal temperatures because of elevation or barometric
	pressure made to nearest 1°C (2°F) (The lab should show the tables and how they determine corrections)?
4.	Sample warmed if necessary and thoroughly stirred?
5.	If sample contains sufficient water to cause foaming or bumping, is it dehydrated at least 250 mL?
6.	Specific gravity of material known?
7.	Weight of 200 mL calculated from specific gravity at 15.6°C?
8.	200 mL of sample weighed into flask within 0.5 g?
9.	Apparatus assembled and water passed through condenser jacket?
10.	Area where test is performed free of drafts?
11.	Heat applied at start time:
12.	First drop leaves flask side arm within 10 ± 5 min. after heat first applied? First drop time:
13.	Drip rate from adapter 50 to 70 drops / min. up to 260°C (500°F)?
14.	Drip rate from adapter 20 to 70 drops / min. from 260 to 316°C (500 to 600°F)?
15.	Volume of distillate recorded at all specified corrected temperatures to 0.5 mL?
	Note: Temperatures are corrected based on the difference between local barometric pressure and 760 mmHg
	The nominal temperatures for recording (at 760 mmHg) are 190, 225, 260, and 316°C.
16.	During dry spells, rate of temperature increase 5°C / minute?
17.	Elapsed time from 316 to 360°C (600 to 680°F) less than 10 minutes?
18.	Heat cut off as soon as temperature reaches 360°C (680°F)?
19.	Flask and thermometer removed for pouring residue?
20.	From cut off of heat to start of pouring: not over 30 seconds?
21.	Side tube of flask substantially horizontal during pouring?
22.	Residue poured into 240 mL (8 oz.) container set on its cover?
23.	Skin pushed aside?
24.	Any residual distillate in condenser drained into receiver?
25.	Residue stirred and poured into test containers as soon as no further vaporization is apparent?
26.	When residue reaches $135 \pm 5^{\circ}$ C ($275 \pm 9^{\circ}$ F), poured for further testing?
COM	MENTS (T78/D402): (T78/D402
COM	(1/6)D+02.

COMMENTS (T79/D3143):

FLASH POINT DETERMINATION USING TAG OPEN CUP

(17/9)	
(D3143)	

		APPARATUS Date:
	Test cu	in.
	(a)	Outer Diameter at the base of the ring 53.2 to 57.0 mm?
	(b)	Overall height 50.0 to 53.2 mm?
	(c)	Base of ring to cup rim 7.1 to 8.7 mm?
	(d)	Mass less than 95 g?
	<u>Bath</u>	
	(a)	Made of copper?
	(b)	Equipped with a constant level overflow?
		ometer and holder
	(a)	Thermometer with a range of -5 to 110°C (20 to 230°F) and conforming to requirements
		for an ASTM 9C / 9F thermometer as prescribed in ASTM E1?
	(b)	ASTM: Or a PRT with a 3- or 4- wire design, 50mm (2 in) greater than immersion depth?
	(c)	Thermometer calibrated according to E77 (mercury) or E644 (PRT)?
	(d)	Holder capable of positioning thermometer as follows:
		(1) Vertically 6.4 mm from inner bottom of cup?
		(2) Midway between center and edge of cup?
		On line through center of cup and pivot of the taper?
	Ignitio	<u>n taper</u>
	(a)	Maintained in fixed horizontal plane by swivel device?
	(b)	Center of orifice 3.2 mm above upper edge of cup?
	(c)	Jet of taper at least 152 mm from center of swivel?
		Jet passes across center of cup at right angles to thermometer?
	(d)	
	(e)	Tip of taper approx. 1.6 mm [AMRL: ± 0.4 mm] in diameter?
	(f)	Flame size comparator bead on apparatus or hole in leveling device?
	(g)	If comparator bead is mounted on taper, the jet tip extends at least 3 mm beyond the bead?
	(h)	Diameter of bead or hole no more than 4 mm?
		ng device
	(a)	Made of suitable metal at least 3.2 mm thick?
	(b)	Two projections 25.4 mm apart and 3.18 ± 0.25 mm in length?
	(c)	Larger hole no more than 4 mm and centered 3.2 mm from bottom and 25.4 mm from end?
	(d)	Center of smaller hole 3.2 mm from bottom of level and 63.5 mm from end of level?
	Heater	•
	(a)	Small gas burner?
r	(b)	Electric heater with variable transformer?
		laneous
	(a)	One of the following available:
	(4)	(1) Draft-free hood in flash room?
		(2) Draft shield?
	(b)	
	(b)	Bath liquid:
		(1) Water for flash points to 79.5°C?
		(2) 1:1 water-glycol solution for points above?
	<i>(c)</i>	ASTM: Cleaning Solvents:
		(1) Technical grade and capable of cleaning and drying test cup?

(T79/D3143)

FLASH POINT DETERMINATION USING TAG OPEN CUP

(T79)	
(D3143)	

	PROCEDURE Date:
1.	Room temperature: 25 ± 5°C?
2.	Tester placed within draft shield or draft-free fume hood?
3.	Tester shielded from strong light?
4.	Test bath filled with either water or 1:1 water-glycol solution?
5.	Temperature of bath media at least 16.5°C (30°F) [ASTM: 10°C (18°F)] below expected flash point?
6.	Glass test cup placed in bath?
7.	Bath filled to approximately 1/8 in. from the top when test cup in place?
8.	Leveling device rested on rim of cup?
9.	Cup filled until the level just touches the pointer of leveling device?
10.	Taper lit and flame adjusted to less than 4 mm in diameter?
11.	AASHTO: Heat applied to bath and temperature rise of sample is $1.0 \pm 0.3 \text{C} (2.0 \pm 0.5 \text{F})$ per minute?
12.	ASTM: Heat applied to bath and temperature rise of sample is $1^{\circ}C$ ($2^{\circ}F$)/min ± 6 seconds?
13.	Sample level adjusted at 10 to 15°C (18 to 27°F) below expected flash point?
14.	Test flame passed across sample at the same temperature?
15.	Test flame applied at successive 1°C (2°F) intervals?
16.	Passed across cup only once at each temperature?
17.	Each pass requires about 1 second?
18.	Flame passed across sample in continuous motion?
19.	Direction reversed with each pass of test flame?
20.	Temperature at time of first distinct flash in interior of test cup recorded as flash point?
21.	ASTM: Recorded flash corrected for barometric pressure if the pressure differs
	from 101.3 kPa (760 mm Hg)?
	Note to Assessors: Aneroid barometers (those used at weather stations and airports) are pre-corrected to
	give sea-level readings and shall not be used. ASTM: Corrected flash reported to the nearest 0.5°C (1°F)?
22.	ASTM: Corrected flash reported to the nearest 0.5°C (1°F)?

COMMENTS (T79/D3143): (T79/D3143)

THIN-FILM OVEN TEST

(T179)	
(D1754)	

Date: _____

|--|

Sample containers	1	2	3	4	5	6	7	8
Diameter: 140 mm?								
Depth approximately 9.5 mm?								
Thickness 0.64 mm [AMRL: at least 0.38 mm]?								
Made of stainless steel?								
Bottom not badly warped?								

1.	<u>Ovens</u>						
	(a)	AASHTO: Inside at least 330 x 330 x 330 mm (13 x 13 x 13 in.)					
	(b)	ASTM: Maximum dimensions of 535 x 535 x 535 mm?					
	(c)	Tightly fitted hinged door?					
	(d)	Hinged door has a window at least 100 x 100 mm (4 x 4 in.), clear, 2 sheets of glass?					
	(e)	Air vents (Assessors check to feel air current with hand over vent)?					
	(f)	Shelf					
		(1) Approximately centered in oven (ASTM: 6 in. above bottom)?					
		(2) Minimum diameter 250 mm (<i>ASTM: Maximum</i> = 450 mm)?					
		(3) Rotates at $5.5 \pm 1 \text{ rpm}$?					
		(4) Made of metal?					
		(5) Air can flow through shelf when samples in place?					
		(6) Shelf constructed or marked so that sample containers can be placed in same					
		position for each test?					
		(7) Each container position symmetrical with respect to shaft and any holes in shelf?					
		(8) ASTM: Maximum tilt during rotation no more than 3% from horizontal?					
	(g)	ASTM 13C thermometer (ASTM: or PRT conforming to ASTM E1137 verified accurate to 163 °C)?					
	1	(1) Readable through door?					
	/ /	(2) Can be positioned vertically?					
		(3) Approximately 6.4 mm (ASTM: 40 mm) above shelf?					
		(4) At midpoint of shelf radius (ASTM: Radially centered over a sample container position?)?					
	(h)	Oven capable of maintaining temperature of 163 ± 1°C?					
	(i)	Oven capable of returning to 162°C within 15 min. after introducing two sample pans?					
2.	Miscell						
	(a)	Rigid heat-resistant insulation board(s) (for reheating of residue)?					
	(b)	Spatula or putty knife?					
	(c)	Class B (0.001 g) balance available if the loss on heating is desired?					
	(d)	240 mL (8 oz.) ointment tin?					
	(e)	Class G2 (0.1 g) balance available to weigh the pans and sample?					

COMMENTS (T179/D1754):

(T179/D1754)

THIN-FILM OVEN TEST

(TT/9)	
(D1754)	

	PROCEDURE Date:
1.	Sample heated to less than 150°C (302°F), stirring frequently with thermometer while avoiding the incorporation of air bubbles
2.	AASHTO: Oven temperature control adjusted so that thermometer reads $163 \pm 1^{\circ}$ C ($325 \pm 2^{\circ}$ F) while
۷.	oven is at equilibrium (with empty sample containers in predetermined positions)?
3.	ASTM: Oven preheated to setting for at least two hours prior to testing?
<i>4</i> .	50.0 ± 0.5 g of material weighed into each of two or more tared TF pans?
5.	At same time, original asphalt properties samples prepared (if desired)?
6.	Samples in TF pans cooled to room temperature?
7.	If percent loss is desired, each TF sample weighed to nearest mg (0.001 g)?
8.	TF pans with sample quickly placed in oven in predetermined positions on shelf after removing empty pans?
9.	Vacant positions on the shelf filled with empty sample containers so every position is occupied?
10.	Oven closed and shelf rotated?
11.	Samples removed 5 hours after oven recovered to 162°C?
12.	Oven holds at 163 ± 1°C?
13.	Total time in oven not more than 5 hours and 15 minutes?
14.	If change in mass is desired:
	(a) Samples cooled to room temp. and weighed to nearest mg (0.001 g)?
	(b) Samples placed on refractory boards and returned to oven?
	(c) Oven closed, shelf rotated for 15 ± 2 minutes, and then samples removed?
15.	ASTM: Oven preheated to setting for at least two hour prior to testing?
16.	ASTM: Pans removed individually and scraped into the ointment tin?
17.	ASTM: Oven door closed, heater power on, and shelf rotating during scraping?
18.	ASTM: Final pan removed within five minutes of the initial pan?
19.	AASHTO: Almost all residues removed from pans by scraping?
20.	AASHTO: Placed in 240 mL (8 oz) container?
21.	Combined residues thoroughly mixed by stirring (AASHTO: a hot plate may be used)?
22.	Material poured into containers and molds for further testing? Tests on residue completed within 72 hours?
23.	Tests on residue completed within 72 hours?
COMM	ENTS (T179/D1754): (T179/D1754)

KINEMATIC VISCOSITY TEST

(T201)	
(D2170)	

				APF	PARATUS	Date:				
		T.74								
1.		Viscon								
		(a)								
		(b)	Viscometer calibration of			DOWNE.				
		(c)			, Lantz – Zeitfuchs:	, BS/IP/RF:,				
		<i>(</i> 1)	Zeitfuchs – Cross Arm:							
		(d)			pipette available?					
		(e)		n good condition?						
2.		Cleane								
		(a)			not set above 500°C or a					
					TM: only if deposits are ob					
		(3) Residue-free acetone?								
		(4) Clean dry air?								
		(5) Completely miscible solvent?								
				latile solvent?						
3.		Pre-he	ating apparatus							
		(a)	Oven or bath at $63 \pm 3^{\circ}$	C $(145 \pm 5^{\circ}F)$ for σ	cutbacks?					
		(b)	Means of heating to 135	5 ± 5 °C (275 ± 10°	F) for asphalt cements?	•••••				
4.		Viscon	neter bath or baths							
		(a)		est temperature ± 0	.1°C (0.2°F)?					
		(b)			02 F)?					
		(c)			vary by more than ±0.1°C					
		(6)			th of the viscometer, or fr					
					at 60°C (140°F)?					
		(d)			nple reservoir or top of ca					
		` /	(e) Thermometer correctly immersed at test temperature?							
		Note: Distilled water has been found to be an acceptable bath liquid for tests at 140°F and white oil has been found to be an acceptable bath liquid for tests at 275°F. If other liquids are used, please note below.								
		veen jo	una io be an accepiable i	am uqua jor iesis	s at 275 F. If other tiquia	s are usea, piease noie i	below.			
5.		Thomm	ometers							
٥.				25C / 25E C	4 1 400E					
		(a)			s at 140°F					
	or	(b)								
	or	(c)			curacy of at least 0.04°F (
		(d)		levice calibrated at	least every six months?	•••••				
6.		<u>Timers</u>	<u> </u>							
		=					-			
				1/10 s	Accurate to 0.45 s in					
			Manufacturer	graduations?	15 min?	Serial Number				
				gradations.	10 111111					
		L								
		L. L		L			_			
		(a)	•	6 months?						
7.		<u>Miscellaneous</u>								
		(a) Air-tight containers of about 30 mL capacity for cutbacks?								
		(b)			cements?					
		(c)	Provision for filtering, d	lrying air when cle	aning tubes?					
CC	MM	ENTS (Γ201/D2170):				(T201/D2170)			

KINEMATIC VISCOSITY TEST

(T201)	
(D2170)	

	PROCEDURE Date:	
1.	What type of samples does the laboratory test? (a) Cutback Asphalt?	
	(b) Asphalt Cement?	
	(c) Both?	
	Note: If only cutback asphalt is tested, please write an informational note on the report.	
2.	Sample heated and stirred until sufficiently fluid to pour?	
	<u>Cutbacks</u> : Sample stirred at room temp. for 30 sec. [if necessary, sealed container placed in	
	bath or oven at 145 ± 5 °F (63 ± 3 °C) until fluid enough to stir] and viscometer	
	charged immediately?	
3.	Minimum of 20 mL transferred to a suitable container?	
	<u>Cutbacks</u> : If not tested immediately, 20 mL sample sealed?	
4.	For pouring, 20 mL sample heated to $275 \pm 10^{\circ}F$ ($135.0 \pm 5.5^{\circ}C$)?	
	<u>Cutbacks</u> : For pouring, if necessary (materials with viscosities above 800 cSt at 140°F) 20	
	ml sealed container heated in bath or oven at 145 ± 5 °F (63 ± 3 °C) for not more than 30 min	
5.	Viscometer selected is clean and dry?	
6.	Viscometer used preheated to test temperature?	
7.	Charged correctly according to design of instrument?	
8.	Viscometer remains in the bath for minimum 10 min. and maximum 30 min.?	
9.	Flow started as prescribed by design of instrument?	
10.	Efflux time measured to within 0.1 seconds?	
11.	If efflux time is less than 60 seconds, is tube with smaller capillary chosen and operation repeated?	
12.	Upon completion of the test, is the tube cleaned according to Sec. 8.8 (either solvent or a glass cleaning oven not set above 500°C)?	
13.	Kinematic viscosity calculated to three significant figures by <i>Time X Calibration Factor</i> ?	···
COMM	FNTS (T201/D2170)· (T201	/D2170)

AASHTO Materials Reference Laboratory

ABSOLUTE VISCOSITY TEST

(T202)	
(D2171)	

		APP	<u>PARATUS</u>	Date:		
Visco	meters					
a)	Viscometer Number:					
b)	Viscometer calibration co	onstant:				
c)				Modified Koppers:		
d)						
Clean		C				
a)		ass cleaning oven	not set above 500°C or a	solvent based system as below		
				served within the viscometer]?.		
	(2) Distilled water?					
	(3) Residue-free ace	etone?				
	(4) Clean dry air?					
	(5) Completely misc	cible solvent?				
/isco	<u>meter bath</u>					
a)	AASHTO: Maintains test	t temperature ± 0	.1°C (0.2°F)?			
b)	ASTM: Test temperatur	$e \pm 0.01$ °C (± 0.0	02 F)?			
c)	Temperature of the bath n	nedium does not	vary by more than ±0.1°C	C (±0.2°F)		
	[ASTM: ± 0.03 °C (± 0.05 °	F)] over the leng	th of the viscometer, or fr	rom viscometer		
	to viscometer in the vario	ous bath positions	at 60°C (140°F)?			
d)	Top timing mark at least 2	20 mm below sur	face of liquid?			
e)						
acur	um system					
a)	Capable of maintaining vacuum of 300.0 ± 0.5 mm?					
b)	Holds vacuum when system is evacuated and closed?					
c)	Standardized at least once	e a year?				
heri	mometer					
a)	ASTM 47F or 47C?		$C + 1 \rightarrow O O AOT $	0.000000		
a) b)	Any other thermometric of					
a) b) c)	Any other thermometric of Thermometer or other de-			0.02°C)?		
a) b) c)	Any other thermometric of Thermometer or other de-					
a) b) c)	Any other thermometric of Thermometer or other desers	vice calibrated at	least every six months?			
a) b) c) F ime r	Any other thermometric of Thermometer or other de-					
a) b) c)	Any other thermometric of Thermometer or other desers	vice calibrated at	least every six months? Accurate to 0.45 s in			
a) b) c)	Any other thermometric of Thermometer or other desers	vice calibrated at	least every six months? Accurate to 0.45 s in			
a) b) c)	Any other thermometric of Thermometer or other desers	vice calibrated at	least every six months? Accurate to 0.45 s in			
a) b) c) T ime r	Any other thermometric of Thermometer or other decrees Manufacturer	vice calibrated at 1/10 s graduations?	Accurate to 0.45 s in 15 min?	Serial Number		
a) b) c) 'ime r	Any other thermometric of Thermometer or other desers Manufacturer Timers calibrated every 6	vice calibrated at 1/10 s graduations?	Accurate to 0.45 s in 15 min?			
a) b) c) 'ime r 'lisce	Any other thermometric of Thermometer or other desers Manufacturer Timers calibrated every 6 cellaneous	1/10 s graduations?	Accurate to 0.45 s in 15 min?	Serial Number		
a) b) c) ime Misce a)	Any other thermometric of Thermometer or other descriptions Manufacturer Timers calibrated every 6 cellaneous Means for melting asphal	1/10 s graduations? months?	Accurate to 0.45 s in 15 min?	Serial Number		
a) b) c) imen Alisce a) b)	Any other thermometric of Thermometer or other descriptions Manufacturer Timers calibrated every 6 Ellaneous Means for melting asphal Containers 20 mL or large	1/10 s graduations? months? t cement available available?	Accurate to 0.45 s in 15 min?	Serial Number		
a) b) c) imen Misce a) b) c)	Any other thermometric of Thermometer or other desers Manufacturer Timers calibrated every 6 ellaneous Means for melting asphal Containers 20 mL or large Oven or bath at 275 ± 10°	1/10 s graduations? f months? t cement available available?	Accurate to 0.45 s in 15 min?	Serial Number		
a) b) c) imen Aisce a) b) c) disce d)	Any other thermometric of Thermometer or other description. Manufacturer Manufacturer Timers calibrated every 6 cellaneous Means for melting asphal Containers 20 mL or large Oven or bath at 275 ± 10 Means for drying and filter.	1/10 s graduations? t cement available available? F available?	Accurate to 0.45 s in 15 min? e?	Serial Number		
a) b) c) imen Misce a) b) c)	Any other thermometric of Thermometer or other description. Manufacturer Manufacturer Timers calibrated every 6 gellaneous Means for melting asphal Containers 20 mL or large Oven or bath at 275 ± 10 Means for drying and filte No. 50 (300-µm) sieve? (1/10 s graduations? t cement available er available? °F available? ering air stream a optional)	Accurate to 0.45 s in 15 min? e?	Serial Number		

ABSOLUTE VISCOSITY TEST

(T202)	
(D2171)	

	PROCEDURE Date:
1.	Sample heated and stirred until sufficiently fluid to pour?
2.	The maximum temperature shall not exceed 100°C (180°F) above expected softening point?
3.	At least 20 mL poured into suitable container?
4.	If suspected that sample contains solid material, sample strained through No. 50 (300-mm) sieve?
5.	Asphalt heated to $275 \pm 10^{\circ} F$ ($135 \pm 5.5^{\circ} C$), stirring occasionally?
6.	Clean, dry viscometer selected for flow time exceeding 60 seconds?
7.	Viscometer preheated at 275 ± 10°F (135 ± 5.5°C)?
8.	Charged to within 2 mm of fill line?
9.	Maintained in preheated oven or bath at $275 \pm 10^{\circ}$ F ($135 \pm 5.5^{\circ}$ C) for 10 ± 2 minutes?
10.	Within 5 minutes of removing the tube from the oven or bath, sample placed in testing bath?
11.	300.0 ± 0.5 mm of mercury vacuum established?
12.	Manometer properly zeroed?
13.	Allowed to sit in bath for 30 ± 5 minutes?
14.	Flow started?
15.	Flow between successive timing marks determined to 0.1 seconds?
16.	First flow time exceeding 60 seconds recorded and marks identified?
17.	Viscometer drained in oven (optional)?
18.	Upon completion of the test, is the tube cleaned according to Sec. 8.1.9 (either solvent or a glass cleaning
	oven not set above 500°C)?
19.	Viscosity calculated to three significant figures by <i>Time X Calibration Factor</i> ?
20.	Temperature and vacuum pressure recorded?
COMM	ENTS (T202/D2171): (T202/D2171)

AASHTO Materials Reference Laboratory

DENSITY (PYCNOMETER METHOD)

(T228)
(D70)

Date: _____

AP	PΑ	\mathbb{R}	AΤ	U	S

PYCNOMETERS	1	2	3
Made of Glass?			
Capacity of 24 to 30 mL?			
Weight not more than 40 g?			
Ground glass stopper diameter 22 to 26 mm?			
Stopper top smooth plane?			
Stopper hole diameter 1.0 to 2.0 mm?			
Bottom of stopper concave?			
Concavity 4.0 to 18.0 mm high at center?			
Pycnometer calibrated?			
Physical condition OK?			
Bottom edge of stopper not chipped?			
Edges of hole in stopper not chipped?			

Note to Assessors: Frosted glass is acceptable, glass etching is unacceptable.

771				
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111		UIII		. o.

1.	Liquid in glass, total immersion thermometer readable to at least 0.1°C or 0.2°F (ex. 63C or 63F)?
or	
2.	Any other thermometric device with at least 0.1°C or 0.2°F subdivisions?
3.	Thermometer calibrated?
4.	Maximum scale error of 0.1°C?
Beakers	
1.	Capacity of 600 mL or larger?
2.	Filled with enough freshly boiled distilled or deionized water at test temperature to
	allow the top of the pycnometer to be immersed to a depth of at least 40 mm?
3.	Bottom of beaker immersed in the water bath at least 100 mm without the top being submerged?
4.	Any convenient method used to insure the beaker does not tip over or restrict the bath circulation (ASTM: beaker clamped in place)?
Water B	<u>ath</u>
1.	Capable of ± 0.1 °C at test temperature?
Balance	
1.	Class B (0.001 g)?
COMM	ENTS (T240/D2872): (T240/D2872)

DENSITY OF SEMI-SOLID BITUMINOUS MATERIALS (PYCNOMETER METHOD)

(AASHTO T228)
(ASTM D70)

PR			

	PROCEDURE Date:				
1.	Sample heated and stirred avoiding incorporation of air bubbles until sufficiently fluid to pour and for				
	not more than 60 min. over flame or hotplate, or 2 hours in an oven?				
2.	ASTM: Sample not heated for more than 60 minutes and 110 ${\mathcal C}$ (230 ${\mathcal F}$) above expected softening point?				
3.	Pycnometer calibrated? (A, B)				
4.	Lab says that calibration was performed by Sec. 10? (Steps 12-18 below)				
5.	Pycnometer clean and dry?				
6.	Pycnometer warmed?				
7.	Pycnometer filled about ³ / ₄ full?				
8.	Sample does not touch sides above final level?				
9.	Air bubbles avoided?				
10.	If air bubbles are present, are they removed with a Bunsen flame?				
11.	Pycnometer cooled to ambient temperature for at least 40 min.?				
12.	Mass of pycnometer and sample determined? (C)				
13.	Pycnometer filled with freshly boiled distilled or deionized water at test temperature and stoppered without allowing any air bubbles to remain in the pycnometer?				
14.	Pycnometer placed in beaker and transferred to the bath [ASTM: beaker clamped in place]?				
15.	Immersed in the water bath for at least 30 minutes?				
16.	Top of pycnometer stopper immediately dried with one stroke of towel, no redrying?				
17.	Outside of pycnometer dried?				
18.	Mass of pycnometer, sample, and water determined? (D)				
19.	All masses determined to the nearest 0.001 g?				
20.	Lab says that book formula used in calculating specific gravity?				
	S.G. = (C - A) / [(B - A) - (D - C)]				
	A = weight of pycnometer $C = $ weight of pycnometer $+ $ asphalt				
	B = weight of pycnometer + water $D = weight of pycnometer + asphalt + water$				
	Specific gravity reported to 0.001?				
21.	Specific gravity reported to 0.001?				

COMMENTS (T228/D2872):

(T240/D2872)

ROLLING THIN-FILM OVEN

(1240)	
(D2872)	

		<u>APPARATUS</u>	Date:
Oven			
Oven:	Chambe	er e	
1.	(a)	Height of 381 mm (15 in.) [± 12.7 mm (1/2 in.) AMRL]?	
	(b)	Width of 483 mm (19 in.) [± 12.7 mm (1/2 in.) AMRL]?	
	(c)	Depth of $445 \pm 13 \text{ mm} (17 \frac{1}{2} \pm 1/2 \text{ in.})$?	
	(d)	Floor and ceiling vented?	
2.	Plenum	· · · · · · · · · · · · · · · · · · ·	
	(a)	(ceiling and walls only) 38 mm (1 1/2 in.) [± 12.7 mm (1/2 in.) AMRL]?	
3.	Windov		
	(a)	Two sheets of heat-resistant glass providing an unobstructed view to interior?.	
	(b)	Window height 203 mm to 229 mm (8 to 9 in.)?	
	(c)	Window width: 305 mm to 330 mm (12 to 13 in.)?	
4.	Heater		
	(a)	ASTM: Top of heating element $1 \pm 1/8$ in. $(25 \pm 3 \text{ mm})$ below oven floor?	
5.	Fan		
	(a)	Outside diameter of fan 133.4 mm (5 1/4 in.) [± 12.7 mm (1/2 in.) AMRL]?	
	(b)	Fan height 73 mm (2 7/8 in.) [± 12.7 mm (1/2 in.) AMRL]?	
	(c)	Centered in width of oven?	
	(d)	Blows air into chamber?	
	(e)	Carriage face to axis distance of 152 mm (6 in.) [± 9.52 mm (3/8 in.) AMRL]?	?
	(f)	Fan set so that it turns in an opposite direction to its vanes?	
6.	Sensor		
	(a)	The sensing element of the thermostat may be placed at any location the oven	
		maintain temperature control?	
7.	Thermo		
	(a)	ASTM 13C thermometer?	······ <u> </u>
	(b)	Distance from thermometer bulb to right side of the oven 51 mm (2 in.)	laboratory
		[± 9.52 mm (3/8 in.) AMRL]?	
	(c)	Bulb level is approximately mid-depth of the oven?	
	(d)	Bulb level with carriage axis within 25 mm (1 in.)?	
	(e)	(Optional) - Optically transparent polymer sheath having a maximum	
		thickness of 0.25 mm (0.01 in.)?	
		Note to Assessors: If a sheath is used, it shall be installed with substantial ph	•
	0.14	sheath and the thermometer. The thermometer should be standardized after in	nstatiation of a sneath.
8.	<i>or</i> Electron	nic Temperature Measurement System	
0.	(a)	Electronic measurement circuitry includes a digital display having a resolution	
	(a)	of 0.1°C (0.2 °F) or better?	
	(b)	AASHTO: Standardized according to the interval in AASHTO R18?	
	(c)	AASHTO: Tip of the sensor located in the same manner as the bulb of a merc	
	(d)	ASTM: Sensor is 3 or 4 wire, Grade A Platinum Resistance Thermometer	
	(4)	conforming to the requirements of Specification E 1137?	······ _
COMM	IENTS (T	Г240/D2872):	(T240/D2872)

Revised 2014-06-02

ROLLING THIN-FILM OVEN

(T240)	_
(D2872)	_

		<u>APPA</u>	<u>ARATUS</u>	(Continu	ed)		Da	ate:		
	Electro	nic Temperature Measurement System	(Continu	ied)						
	(e) ASTM: Calibrated as a unit containing both the temperature sensor and the									
		electronic measurement circuitry pri							•••••	
	(f)	ASTM: Verified annually as a unit								
		operating temperature while the veri								
	()	with the normal test sensor?								
	(g)	ASTM: Calibrations and verification								
		Note: If the difference between the ve fication will be regarded as having far								
9.	Carriag		иеи, ини	іне іетр	reraiure i	neusurei	neni sysi	ет знан	ve recui	wraiea.
	(a)	Diameter of 305 mm (12 in.) [± 12.7	mm (1/2	in.) AMI	RL1?					
	(b)	Holds 8 containers?								
	(c)	Rotation rate 15.0 ± 0.2 rpm (15 rev/o								
	(d)	Face of carriage to back oven wall is								
10.	Air Jet	C		`	, -	`	ŕ			
	(a)	Copper tube, not covered by foil or or								
	(b)	Orifice diameter 1.02 mm (0.04 in.) a								
		Assessors: The orifice diameter openi	ng can b	e checked	d by the lo	aborator	y if it app	ears to b	e blocke	d using
	a #60 d									
	(c)	Distance from Orifice to open end of $(1/4 \pm 1/16 \text{ in.})$ [ASTM: 1/4 in. (6.4)	mm)]?							
	(d)	Directed along the container axis and	blowing	into the	center of	the conta	iner oper	ning?		
11.	Flow M									
	(a)	Standardized at least every 12 months	s [ASTM	: calibra	ited perio	dically] [*]	?	•••••		
12.	Air Sup									
	(a)	Regulated, dust free, and equipped w								
12	(b)	Flow meter rate set at 4000 mL/min o								
13. 14.	Cooling	melt sample?	H-AI-	ang-in	renei			immi.	arron	·y
14.	(a)	Wire or sheet metal and made of alun	ninum or	ctainless	steel?					
	(a) (b)	Allows containers to cool horizontally					•••••	•••••	••••••	•••
	(0)	to cool in the same horizontal plane								
	(c)	At least 25 mm (1 in.) clearance betw								
Sa	mple Con	tainers	1	2	3	4	5	6	7	8
Le	ength: 138	3.2 to 141.2 mm (5.44 to 5.56 in.)?								
		o 65.2 mm (2.473 to 2.567 in.)?								
		0.3 to 33.3 mm (1.19 to 1.31 in.)?								
_		0.7 to 2.3 mm (0.03 to 0.09 in.)?								

COMMENTS (T240/D2872):

(T240/D2872)

COMMENTS (T240/D2872):

ROLLING THIN-FILM OVEN

(1240)	
(D2872)	

	APPARATUS (Continued) Date:	
15.	Balances	
	(a) Class B (0.001 g) for change in mass?	
	(b) Class G2 (0.1 g) for weighing residue and sample?	
	(c) AASHTO: Standardized every 12 months (in accordance with R18)?	
16.	AASHTO Only: Electronic Level (Optional)	
	(a) At least 125 mm (5 in.) long?	
	(b) $30 \pm 3 \text{ mm } (1.2 \pm 0.1 \text{ in.}) \text{ wide?}$	
	(c) Bearing surfaces ground flat?	
	(d) Hold button the freeze the display?	
	<u>PROCEDURE</u>	
Oven	n Preparation:	
1.	AASHTO: Oven adjusted so that the horizontal axis of the glass containers are level within $\pm 1.0^{\circ}$?	
	Note: T240 recommends that the levelness of the carriage is checked by the procedure found in Appendix	Α.
2.	AASHTO: Carriage bearing checked for wear every 6 months and when the levelness of the	
	carriage is checked?	
3.	Fan started (The fan shall remain on whenever the oven heater is on and the oven door is closed.)?	
١.	Oven preheated at least 2 hours [ASTM: 16 hours]?	
5.	Temperature adjusted such that $163.0 \pm 1.0^{\circ}$ C ($325 \pm 1.8^{\circ}$ F) [ASTM: $163.0 \pm 0.5 \%$ ($325 \pm 1 \%$)]	
	is maintained during testing?	•••••
² repa	aration of the Sample:	
l.	As received, sample free of water?	
2.	Melted in oven set not over 163°C (325°F) [ASTM: 150°C (302°F)]?	
3.	Heated only long enough to be completely fluid in a container with the cover loose?	
١.	Manually stirred without incorporating air bubbles prior to pouring?	
5.	35.0 ± 0.5 g poured into each of the required containers?	ļ:A
) .	Immediately after pouring, sample container turned to a horizontal position?	
7.	Cylinder rotated slowly at least one full rotation, attempting to pre-coat its	
	cylindrical surface (Note: complete pre-coating may not be possible for certain binders)?	
3.	Care taken to prevent the sample from flowing out of the container?	
).	AASHTO: Sample does not coat the central part of the open end of the container?	
0.	Placed in a clean cooling rack for 60 to 180 minutes? (Start Time:)	
11.	Cooling rack in draft-free location at room temperature, away from ovens or other sources of heat?	• • • • •
12.	Two separate bottles used for change in mass determination?	
13.	Bottles weighed in a vertical position to at least the nearest 0.001 g (after cooling period)?	

(T240/D2872)

ROLLING THIN-FILM OVEN

(T240)	
(D2872)	

	PROCEDURE (Continued) Date:
Polling	Thin-Film Oven Test:
1.	Oven at temperature prior to loading sample bottles?
2.	Sample bottles placed in carriage so it is balanced?
3.	Unused spaces filled with empty containers (all containers must be filled for referee test)?
4.	Door closed and carriage/fan started?
5.	Air flow started at a rate of 4000 ± 300 mL/min [ASTM: ± 200 mL/min]? (Start Time:)
6.	Temperature recovery within 10 minutes $163.0 \pm 1^{\circ}\text{C}$ (325.0 ± 1.8°F)
	[ASTM: $163.0 \pm 0.5 \%$ (325 $\pm 1 \%$)] (at thermometer)?
7.	Samples removed 85 minutes after being placed in oven?
	Note to Assessors: Remember to have a timer ready to time scraping sample bottles starting after change-in-mass
	bottles are removed from the oven.
Change-	In-Mass Samples:
1.	Samples removed first 85 minutes after being placed in oven, placed horizontally in a
	cooling rack, and allowed to cool for 60 to 180 minutes? (Start Time:)
2.	Bottles weighed in a vertical position to at least the 0.001 g?
3.	Percent change in mass calculated (see equation next page)?
4.	Noted (AASHTO: and reported) if any sample flowed out of the bottle?
5.	AASHTO: If any sample flowed out of the container, that container is not used for mass loss determination?
6.	AASHTO: Two containers used for referee testing?
Residue	Samples:
1.	Bottles removed one at a time (After change in mass bottles)?
2.	Free flowing residue poured into a container?
3.	Remainder of residue in bottle scraped (Circumferential scraping recommended)?
4.	Average of 90% or more of residue scraped out of bottles?
5.	While residue is being transferred, is oven door closed, power, air, and carriage all on?
6.	Final container removed within 5 minutes of the initial container?
7.	All residues poured into one container?
8.	Container at least 30 % greater than total expected volume of residue?
9.	Residue in container stirred gently without incorporating air bubbles?
10.	ASTM: Residue tested within 72 hours or discarded?
Percent	Change-In-Mass (if performed)
	$(\mathrm{M_i}-\mathrm{M_f})$ / $\mathrm{M_b}$
	M_i = Initial mass of bottle + sample
	M_f = Final mass of bottle + sample
	$M_b = Mass of sample$
1.	Percent change in mass reported to the nearest 0.001 percent?
2.	Mass loss reported as a negative number, mass gain reported as a positive number?
COMM	ENTS (T240/D2872): (T240/D2872)

SPECIFIC GRAVITY OR API GRAVITY OF LIQUID ASPHALT BY HYDROMETER METHOD

(1295)	
(D3142)	

Date: _____

<u>APPARATUS</u>

1. <u>Hydrometers</u>

COMMENTS (T295/D3142):

TABLE 1 Recommended Hydrometers (Partial Listing)

ASTM	Туре	Units	Range	;		Scale
Designation	Туре	Omts	Total	Each unit	Interval	Accuracy
1H to 4H	long plain form	API	-1 to 41	12	0.1	0.1
21H to 28H	short plain form	API	0 to 41	6	0.1	0.1
85H to 90H	long plain form	Sp. Gr. 60/60F	0.800 to 1.100	0.050	0.0005	0.001
105H to 108H	intermediate plain form	Sp. Gr. 60/60F	0.800 to 1.000	0.050	0.001	0.001
315H to 320H	long plain form	Kg/m ³	800 to 1100	50	0.5	0.5

(a)	Thermometer with maximum scale error of 0.1°C (0.25°F) [such as ASTM 12C/12F or
	IP 64C/64F] or any other thermometric device of equal accuracy?
(b)	ASTM: Thermometer standardized every 12 months?
(c)	ASTM: If not conforming to requirements for 12C/12F, calibrated
	by ASTM Test Method E220?
(d)	ASTM: If thermohydrometer used, temperature scale range from
	20 to 65°C or 60 to 220°F (designation H)?
Hydr	ometer cylinder
(a)	Made of glass, plastic, or metal?
(b)	Made of glass, plastic, or metal? If plastic: resistant to discoloration?
(c)	Inner Diameter at least 20 mm (¾ in) greater than Outer diameter of hydrometer used?
(d)	Hydrometer floats with at least 25 mm (1 in) clearance between bottom of hydrometer
\	and bottom of cylinder? r bath
Wate	
(a)	At test temperature within ± 0.5 °C?
(b)	Depth approximately same as sample in hydrometer cylinder?
(c)	Thermometer or electronic temperature measuring device accurate to 0.25°C (0.5°F)?
Oven	<u>s</u>
(a)	At test temperature within ± 3°C?
(b)	Thermometric device accurate to 1°C (2°F)?

(T295/D3142)

SPECIFIC GRAVITY OR API GRAVITY OF LIQUID ASPHALT BY HYDROMETER METHOD

(T295)	
(D3142)	

<u>APPARATUS</u>	Date:

1. Select test temperature in accordance with Section 9 (see table below)?.....

Recommended Testing Temperatures

Grade	Testing Temperature °C (°F)
MC-30	Room
SC-70, MC-70, RC-70	40 (104)
SC-250, MC-250, RC-250	60 (140)
SC-800, MC-800, RC-800	80 (176)
SC-3000, MC-3000, RC-3000	100 (212)

2.	Samp	le heated in oven to within ± 3°C (5°F) of test temperature?							
3.	Sample container loosely covered?								
4.	Hydrometer, cylinder, and thermometer brought to test temperature?								
5.		le transferred to clean hydrometer cylinder carefully?							
6.	Air b	ubbles removed from sample surface by touching with a piece of clean filter paper?							
7.	If me	tal cylinder, level within 5 mm (1/4 in) of top of cylinder?							
8.	If test	is performed at other than room temperature:							
	(a)	Constant temperature bath within ± 0.5 °C (1.0°F) of test temperature?							
	(b)	Sample in cylinder placed in constant temperature bath?							
	(c)	Sample temperature equilibrated with bath temperature?							
	(d)	Thermometric device immersed properly in sample?							
	(e)	Sample temperature steady and recorded to nearest 0.2°C (0.5°F)?							
9.	Hydro	ometer gently lowered into sample, avoiding wetting stem above the immersion level?							
10.									
11.	Air b	ometer becomes completely stationary?ubbles allowed to rise to sample surface?							
12.	Hydro	ometer floats freely, not touching cylinder wall?							
13.		ometer read to nearest scale division?							
14.	Observation corrected for meniscus height?								
15.	Samp	le immediately stirred, keeping thermometric device correctly immersed, and temperature observed?							
16.	Temp	erature recorded to nearest 0.2°C (0.5°F)?							
17.	If tem	perature differs from previous reading by more than 0.5°C (1.0°F) is hydrometer							
	readii	ng repeated until the temperature becomes stable within 0.5°C (1.0°F)?							
18.		opriate corrections made to the observed hydrometer reading?							
19.		corrected hydrometer reading recorded to the nearest 0.1° API, 0.001 specific gravity, or 0.5 kg/m ³ ?							
20.		temperatures observed (before and after) recorded to the nearest 0.5°C (1.0°F)?							
21.	Corre	cted values converted to standard temperature using Guide D1250 and ASTM D1250 (Vol. 5.01)?							
	(a)	API gravity use Table 5A, then (ASTM: if desired) Table 3 to obtain the							
		density at 15.6°C (ASTM: or specific gravity at 60/60°F)?							
	(b)	Specific gravity use Table 23A, then (ASTM: if desired) Table 21 to obtain the							
		density at 15.6°C (ASTM: or specific gravity at 60/60°F)?							
	(c)	Density scaled hydrometer use Table 53A, (ASTM: then if desired use							
		Table 21 to obtain specific gravity 60/60°F or API gravity at 60°F.)?							
22.	Repo	rted at 15.6°C to nearest 1 kg/m ³ (ASTM: or specific gravity 60/60°F or degrees API gravity)?							

COMMENTS (T295/D3142):

(T295/D3142)

COMMENTS (T300):

FORCE DUCTILITY OF BITUMINOUS MATERIALS

/ A	ASHTO T300)	
L A	83010 1300	

Date: _____

<u>A</u>	PP	Άŀ	₹A'	ΤU	JS

MOI	. DC		1	2	2	1	_	(7	0
MOLDS Design conforms to Fig. 1?			1	2	3	4	5	6	7	8
		_								
		- 10.1 mm?								
Widt	h at min.	cross-section: 9.9 - 10.1 mm?						•		1
Bras	s?									
MO	LD PLAT	ES								
Non-	absorbent	?								
Flat	and level?									
							•			
1.	Release				`					
	(a)	As specified in Note 1? (Identify:)	••••••		•••••	• • • • • • • • • • • • • • • • • • • •	•
2.	Water 1	Rath								
2.	(a)	Maker:								
	(b)	Depth of bath not less than 50 mm (2 in.)?)							
	(c)	Bath capable of maintaining temperature v								
	(d)	Mold can be immersed to a depth of at lea								
	(e)	Volume of water not less than 10 L?								
	(f)	Water free of oil and slime or other organi	ic growtl	h?						•
3.	Testino	g Machine								
٥.	(a)	Maker:								
	(b)	Serial No. (or I.D. No.)?		_						
	(c)	Space for at least 25 mm of water above a	nd belov	v sample	at start	of test?				
	(d)	Machine functions without undue vibratio								
	/ /	AAGUTONA		-	r		1.1			
4.	Agents	for Adjusting Specific Gravity of Test Bath	(inform	ational o	nly)	ence	: Lak	oora	ator	V .
	(a)	Specify:			_					
5.	Thermo	ometer								
	(a)	For tests at 4°C (39.2°F) use ASTM There	mometer	· 63C (63	3F)?					
	(b)	For tests performed at other temperatures:								
	(c)	Thermometer calibrated in accordance with								
6.	Heater									
0.	(a)	Oven or hot plate (electric or gas)?								
	(u)	oven of not plate (electric of gas).	•••••••	••••••	••••••	•••••••	•••••••	••••••	• • • • • • • • • • • • • • • • • • • •	•
7.	Miscell	laneous Equipment								
	(a)	Straight edged trimmer at least 1 1/2 in. w								
	(b)	Container?						•••••	•••••	•

(T300)

FORCE DUCTILITY OF BITUMINOUS MATERIALS

(AASHTO T300)				
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8. Calibrated Force Adapter (a) Stainless steel platform and LVDT? (b) Accurate to 0.04 N (0.01 lbs)? (c) Utilizes existing pins on the ductility machine? 9. Digital Indicator (a) Power supply and digital display that serves the LVDT sensor? (b) Interfaces with chart recorder, computer or other readout? 10. Chart Recorder (Optional)? PROCEDURE 1. Sample Preparation (a) Asphalt cement heated preventing local overheating? (b) Sample thoroughly stirred prior to pouring? 2. Mold assembled on plate prepared with release agent? 3. Interior surfaces of mold sides treated with release agent? 4. Mold filled by pouring a thin stream back and forth from end to end? 5. Mold filled until more than level full? 6. Disarrangement of mold parts avoided during filling? 7. Sample, mold, and plate cooled at room temperature? 8. Cooling time 30-40 minutes (at room temperature) 9. Placed in water bath at test temperature for 30 minutes? 10. Excess material cut off with hot straight edged putty knile or spatula? 11. Mold level full? 12. Specimen not pulled away from mold or base plate? 13. Sample, mold and plate and side piaced again in water bath at test temperature? 14. Conditioned in water bath for 85-95 min.? 15. Mold taken off plate and side piaces of mold detached? 16. Briquette leaced in testing machine? 17. Briquette leaced in testing machine? 18. Clips pulled apart at a specific rate? 19. Specimen not pulled away from mold or base plate? 20. Specified minimum ductility met? 21. Briquette tested immediately? 22. Limit of travel of the machine reached? 23. Is gravity adjusted if thread does contact top or bottom? 24. Distance clips pulled measured and recorded? 25. Test not considered normal if the material comes in contact with the surface of the water or the bottom of the bath? 26. Force ductility ratio calculated as the ratio of the force at the second peak (fs) divided by the force at the first peak (fs)?			APPARATUS (Continued)	Date:
(a) Power supply and digital display that serves the LVDT sensor? (b) Interfaces with chart recorder, computer or other readout? PROCEDURE	8.	(a) Stainless steel platform and LV(b) Accurate to 0.04 N (0.01 lbs)?		
PROCEDURE 1. Sample Preparation (a) Asphalt cement heated preventing local overheating? (b) Sample thoroughly stirred prior to pouring? 2. Mold assembled on plate prepared with release agent? 3. Interior surfaces of mold sides treated with release agent? 4. Mold filled by pouring a thin stream back and forth from end to end? 5. Mold filled until more than level full? 6. Disarrangement of mold parts avoided during filling? 7. Sample, mold, and plate cooled at room temperature? 8. Cooling time 30-40 minutes (at room temperature? 9. Placed in water bath at test temperature for 30 minutes? 10. Excess material cut off with hot straight edged putty knife or spatula? 11. Mold level full? 12. Specimen not pulled away from mold or base plate? 13. Sample, mold and plate placed again in water bath at test temperature? 14. Conditioned in water bath for 85-95 min.? 15. Mold taken off plate and side pieces of mold detached? 16. Briquette placed in testing machine? 17. Briquette tested immediately? 18. Clips attached to the pins or hooks of the force adapter and the testing machine? 19. Clips pulled apart at a specific rate? 20. Specified minimum ductility met? 21. Briquette ruptures? 22. Limit of travel of the machine reached? 23. Is gravity adjusted if thread does contact top or bottom? 24. Distance clips pulled measured and recorded? 25. Test not considered normal if the material comes in contact with the surface of the water or the bottom of the bath? 26. Force ductility ratio calculated as the ratio of the force at the second peak (f2) divided by the force at the first peak (f1)? 27. If no second peak is apparent, is the force at minimum specified ductility, point of rupture, or 30 cm used as (f5)?	9.	(a) Power supply and digital displa		
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(a) Asphalt cement heated preventing local overheating? (b) Sample thoroughly stirred prior to pouring? 2. Mold assembled on plate prepared with release agent? 3. Interior surfaces of mold sides treated with release agent? 4. Mold filled by pouring a thin stream back and forth from end to end? 5. Mold filled until more than level full? 6. Disarrangement of mold parts avoided during filling? 7. Sample, mold, and plate cooled at room temperature? (<u>PROCEDURE</u>	
Interior surfaces of mold sides treated with release agent? Mold filled by pouring a thin stream back and forth from end to end? Mold filled until more than level full? Disarrangement of mold parts avoided during filling? Sample, mold, and plate cooled at room temperature? (1.	(a) Asphalt cement heated prevent		
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8. Cooling time 30-40 minutes (at room temp.)? (
9. Placed in water bath at test temperature for 30 minutes? (Sample, mold, and plate cooled at room	temperature? ()	······ <u></u>
10. Excess material cut off with hot straight edged putty knife or spatula?		Cooling time 30-40 minutes (at room te	mp.)? ()	······ <u></u>
11. Mold level full?				
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27. If no second peak is apparent, is the force at minimum specified ductility, point of rupture, or 30 cm used as (f_2) ?	26.	Force ductility ratio calculated as the ratio	tio of the force at the second peak (f_2) divided by	the force
	27.	If no second peak is apparent, is the force	ce at minimum specified ductility, point of ruptu	re,
	28.			

COMMENTS (T300):

(T300)

COMMENTS (T301/D6084):

ELASTIC RECOVERY OF BITUMINOUS MATERIALS BY DUCTILOMETER

<u>APPARATUS</u>

(AASHTO T301)
(ASTM D6084)

Date: _____

Molds		1	2	3	4	5	6
Design	conforms to Fig. 1?						
Width	at midpoint: 10.0 ± 0.1 mm?						
Brass?							
End pi	eces similar to T51/D113?						
Side P	ieces						
Thickn	tess: 10.0 ± 0.1 mm?						
Base P	Plates						
Nonab	sorbent?						
Flat an	d level?						
2.	Release agent as specified in Note 1 (any of the following) (a) Glycerin and dextrin or talc [ASTM: or kaolin] (b) AASHTO: Dow-Corning Silicone Stop-Cock Gree) (c) AASHTO: Castor-oil-Versamid 900 (100:1 mixt) (d) Other release agent that give comparable results Water Bath (a) Maker: (b) AASHTO: Depth of bath not less than 50 mm? (c) Specimens immersed to a depth of at least 25 mm (d) ASTM: Perforated shelf at least 5 cm from both (e) Bath capable of maintaining temperature within (f) Volume of water not less than 10 L? (g) AASHTO: Free from oil, slime, or other organic	AASHTO: ease? ure by weig (Identify: n (ASTM: 1.0°C (1.0°C)	ght, heated 10 cm)? h? F) (ASTM	l and stirre			
3.	Testing Machine (a) Maker: (b) Serial No. (or I.D. No.)? (c) Water tank covers the specimen with at least 25 r (d) ASTM: Water tank maintained within ±0.5 °C (e) (e) Machine capable of maintaining specified speed (f) Machine functions without undue vibration? (g) ASTM: Machine has means of measuring elon	nm of wate 0.9 F) of the within 5 pe gation in c	he test temercent (e.g.	5.00 ± 0.2	? 25 cm/min	1.)?	··
4.	Agents for Adjusting Specific Gravity of Test Bath (inform	nauonai or	ny) speci	ту:			

(T301/D6084)

COMMENTS (T301/D6084):

ELASTIC RECOVERY OF BITUMINOUS MATERIALS BY DUCTILOMETER

(AASHTO T301)	
(ASTM D6084)	

		APPARATUS (Continued) Date:								
1.	Thermo	meter								
1.	(a)	Γhermometer (a) AASHTO: ASTM 17C or 17F for tests performed at 25 ℃ (77 ℉)?								
	(b)	ASTM: 63C or 63F?								
	(c)	Thermometers of appropriate range and equal accuracy to the thermometer required for the test for temperatures other than those covered by the specified thermometers?								
	(d)	Thermometer calibrated in accordance with ASTM E77?								
	(e)	An equivalent thermometric device that has been standardized in accordance with Test Method E220 or Test Methods E644?								
	(f)	ASTM: If 77 F (25 C) penetration bath used, any thermometer or thermometric device with 0.1 C (0.2°F) subdivisions?								
2.	Heater									
	(a)	AASHTO: Oven or hot plate (electric or gas)?								
	(b)	ASTM: Oven capable of maintaining 135 ± 5.5 °C $(275 \pm 10$ °F)?								
3.	Miscella	aneous Equipment								
	(a)	Straight edged trimmer at least 1 1/2 in. wide?								
	(b)	Scissors capable of cutting the bituminous material at test temperature?								
		<u>PROCEDURE</u>								
Sample	Preparati	ion A B A B A B A B A B A B A B A B A B A								
1.		cement heated preventing local overheating (ASTM: in covered container)?								
2.		Heated in an oven at 135 \pm 5°C (275 \pm 10°F)?								
	Note: 1	Higher temperatures may be used if not sufficiently fluid at 135°C.								
3.		O: If polymer modified emulsion residue tested, obtained from evaporation at 162.8 \pm 2.5 °C?								
4.		Sample strained through a 300- μ m (No. 50) sieve preheated at 135 \pm 5°C (275 \pm 10°F)								
	if it is si	uspected that foreign matter is present?								
_		Higher viscosity samples may be strained through an 850-µm (No. 20) sieve.								
5.		If the sample is an asphalt cement or a polymer-modified asphalt cement sample to 135 ±5 °C (275 ±10 °F)?								
6.		If foreign matter is suspected to be in the sample, sample strained through a 300-µm (No. 50)								
0.		eheated at 135 \pm 5°C (275 \pm 10°F)?								
7.		thoroughly stirred prior to pouring?								
8.		ssembled on plate prepared with release agent?								
9.	Interior	surfaces of mold sides treated with release agent?								
10.		lled by pouring a thin stream back and forth from end to end?								
11.		lled until more than level full?								
12.	Disarra	ngement of mold parts avoided during filling?								

(T301/D6084)

ELASTIC RECOVERY OF BITUMINOUS MATERIALS BY DUCTILOMETER

(AASHTO T301)
(ASTM D6084)

	PROCEDURE Date:	
<u>Testing</u>		
1.	Sample, mold and plate cooled at room temperature? ()	
2.	Cooling time 35 ± 5 minutes (at room temp.)? ()	
3.	Placed in water bath at test temperature for 30 to 35 minutes (ASTM: 30 ±5 minutes)	
4.	Excess material cut off with hot straight-edged putty knife or spatula?	
5.	Mold level full?	
6.	Specimen not pulled away from mold or base plate?	
7.	Sample, mold and plate placed again in water bath at test temperature? ()	
8.	Conditioned in water bath for 90 ± 5 minutes? ()	
9.	Mold taken off plate and side pieces of mold detached (ASTM: without distorting or fracturing)?	
10.	Briquette placed in testing machine?	
11.	Water in testing machine within 0.5°C (0.9°F) of test temperature throughout the test?	
12.	Briquette tested immediately?	
13.	AASHTO: Clips pulled to an elongation of 20 cm	
14.	ASTM: 10 ± 0.25 cm for Procedure A and 20 ± 0.25 cm for Procedure B?	
15.	AASHTO: Elongation stopped and specimen held in position for five minutes?	
16.	ASTM: No holding time specified for Procedure A and hold for five minutes for Procedure B?	
17.	Immediately cut into halves at the midpoint?	
18.	Specimen remains in testing machine undisturbed for 60 minutes? ()	
19.	Carriage is moved back until the two ends of the specimen just touch?	
20.	If ends have sagged, are they carefully lifted to their original level prior to adjusting carriage?	
21.	Total length of specimen with severed ends just touching recorded in cm?	
22.	ASTM: Test not considered normal if the sample comes in contact with the surface of the	
	water, the bottom of the bath, or if the sample fractures before reaching elongation?	
23.	Is gravity adjusted if thread contacts bottom or surface and the test rerun?	
24.	Average of three normal tests (nearest whole percent) reported as the elastic recovery?	
25.	If after three tests, a normal test cannot be obtained, is it reported as being unobtainable?	
	AASH 10 Materials Reference Laboratory	
	$Recovery,\% = \frac{Elongation}{Original} - \frac{Elongation}{EndsJustTouching} X100$	
	$Elongation_{Original}$	
	0	
26.	Sample thermal handling history reported?	
COMPA		0.45
COMM	ENTS (T301/D6084): (T301/D608	ŏ4)

DETERMINING THE FLEXURAL CREEP STIFFNESS OF ASPHALT BINDER USING THE BENDING BEAM RHEOMETER (BBR)

(AASHTO T313)	
(ASTM D6648)	

			<u>APPARATUS</u>	Date:
		Manufacturer:	Software:	
1.	Load	ing Frame and Shaft		
	(a)			
	(b)		with the loading shaft?	
	(c)		with load cell and LVDT and able to app	
	(d)	Loading shaft blunt-nosed	with a spherical radius of 6.25 ± 0.30 mm	m [ASTM: 6.30 ±0.30 mm]?
2.	Load	ing System		
	(a)	Capable of applying a cor	tact load of 35 ± 10 mN?	······
	(b)	Rise time from 35 ± 10 m	N to 980 ± 50 mN less than 0.5 seconds?	
	(c)		nds, test load of 980 ± 50 mN applied & l	
	(d)		conds, loads within ± 10 mN from averag	
3.	Load	Cell		
	(a)		0 mN and resolution of 2.5 mN (0.0025 r	nN)?
	(b)		ne controlled temperature bath?	
4.	Linea	ar Variable Differential Trans	ducer (LVDT)	
	(a)		mm and resolution of 2.5µm (0.0025 mm)	
	(b)	Mounted axially with and	above loading shaft?	
5.	Samp	ole Supports		_
	(a)		crosion resistant metal) supports?	
	(b)		s is 102.0 ± 1.0 mm?	and the second s
	(c)	Width of each supporting	area is 9.5 ± 0.25 mm with contact radius	$3 \text{ of } 3.0 \pm 0.3 \text{ mm}$?
	(d)	Vertical alignment pins of	n each support 2 - 4 mm in diameter?	
	(e)	Front face of alignment p	ns 6.75 ± 0.25 mm from the middle of the	e supports?
6.	Temp	perature Transducer [ASTM:		
	(a)		d (check records)?	
	(b)		perature to 0.1°C over the range of -36 to	
	(c)	Mounted within 50 mm o	f the midpoint of the test specimen suppor	rts?
7.		rolled-Temperature Fluid Bat		
	(a)		temperature between -36 and 0°C within	
	(b)		m test temperature allowed due to introdu	1
	(c)		ng bath) does not cause fluctuations great	

Revised 2014-06-02

DETERMINING THE FLEXURAL CREEP STIFFNESS OF ASPHALT BINDER USING THE BENDING BEAM RHEOMETER (BBR)

(AASHTO T313)	
(ASTM D6648)	

		APPARATUS (Continued) Date:				
8.	Data A	quisition System Resolves load to nearest 2.5 mN, beam deflection to nearest 2.5 μm, & temperature. to 0.1°C?				
	(b)	Provides a record of load and deflection measurements at 0.0, 0.5 8.0, 15.0, 30.0, 60.0, 120.0, and 240.0 seconds [ASTM: 8.0, 15.0, 30.0, 60.0, 120.0, and 240.0 seconds]?				
9.	Test S	ecimen Molds				
	(a)	Made from aluminum [ASTM: aluminum, stainless steel, or silicone rubber]?				
	(b)	Interior dimensions when assembled of 6.35 ± 0.05 mm thick, 12.70 ± 0.05 mm wide, and 127 ± 2 mm long [ASTM: 127 ± 5 mm long]?				
	(c)	AASHTO: Two spacers with thicknesses that do not vary from each other by more than 0.05 mm?				
	(d)	ASTM: The thicknesses of the two spacers are measured with a micrometer to verify that they meet the tolerances listed above and the measurements recorded as part of the laboratory's quality control program?				
10.	Sheeti					
	(a)	Clear plastic sheeting, 0.12 ± 0.04 mm [ASTM: 0.08 to 0.15 mm] thick?				
	(b)	ASTM: Silicone coated release paper for metal molds (Optional)?				
		(1) 4.0 to 5.0 mils thick?				
		(2) Coated on both sides?				
	(c)	ASTM: If silicone molds are used, silicone rubber sheeting for lining the glass plate?				
	(d)	Sheeting not distorted from hot asphalt or pulled away from the mold during cooling?				
11.	Releas	Agents				
	(a)	For coating the end pieces of the mold, one of the following.				
	()	(1) AASHTO: A mixture of glycerin and talc or Kaolin (50/50 w/w) is suitable?				
	1 1	(2) ASTM: A mixture of glycerin and Dextrin, talc, or Kaolin?				
		(3) ASTM: Petroleum-based grease?				
		(4) ASTM: A mixture of Versamid and mineral oil?				
		(5) ASTM: Other materials that do not affect the properties of the binder?				
		Note: No silicone based products shall be used.				
	(b)	For adhering the plastic strips to the metal mold.				
		(1) AASHTO: Petroleum-based grease?				
		(2) ASTM: Any of the materials in the section above?				
12.	Bath F	aid				
•	(a)	Not absorbed by or does not affect the properties of the binder?				
	(b)	Optically clear and density not exceeding 1.05 g/cm ³ at test temperature?				
	(c)	Any fluid including ethanol, methanol, or glycol-methanol solution				
	• /	(60% glycol, 15% methanol, 25% H ₂ 0) [ASTM: and stabilized isopropanol]				
		but no silicone fluids or fluids containing silicone?				

DETERMINING THE FLEXURAL CREEP STIFFNESS OF ASPHALT BINDER USING THE BENDING BEAM RHEOMETER (BBR)

(AASHTO T313)
(ASTM D6648)

1.	_	Compliance (thick) Beam				
	(a)	Dimensions of 6.4 ± 0.1 mm thick, 12.7 ± 0.25 mm wide and 127 ± 5 mm long?				
	(b)	ASTM: Dimensions of 6.4 \pm 0.3 mm thick, 12.7 \pm 0.3 mm wide, and 127 \pm 5 mm long?				
	(c)	ASTM: When used to measure thickness of test specimens, compliance beam thickness measured to the nearest 0.01 mm?				
2.	Perfo	rmance (thin) Beam				
	(a)	Dimensions of 1.3 ± 0.3 mm thick, 12.7 ± 0.1 mm wide, and 127 ± 5 mm long?				
	(b)	Manufacturer provides certificates with elastic modulus reported to 3 significant figures, thickness to 0.01 mm and width to 0.05 mm?				
	(c)	Dimensions used to calculate the modulus during overall system check?				
3.	Stand	ard Masses				
	(a)	One or more masses totaling 100.0 ± 0.2 g for verifying load cell?				
	(b)	Two masses of 2.0 ± 0.2 g for verifying load cell?				
		AASHTO note: a coin may be used if the mass is confirmed to be 2.0 ± 0.2 g.				
	(c)	Four equal masses of known mass ± 0.2 g for calibration of load cell?				
	(d)	Two or more masses of known mass \pm 0.2 g for conducting overall system check?				
	(e)	All above masses verified at least once every three years?				
1.	Stepp	ed-thickness gauge block				
	(a)	Of known dimensions and used to verify the calibrations of the LVDT (ASTM: thickness				
		measured to ±5 μm)?				
5.	Thern	nometer [ASTM: Calibrated Thermometric Device]				
	(a)	Calibrated in accordance with E77 at least once per year?				
	(b)	Calibrated in accordance with E77 at least once per year?				
	(c)	Either of the following:				
		(1) A partial immersion liquid-in-glass thermometer with an ice point?				
		(2) An electronic thermometer of equal accuracy and resolution [ASTM: a thermometric				
		device based upon a platinum or thermistor sensor]?				
		Note: A suitable liquid-in-glass thermometer is an ASTM 133C-00. ASTM 62C thermometers are				
		generally standardized as total immersion and should not be used for this test method unless specifically				

DETERMINING THE FLEXURAL CREEP STIFFNESS OF ASPHALT BINDER USING THE BENDING BEAM RHEOMETER (BBR)

(AASHTO T313)	
(ASTM D6648)	

			<u>PROCEDURE</u>	Date:		
1. 2.		orts, loading head, and bath fluid cle brought to selected test temperature				
		d Standardization	11	6 11 .		
1.	air-be verific	acement transducer (LVDT), load co aring, and contact load verified dail cation of zero load cell reading performance. Verifications and calibrations may	y before conducting tests (ormed (step 6)]?	steps 2, 3, 4, 5, 7, and 8) [A	ASTM: and	
	woie.	verifications and canorations may	ve combined at the option	of the manafacturer		
2.	Verifi	cation of Calibration of the Displac	ement Transducer (LVDT			
	(a)	Stepped gage block of known di	mensions used?			
	(b)	100 ± 0.2 g mass applied to the l				
	(c)	Agreement of \pm 5 μ m (0.005 mm measured values and certified di				
	(d)	If values differ by more than ± 5				
	. ,	Note: If tolerance cannot be me				
		manufacturer	•			
		Known Values (mm)	Measured Values (mm)	Within 0.005 mm		
		(from certificate)		[ASTM: 0.015 mm]?		
	1					
		AASHTO	Materials Re	ference Lab	oratory	
3.	Verifi	cation of Air-Bearing	71010110110 110	10101100 2010	or dicory	
•	(a)	Thin beam placed on the support	ts and a 35 \pm 10 mN load a	pplied with the zero load res	gulator?	
	(b)	Reading of the LVDT observed?				
	(c)	Shaft lifted upwards approximate				
		when released? (LVDT reading:)				
	(d)					
		zero load regulator?				
	(e)	An approximately 2 g weight add				
	(f)	Shaft slowly falls downward?				
		If not, too much friction is present				
		/DT adjusted to reduce friction. If i	maintenance does not redu	ce friction, aiscontinue use	ana consult	
	ine me	anufacturer.				
СОМ	MENTS	(T313/D6648):			(T313/D664	

DETERMINING THE FLEXURAL CREEP STIFFNESS OF ASPHALT BINDER USING THE BENDING BEAM RHEOMETER (BBR)

(AASHTO T313)	
(ASTM D6648)	

		PROCEDURE (Continued) Date:
Verif	ication an	nd Standardization (Continued)
4.		ication of Load Cell: Contact Load
т.	(a)	Load shaft rested against compliance beam?
	(b)	20 ± 10 mN load applied to the beam using the zero load regulator?
	(c)	2.0 ± 0.2 g mass added to loading platform?
	(d)	Load display shows an increase of 20 ± 5 mN?
	(e)	A second 2.0 \pm 0.2 g mass added and load display increases by another 20 \pm 5 mN?
	(f)	If not, calibration conducted?
		If tolerance cannot be met after calibration, the lab should discontinue use and consult the manufacturer
5.	Verifi	ication of Load Cell: Test Load
	(a)	Load shaft rested against compliance beam?
	(b)	20 ± 10 mN load applied to the beam using the zero load regulator?
	(c)	100 g mass added to the loading platform?
	(d)	Load display shows an increase of 981 ± 5 mN?
	(e)	If not, calibration conducted?
		If tolerance cannot be met after calibration, the lab should discontinue use & consult manufacturer
6.	Verifi	ication of Zero Load Cell Reading
	(a)	Thick steel beam placed on the loading platform?
	(b)	A 100 g mass placed on the platform?
	(c)	Reading of the LVDT observed? (LVDT reading) This is the test starting position
	(d)	Thick steel beam and 100 g mass removed?
	(e)	Loading shaft adjusted to the test starting position while free floating?
	(f)	The load indicated is 0 ± 5 mN?
	Note:	If the tolerance cannot be met after calibration, the lab should discontinue use and consult the
	manu	facturer. AASHTO Materials Reference Laboratory
7.	Daily	Overall Systems Performance Check
	(a)	Thin beam placed on the sample supports and manufacturer's instructions followed?
	(b)	$50.0 \text{ or } 100.0 \pm 0.2 \text{ g mass applied to the beam?}$
	(c)	An additional 100 to 300 ± 0.2 g applied (depending on the manufacturer's instructions)?
	(d)	Calculated modulus within 10% of the modulus reported by the manufacturer of the BBR?
	(e)	Reported value from Manufacturer's certificate:
	(f)	
	(f)	Reported valued from Performance Check:

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DETERMINING THE FLEXURAL CREEP STIFFNESS OF ASPHALT BINDER USING THE BENDING BEAM RHEOMETER (BBR)

(AASHTO T313)	
(ASTM D6648)	

	PROCEDURE (Continued) Date:
cation on	d Standardization (Continued)
	cation of Calibration of the Temperature Detector
	Temperature verified each day and whenever the test temperature is changed?
` /	Thermometer immersed in bath close to the thermal detector?
	Calibrated thermometer reading agrees with the thermal detector within $\pm 0.1^{\circ}$ C?
	If not, is calibration conducted?
(u)	If not, is canoration conducted?
	cation of front-to-back alignment of the loading shaft (Assessors: Check Records)
	AASHTO: Alignment checked every 6 months?
	ASTM: Alignment checked when installed, disturbed, or the alignment is thought to be suspect?
(c)	Checked by one of the following means:
	(1) Checked with an alignment gauge supplied by the manufacturer?
	(2) Checked by measurement as follows?
	1. 25-mm long strip of white paper slightly narrower than the beam taped
	to the thick beam with scotch tape?
	2. With the frame out of bath, beam placed on the supports with a small piece
	of carbon paper placed atop white paper near beam midpoint?
	3. With the air pressure applied to air bearing, shaft pushed downward to
	make an imprint on the paper?
	4. Distance between the center of the imprint and the two edges of the beam
	measured with vernier calipers?
	5. Difference between measurements 1.0 mm or less?
	Note: If the alignment does not meet the specification, the manufacturer should be consulted.
ration of	Molds (either of the following)
	molds (AASHTO and ASTM)
(a)	ASTM: Visually inspected to verify molds are free of nicks, dings, and burrs?
(b)	Very thin layer of petroleum-based grease used to hold plastic strips to the aluminum?
(c)	Molds assembled correctly using O-rings?
(d)	Air bubbles eliminated between plastic film and aluminum [ASTM: with firm finger pressure]?
(e)	Inside faces of the two end pieces covered with glycerol-talc mixture?
(f)	Mold assembly kept at room temperature until asphalt is poured?
Silico	ne molds (ASTM only)
	Any remaining binder, grease, or other residue wiped out of the molds with a clean, dry cloth?
(u)	Note: A cloth moistened with an essentially residue free solvent (e.g. acetone or heptane)
	is satisfactory for this purpose. If a solvent is used to wipe down the molds, allow the molds
	to dry at ambient temperature for at least 10 minutes prior to use.
<i>a</i> >	Assembled correctly according to Figure 5?
(b)	
	Verifi (a) (b) (c) (d) Verifi (a) (b) (c) Metal (a) (b) (c) (d) (e) (f)

DETERMINING THE FLEXURAL CREEP STIFFNESS OF ASPHALT BINDER USING THE BENDING BEAM RHEOMETER (BBR)

(AASHTO T313)	
(ASTM D6648)	

	PROCEDURE (Continued) Date:			
Prena	ration of Material			
1.	Asphalt binder heated in an oven set at up to 165°C [ASTM: 168±5°C] until sufficiently fluid to pour?			
2.	AASHTO: When testing modified binders, the oven may be set at up to 180% ?			
3.	ASTM: May be heated to higher temperatures if material does not pour easily when heated at 173°C.			
	Oven temperature noted on report?			
4.	Heating time minimized?			
	g of Molds			
1.	Metal molds (AASHTO and ASTM)			
	(a) Sample container held 20 to 100 mm from the top of mold?			
	(b) Mold filled by pouring stream of asphalt from one end toward other in a			
	single pass, slightly overfilling?			
2.	Silicone molds (ASTM only)			
	(a) If the viscosity of the binder warrants, silicone mold preheated in a 135°C for up to 30 minutes?			
	(b) Mold filled from the top in a slow steady manner?			
	(c) Entrapment of air bubbles avoided?			
	(d) Mold filled to the top with no appreciable overfilling?			
Prepa	ration and Conditioning of Specimens			
1.	Samples cooled at ambient temperature for 45 to 60 minutes?			
2.	Excess material cut off with a hot knife or a heated spatula?			
	ASTM note: "Buttering" maybe be performed on the samples from stiffer grades of binder by bringing the			
	heated trimming tool into momentary contact with the top of the specimen just enough to flatten the surface.			
3.	AASHTO: Specimens cooled in a freezer or ice bath at -5 \pm 7 °C for 5 – 10 minutes?			
4.	ASTM: Specimens cooled for no longer than 5 minutes to stiffen the specimen sufficiently to			
	be readily demolded?			
5.	ASTM: Not exposed to temperatures within 10°C of the test temperature?			
6.	Specimens immediately demolded after cooling by disassembling the metal mold [ASTM: or by			
	removing the specimen from the silicone mold]?			
7.	Distortion and warping of the specimens avoided?			
	Note: If the plastic sheeting does not fully separate from the beam, the final portion of the sheeting			
	may be removed while immersing the sample in the bath to avoid distortion.			
8.	Specimens immediately submerged in the bath at test temperature?			
9.	Conditioned for 60 ± 5 minutes?			
10.	ASTM: Test specimen thickness taken as 6.35 mm or measured by one of the methods in Section 13.2?			

DETERMINING THE FLEXURAL CREEP STIFFNESS OF ASPHALT BINDER USING THE BENDING BEAM RHEOMETER (BBR)

(AASHTO T313)	
(ASTM D6648)	

		PROCEDURE (Continued) Date:
Factio	ng of Spe	cimens
1 esui 1		ections and test loads checked using the compliance beam?
	(a)	Using the test load regulator, 980 ± 50 mN added to the thick beam?
	(b)	Switched to the zero load regulator, and 35 ± 10 mN contact load added to the thick beam?
	(c)	Switched between the test and the contact load at least 4 times [ASTM: until consistent readings are
	(0)	obtained] to ensure that the loads are maintained?
	(d)	When switching between contact and test loads, loading shaft remains in contact with steel beam
		(Check by observing LVDT readings as loads are switched)?
	(e)	ASTM: Successive contact load readings vary by no more than 10 mN?
		Note: If the loads do not recover, the device may require cleaning or calibration of the load cell,
		and if it still does not work, the lab should consult the manufacturer.
2.		men information entered into the test system?
3.	-	men placed on supports against the alignment pins? [ASTM: if the thickness is not determined
	•	placement]
1.		fluid maintained at test temperature \pm 0.1° C during testing?
	(a)	If not, is the test rejected?
5.		act load of 35 \pm 10 mN applied to beam?
	(a)	Contact load applied by gently increasing and does not exceed 45 mN?
_	(b)	Time to apply and adjust the contact load no greater than 10 s?
5.		matic test system activated and proceeds as follows:
	(a)	980 ± 50 mN initial seating load applied for 1.0 \pm 0.1 s?
	(b)	Load reduced to 35 \pm 10 mN and beam allowed to recover for 20.0 \pm 0.1 s?
		(1) If not, is test rejected? Test load of 980 ± 50 mN applied?
	(c)	
	(d)	Maintained ± 50 mN of the average test load for the first 5 s?
	(e)	Maintained \pm 10 mN from the average of the test load from 5.0 s to 240.0 s for the remainder of the test?
	(6)	the remainder of the test?
	(f)	Test load removed and test terminated?
	(g)	Load on the beam returns to 35 ± 10 mN at the end of the test?
	(1.)	Note: If not, the test is invalid and the rheometer should be recalibrated.
7	(h)	Time required for rise from 35 ± 10 mN preload to 980 ± 50 mN test load less than 0.5 seconds?
<i>/</i> .		of measured load versus measured deflection generated [ASTM: intervals of 0.5 s or less] ng with the seating load?
	startii	ig with the scatting 10au :

DETERMINING THE FLEXURAL CREEP STIFFNESS OF ASPHALT BINDER USING THE BENDING BEAM RHEOMETER (BBR)

(AASHTO T313)	_
(ASTM D6648)	_

ASSHTO 1. Date and time when test initiated & load applied? 2. File name of test data, name of operator, & specimen identification number? 3. Conditioning time for specimens
1. Date and time when test initiated & load applied? 2. File name of test data, name of operator, & specimen identification number? 3. Conditioning time for specimens. 4. Any flags issued by software during test. 5. Correlation coefficient, R² for log stiffness versus log time expressed to nearest 1 x 10⁻ 6. Constants A, B, & C to three significant figures. 7. Percent difference between measured & estimated stiffness modulus. 8. Load and deflection for times 0.0 and 0.5 seconds (to nearest 1 mN)? 9. Report data for time intervals of 8.0, 15.0, 30.0, 60.0, 120.0, and 240.0 seconds including: (a) Loading time (nearest 0.1 sec.)? (b) Load (nearest 1.0 mN)? (c) Beam deflection (nearest 1 μm)? (d) Measured stiffness modulus (MPa to three significant figures)? (e) Estimated stiffness modulus (MPa to three significant figures)? (f) Estimated m value (nearest 0.001)? 10. Report load and deflection for times 0.0 and 0.5 seconds (to nearest 1 mN)? ASTM 1. Test Specimen Information (a) BBR file name and specimen ID number? (b) Test specimen width and thickness (defaults are 12.70 and 6.35 mm, respectively)? (c) Date of test? (d) Version of software used? (e) Calibration information (f) Date of the last temperature, load cell, and LVDT calibrations? (g) Load cell and LVDT calibration constants (mN/bit and μm/bit, respectively, to three significant figures)? (h) Date of the last modulus and compliance checks? (i) Measured modulus and of the steel beam and compliance of the loading system (GPa and μm/N)
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(f) Estimated m value (nearest 0.001)?
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 (c) Date of test?
 (d) Version of software used?
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(i) Measured modulus and of the steel beam and compliance of the loading system (GPa and μm/N,
respectively to three significant figures)?
(a) Time test load applied
(c) Minimum and maximum load recorded during test (to the nearest 1 mN)?
(d) Contact load at t = 0, just prior to application of the test load (to the nearest 1 mN)?
(e) Test load after 0.5 s loading time (to the nearest 1 mN)?
3. Test Results (report the following test results for time intervals of 8.0, 15.0, 30.0, 60.0, 120.0, 240.0 s)
(a) Loading time in seconds (nearest 0.1 s)?
(b) Test load (mN to nearest 1 mN)?
(c) Test specimen deflection (mm to nearest 1 µm)?
(d) Measured stiffness modulus, Eq. 3 (MPa to three significant figures)?
(e) Estimate stiffness modulus, Eq. 5 (MPa to three significant figures)?
(f) Percent difference between estimated and measured stiffness?
(g) Estimated m-value, Eq. 6 (to nearest 0.001)?

COMMENTS (T313/D6648):

DETERMINING THE FLEXURAL CREEP STIFFNESS OF ASPHALT BINDER USING THE BENDING BEAM RHEOMETER (BBR)

(AASHTO T313)	_
(ASTM D6648)	_

		DENDING BEAM RHEOMETER (BBR)	
		REPORT (Continued)	Date:
4.	Sumn	nary Data	
	(a)	Regression coefficients and R ²	
	(b)	Average load at 0.5 s and every 0.5 s thereafter up to 240.0 s	
	(c)	Maximum deviation of the load from 0.5 to 5.0 s (mN)?	
	(d)	Maximum deviation of load from 5.0 to 240.0 s (mN)?	
	(e)	Deflection at zero time (mm)?	
	(f)	Deflection at 0.5 s (mm)?	
COM	MENTS	(T313/D6648):	(T313/D6648)



DETERMINING THE FRACTURE PROPERTIES OF ASPHALT BINDER IN DIRECT TENSION (DT)

(AASHTO T314)
(ASTM D6723)

Date: _____

APPARATUS

(a) Distance between the loading pins capable of accommodating specimens with a total length of at least 100 mm (including the plastic end tabs)?	
 (b) If a fluid based system is used, gripping system (loading pins and platens) completely submerged under the cooling fluid at a minimum depth of 25 mm (1 in.)? (c) If an air based system is used, testing frame equipped with two columns having sufficient space between them so that an insulated temperature control chamber can be placed between 	
under the cooling fluid at a minimum depth of 25 mm (1 in.)?	
(c) If an air based system is used, testing frame equipped with two columns having sufficient space between them so that an insulated temperature control chamber can be placed between	
space between them so that an insulated temperature control chamber can be placed between	
the two columns?	
The following lines are for informational purposes only:	
(d) Fluid bath or an insulated environmental chamber?	
(e) Real-time load measuring and recording devices?	
(f) Real-time elongation measuring and recording devices?	
(g) Real-time temperature detection and recording devices?	
(h) Real-time data acquisition and display devices?	
(i) Electro-mechanical or servo-hydraulic loading unit capable of applying and measuring:	
(1) Tension and compression forces of at least 500 N?	
(2) Actuator travel of 20 mm (0.78 in.)?	
(j) Closed feed-back loop displacement-controlled tensile loading machine?	
(k) System stiffness at least 3 MN/m, including the load cell and loading pins?	
(l) Displacement transducer:	
(1) Capable of measuring and controlling grip separation?	
(2) Provides feedback for strain control with a displacement resolution of 1.0 μm?	
(m) System capable of closed loop elongation control accurate to at least 1 percent of the	
commanded specimen elongation rate using either:	
(1) Feedback from a displacement transducer mounted between the loading pins?	
(2) A non-contact extensometer measuring elongation of the specimen?	
(n) Closed loop elongation rate control algorithm capable of real time compliance correction?	
(o) Tensile loading machine having a controlled-displacement loading frame capable of producing at least a 500 N load?	ry
(p) Loading frame table mounted?	
(q) Load cell having a minimum capacity of 500 N and a sensitivity of 0.1 N?	
2. Specimen Gripping System	
(a) Produces a self-aligning uniaxial test load?	••••
(b) Accepts the specified plastic end tabs of the specimen?	
(c) Consists of two grips?	
(d) Grips include a specially-shaped pin that is rigidly mounted to the loading platens of	
the testing machine?	
(e) One grip remains stationary while the other is displaced at the desired elongation rate?	

COMMENTS (T314/D6723):

DETERMINING THE FRACTURE PROPERTIES OF ASPHALT BINDER IN DIRECT TENSION (DT)

(AASHTO T314)	
(ASTM D6723)	

APPARATUS (Continued)

			<u>APPARAT</u>	TUS (Continued)	Date:							
3.	Temp	erature C	ontrol Chamber									
	(a)	Chamber having sufficient space for storage of at least 12 specimens (ASTM: at least 8 specimens) on a rack?										
	(b)	-	erature control range for the cooling		-							
					desired test temperature?							
	(c)		num temperature fluctuation from the	-	• • •							
					······							
	(d)		perature gradient between grips does not exceed ± 0.1°C?									
	(e)											
		(1)	Located in the bath in the proxim	ity of the test area?								
4.	Coolii	ng Syster	n (either of the following)									
	(a)	Air B	ased									
		(1)	System controlled mechanically	or by liquid nitrogen?	<u></u>							
		(2)	Equipped with a dehumidifying s of the chamber and the test speci		ation of frost on the interior							
		(3)	Equipped with a front-opening de	oor for maintenance and	standardization purposes and							
			an access port allowing for the in position specimens?		hand and forearm to							
		(4)	Chamber and access port designed									
		(5)			removed from the chamber?							
		(5)	Visual access to the interior of the specimens and for monitoring the		proper mounting of the test							
		(6)	Specimen elongation measured v	vith an optical laser, optic	cal glass windows on two sides							
			of the temperature chamber so th	at a laser beam can be pa	assed through the chamber							
	1 .											
	()	(7)	Plexiglas, Teflon, or other plastic									
			environmental chamber?	mals Reterd	ence Laboratory							
		(8)	Freezer capable of maintaining a	temperature of -15 ± 5 °C	C?							
	(b)	Fluid	Based									
		(1)										
		(2)	Aqueous mixture of 42% potassi	um acetate and 58% deio	onized water by weight?							
COM	MENTS ((T314/D	5723):		(T314/D6723)							

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DETERMINING THE FRACTURE PROPERTIES OF

(AASHTO	T314)
(ASTM D	6723)

		ASPHALT BINDI	ER II	N DI	REC	T TI	ENSI	ION	(DT)			(A	STM	1 D67
		APPA	RAT	US (Cont	inuec	<u>l)</u>				D	ate:		
5.	Specir (a)	men Molds Made of aluminum?												
		Dimensions	1	2	3	4	5	6	7	8	9	10	11	12
		Length: 99.924 to 99.975 mm?												
		Width at point of contact: 26.880 to 27.120 mm?												
		Width from point of contact to shoulder: 9.880 to 10.120 mm?												
		Width from face to face: 5.760 to 6.240 mm?												
		Thickness of mold: 5.880 to 6.120 mm?												
		Inside length of base:												

(b)	Release agent	(mixture of 20	g of glycering	and 20 g of	talc (USP)?	 	
/ \	D 1			TT C1 (/	COTTO # 171	 •	T. O.

- Paper coated on both sides with 0.3 microns Teflon (ASTM: silicone coated papers may be used)? ... __ (c)
- (d) Bottom plate?....

6. Acrylic End Tabs

100.025 – 100.127 mm?

End tabs machined from Phenolic G-10 material?....

Dimensions	1	2	3	4	5	6	7	8	9	10	11	12
Length: 29.4 to 30.6 mm?	ате	erie	ais	K	ете	ere	ene	ce	La	DO	ога	το
Width: 19.4 to 20.6 mm?												
Height: 5.4 to 6.6 mm?												
Hole lined with stainless ring?												
Diameter of ring: 9.95 to 10.05 mm?												

COMMENTS (T314/D6723):

DETERMINING THE FRACTURE PROPERTIES OF ASPHALT BINDER IN DIRECT TENSION (DT)

(AASHTO T314)
(ASTM D6723)

		APPARATUS (Continued) Date:
7.	Data A	Acquisition Systems
	(a)	Load and elongation monitored with a data acquisition system such that they can be resolved to 1 percent of the failure load and elongation, respectively?
	(b)	Once the test has started, data acquisition system detects the point in time when the load starts to change as a result of elongation in the sample?
	(c)	Data acquisition and display system displays a stress-strain curve in units of stress (MPa) versus percent strain?
	(d)	If an x-y recorder is used, the units may be in volts but the test file shall contain the calibration factor in MPa/Volt and percent strain/volt for both x and y axes?
	(e)	Display device displays the strain failure once the test is complete,
	(f)	Stress and strain displayed to the nearest 0.1 percent?
	(g)	If the data acquisition component consists of an IBM-compatible computer: (1) Does it have three A/D channels: one for load, one for elongation, and one
		for temperature?(2) Data stored in ASCII format?
8.	Misce	llaneous
	(a)	Calibrated liquid-in-glass thermometer of suitable range with subdivisions of 0.1°C for verification of the temperature transducer?

[AMRL Policy: AAP will accept 0.04°C for liquid-in-glass thermometers, please write the finding as informational if the uncertainty is less than or equal to 0.04°C.]

Note: The thermometric device used for <u>calibration</u> of the temperature transducer (at least once per

(b) Forced-air convection oven capable of maintaining $160 \pm 5^{\circ}$ C for heating the asphalt?.....

(d) Cotton cleaning cloths for wiping molds, end tabs, and plates?....

COMMENTS (T314/D6723):

DETERMINING THE FRACTURE PROPERTIES OF ASPHALT BINDER IN DIRECT TENSION (DT)

(AASHTO T314)	_
(ASTM D6723)	_

		CALIBRATION AND VERIFICATION Date:
	Load	Cell and Displacement Transducer
	(a)	Verified at least every 6 months and whenever measurements are suspect (check records
		of verification for agreement with certificate values)?
	(b)	Verification standard having a spring rate of approximately 135 N/m used?
	(c)	Supplied by manufacturer along with stress-strain characteristics?
	(d)	Equipment on which the force and elongation for the spring was determined NIST traceable?
		Note: this usually relates to the initial calibration of the DTT at the factory, with a subsequent determination
		of the spring properties on that DTT. There is no requirement for the spring itself to be NIST-traceable.
	Verifi	cation of the Elongation Rate
	(a)	Performed in conjunction with the load cell and displacement transducer verification?
	(b)	Measurements made at -18°C?
	(c)	Elongation plotted as a function of elapsed time?
	(d)	Resulting plot a straight line with a slope of 1.00 mm/min.?
	X 7 . C'	
		cation of the Temperature Detector
	(a)	Performed at least once per year [ASTM: at least every six months]?
	(b)	Output from the RTD compared to a calibrated liquid-in-glass thermometer [ASTM: a
	(.)	NIST-traceable RTD digital thermometer may be used]?
	(c)	Calibrated at each temperature used?
	(d)	Direct contact between RTD and temperature calibration device?
	(e)	If not agreeing within \pm 0.1°C, correction applied or further calibration or maintenance performed?
		/ / AIVIE/I
		PROCEDURE
	١.	
•	_	ration of Sample
	(a)	If binder is unaged, sample obtained in accordance with T40 [ASTM: D140]?
	(b)	AASHTO: Sample degassed prior to testing as described in R28? [ASTM: this is still in the ASTM
		PAV specification and should be performed for ASTM testing as well]
	(c)	Sample heated until sufficiently fluid to pour?
	(d)	Specimen forming surfaces of the two mold plates coated with the release agent such that no part
		of the metal surface is exposed?
	(e)	Single precut sheet of release paper placed on the bottom plate of the mold?
	(f)	Side plates (coated with release agent) and plastic end tabs assembled on the bottom of the plate?
	(g)	AASHTO: Assembly placed into an oven for no more than 3 minutes?
	(h)	ASTM: Mold assembly placed in sand bath (preheated in the oven) in oven for 7 ± 2 min?
	(i)	Oven set at the same temperature the sample was heated to for pouring?
	(j)	Hot asphalt binder poured into the mold starting from one end of the cavity and moving across the cavity in a single pass?
	(k)	Specimen poured in a continuous stream to avoid entraining air bubbles or gaps?
	(1)	Sample poured until asphalt is slightly above the top surface of the mold?
	(m)	AASHTO: Pouring operation completed as quickly as possible to avoid excessive drop in
	()	temperature of the asphalt?
	(n)	ASTM: Only two molds heated and poured at a time
	(o)	ASTM: Molds allowed to anneal in sand bath for 5 ± 1 min before being placed on the bench top?

Entire assembly allowed to cool on a bench top at ambient temperature for 30 to 60 minutes?.....

Specimen not quenched to achieve an ambient temperature?....

COMMENTS (T314/D6723):

(1)

(p)

DETERMINING THE FRACTURE PROPERTIES OF ASPHALT BINDER IN DIRECT TENSION (DT)

(AASHTO T314)
(ASTM D6723)

PROCEDURE (Continued)

		PROCEDURE (Continued) Date:
2.	Trimn	ning of the Specimen
	(a)	Excess asphalt binder trimmed off with a straightedge heated to approximately 165°C so that
	()	the asphalt binder is flush with the top of the mold?
	(b)	Care taken during the trimming operation so that:
	. /	(1) The asphalt binder is not pulled away from the mold?
		(2) The bond between the plastic end tabs and the asphalt binder is not damaged?
	(c)	Debris and extraneous asphalt binder removed from holes or slots in the plastic end tabs?
	(d)	Specimen allowed to sit at ambient temperature for 10 to 15 minutes?
3.	Demo	lding and Conditioning of the Specimen
	(a)	Two side plates of an unused aluminum mold placed in cooling bath or air chamber and cooled
		to test temperature for use as holders when transferring specimens?
	(b)	Bottom plate of an unused aluminum mold placed upside down on the work bench and two
		release papers placed so that they overlap lengthwise and cover the plate?
	(c)	Specimen and the two side plates gently slid towards one edge of the bottom plate until the
		side plate nearest to the edge is halfway across the edge?
	(d)	Overhanging side plate pivoted downwards and separated using gentle pressure?
	(e)	Just released side plate replaced?
	(f)	Specimen slid, with both sides in place, to the other edge of the bottom plate?
	(g)	Demolding procedure repeated except one plate removed completely and one cold side plate
	(1-)	from bath or chamber mounted on that side?
	(h)	Other side plate demolded and other cold side plate from bath or chamber mounted?
	(i)	Cold specimen mold assembly turned upside down and placed on the transfer plate?
	(j)	Bottom plate (now on top) removed from upside down specimen assembly by
	(k)	sliding plate off assembly?
	(1)	Release paper gently removed while holding the two plates?
	(n)	
	(III)	Specimen, along with the two cold side plates, immediately placed into the cooling bath or chamber onto one of the trays?
	(n)	Specimen side plates removed after a two minute cool-down period?
	(o)	Specimen flipped in test bath or chamber so that the trimmed side is up?
	(p)	Transfer plate with release paper never placed in bath?
	(q)	Cooling bath stabilized to within $\pm 0.1^{\circ}$ C of the test temperature?
	(r)	Specimen conditioned at the test temperature for 60 ± 10 [ASTM: 60 ± 5] minutes?

COMMENTS (T314/D6723):

DETERMINING THE FRACTURE PROPERTIES OF ASPHALT BINDER IN DIRECT TENSION (DT)

(AASHTO T314)
(ASTM D6723)

				PROCEDURE (Continued)	Date:			
4.	Test F	Procedure f	or Fluid Ba	ath Cooling Systems				
	(a)							
	(b)	Seating of the specimen checked by running the tongs lightly along the end tabs?						
	(c)			ng software controlled or manual set up procedures				
	(-)	(1)		automatically removes the slack between the speci				
		· /		e strain, and starts the test when load reaches a value				
		(2)	Manual se					
		· /		Thumb wheel moved until 2 ± 0.3 N is shown on the	he monitor?			
				Seating of the specimen checked with the tongs on				
				erroneous readings?				
				Slack again removed until a load of 2 ± 0.3 N obta				
				Strain quickly zeroed and test started?				
		(3)		te set to 3 percent per minute (an elongation rate of				
5.	Test F	Procedure f	or Forced A	Air Cooling Systems				
	(a)	Specimen mounted on the pins using the environmental chamber hand access port after conditioning?						
	(b)	Specimen mounted using rubber surgeon's gloves, taking care not to touch the asphalt binder						
				e chamber door?				
	(c)	If a was		I to secure the end tabs to the pins:				
		(1)						
		(2)		nm thick with an outer diameter of approximately 1				
		(3)		onto the pins?				
	(d)			the specimen which results in an elongation suffic				
	(e)			d and the load relaxed until it is no longer detachab				
	(f)	Desired	l strain rate	selected and specimen loaded to failure?	······			
_	1	/						
6.	If Testing for Compliance to AASHTO M320 / ASTM D6373							
	(a)	Strain r	ate selected	d that gives an effective gage section elongation ra	te of 1.00 ± 0.01 mm/min?			
	(b)	If the te	est specime	n fails outside of the gage area of the specimen, tes	st discarded?			
7	Esilee	I.J4:C	.4:					
7.		e Identifica		studie at failure recorded as the studie at most load	(marinum atmass)?			
	(a)			strain at failure recorded as the strain at peak load ses not fracture, strain at failure recorded as the strain				
	(b)			ss?ss?	1 0			
	(c)			er can be stretched to 10 percent without fracture,				
	(0)			"greater than 10 percent"?				
	(d)			ils in the throat section, noted that failure occurred				
	(u)	11 1110 5	occinicii tai	ns in the timbat section, noted that failule occurred	m me moat:			

If end tabs are to be reused, soaked in solvent or sprayed with a degreasing cleaner, and wiped with a soft cloth?....

COMMENTS (T314/D6723):

Cleaning (a)

(b)

(c)

8.

COMMENTS (T314/D6723):

(T314/D6723)

DETERMINING THE FRACTURE PROPERTIES OF ASPHALT BINDER IN DIRECT TENSION (DT)

(AASHTO T314)
(ASTM D6723)

CALCULATIONS AND REPORT	Date:

Calcula	tions
	·
(a)	Is the failure stress computed by the following equation?
	$\sigma_f = P_f / A$ where $A = 36 \times 10^{-6} \text{m}^2$
(b)	Is the failure strain computed by the following equation?
	$\varepsilon_{\rm f} = \sigma_{\rm f} / L_{\rm e}$ where $L_{\rm e} = 33.8 \ \rm mm$
(c)	Total of six specimens tested?
(d)	Two specimens with lowest values of failure stress, strain, and energy discarded?
(e)	Mean and standard deviation values for the four remaining failure values calculated?
	·
Report	
-	The test temperature to the nearest 0.1°C?
` '	The rate of elongation to the nearest 0.01 mm/min.?
` '	The failure strain to the nearest 0.01 percent [ASTM: average failure strain and
(0)	standard deviation]?
(1)	
(d)	The failure stress to the nearest 0.01 Mpa [ASTM: average failure stress and standard deviation]?
(e)	The peak load to the nearest N?
	(c) (d) (e) Report (a) (b) (c) (d)



COMMENTS (T315/D7175):

DETERMINING THE RHEOLOGICAL PROPERTIES OF ASPHALT BINDER USING A DYNAMIC SHEAR RHEOMETER (DSR)

(AASHTO T315)
(ASTM D7175)

		<u>APPARATUS</u>	Date:			
Manuf	facturer: _	Software:	Model:			
1.	Metal 7	Test Plates				
	(a)	One set of plates 8.00 ± 0.02 mm in diameter?	······			
	(b)	One set of plates 25.00 ± 0.05 mm in diameter?				
	(c)	Base plates are either flat or have a raised portion a minimum of 1 radius as the upper plate?				
	(d)	Plates are concentric with each other when mounted in the DSR?				
2.	Enviro	nmental Chamber				
	(a)	If air is used as a media, is a suitable drier included to prevent conformation of ice on the plates?	······			
	(b)	Controls the temperature of the specimen to an accuracy of 0.1°C				
	(c)	Chamber completely encloses top and bottom plates?				
	(d)	If a liquid circulating bath is used, is the flow rate of the bath med temperature settings have been adjusted to the desired value?				
	(e)	Are media lines periodically inspected, cleaned, and replaced if no	ecessary to remove obstructions?			
3.	Temperature Controller					
	(a)	Capable of maintaining specimen temperatures within ± 0.1°C for from 3 to 88°C [ASTM: 4 to 88°C]?	test temperatures ranging			
4.	Interna	1 Temperature Measurement Device [AASHTO: Platinum Resistand	ce Thermometerl			
	(a)	Mounted within the environmental chamber as an integral part of to the fixed plate?	the DSR and in closer proximity			
	(b)	Range of 3 to 88°C [ASTM: 4 to 88°]?				
	(c) (d)	Resolution of 0.1°C? Provides a continuous readout of temperature during the mounting	ence Laboratory			
		specimen?	······			
		Note: Platinum resistance detectors meeting DIN Standard 43760 The PRT shall be calibrated as an integral unit with its respective				
5.	Loadin	g Device				
	(a)	Applies a sinusoidal oscillatory load to the specimen at a frequence (1) If other frequency, accurate to 1 percent?				
	(b)					
	(c)	AASHTO: If strain controlled, torque accurate to 100 µrad or if to 10 mN•m?	stress controlled, torque accurate			
	(d)	AASHTO: Total system compliance at 100 Nom torque less than 2				
	(e)	Manufacturer provides a certificate certifying that the frequency, s measured with an accuracy of 1% or less in the range of this measured.				

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(AASHTO T315)
(ASTM D7175)

		APPARATUS (Continued) Date:					
6.	AASHTO Only: Control and Data Acquisition System						
	(a) Provides a record of temperature to 0.1 °C?						
	(b)	Provides a record of frequency to 1 percent?					
	(c)	Provides a record of torque to 10 mN•m?					
	(d)	Provides a record of deflection angle to 100 μrad?					
	(e)	Measures and records G*, in the range of 100 Pa to 10 MPa, to an accuracy of 1.0 % or less?					
	(f)	Measures and records phase angle, in the range of 0 to 90°, to an accuracy of 0.1°?					
7.	Silicor	ne Rubber Molds (Optional)					
	(a)	Thickness greater than 5 mm?					
	(b)	AASHTO: Diameter of cavity approximately equal to diameter of upper test plate?					
	(c)	AASHTO: For a 25-mm test plate with a 1-mm gap: mold cavity approximately 18 mm in					
	diameter and 2.0 mm deep?						
	(d)	AASHTO: For an 8-mm test plate with a 2-mm gap: mold cavity approximately 8 mm in					
	` '	diameter and 2.5 mm deep?					
		AMRL: The mold must be large enough for excess material.					
8.	Specin	nen Trimmer					
	(a)	Straight edged?					
	(b)	AASHTO: at least 4 mm wide?					
9.		g Material					
	(a)	Clean cloth, paper towels, cotton swabs, or other suitable material for wiping test plates?					
10.	Cleaning Solvents						
	(a)	Mineral oil, citrus-based solvents, mineral spirits, toluene, or similar solvent for cleaning the plates?					
	(b)	AASHTO: Acetone for removing solvent residue from the surfaces of the plates?					
	(c)	ASTM: An organic solvent that does not leave a residue, such as heptane, acetone, or ethyl alcohol may be used for removing solvent residue from the test plates?					
COMM	ENTS (T315/D7175): (T315/D7175)					

Revised 2014-06-02

(AASHTO T315)
(ASTM D7175)

			APPARATUS (Continued)	Date:			
11	Deferer	aa Tharmamatar					
11.	Reference Thermometer (a) Either a NIST-traceable liquid-in-glass or a NIST-traceable electronic thermometer used to verify the						
	(b)	portable thermometer?					
	(b) (c)	Liquid-in-glass thermometer					
	(C)	(1) Partial imm	nersion NIST-traceable with a suitable range [AS and subdivisions of 0.1°C				
			e point and calibrated in accordance with ASTM				
		(3) Optical vie	ewing device for use with the thermometer that en parallax (optional)?	nhances readability and			
		(4) ASTM: V	erified at least once a year?				
	(d)	Electronic Thermon	meter [ASTM: Digital Temperature Measureme	ent Device]			
		(1) With an ac	ccuracy of ± 0.05 °C and a resolution of 0.01 °C?.				
			at least annually using a NIST-traceable referenced test method E77?				
			tinum resistance detectors meeting DIN Standard ded. The PRT shall be calibrated as an integral circuitry.				
12.	Portabl	e Thermometer [AST	M: Portable Temperature Measurement Devic	e l			
12.	(a)						
	(-1)), , , , , , , , , , , , , , , , ,				
	(b)		2.0 mm such that it can inserted between the test				
	(c)	If the reference ther	mometer is used as the portable thermometer, do	oes it fit within the dummy			
	1 -	specimen as require	zd?	·····			
	/ /	AASI	CALIBRATION AND VERIFICATION	nce Laboratory			
1.		ate Diameters					
	(a)		ed to the nearest 0.01 mm?				
	(b)		to the nearest 0.02 mm (at the average of 3 diff				
	(c)		of measurements so the measurements are clearly				
	(d)		neasured dimensions entered into the DSR softwo				
			If the top and bottom plates differ in diameter, tentered?	· ·			
2.	Portable Thermometer						
	(a)		aboratory reference thermometer at least every si	x months?			
	(b)		over a range with approximately 6°C increments				
	(c)	If difference is 0.1°	C or more, are corrections written in log as part of	of the lab quality			
			the difference is $0.5^{\circ}C$ or more, the portable ther				
COMN	MENTS (1	Γ315/D7175):		(T315/D7175)			

(AASHTO T315)
(ASTM D7175)

	CALIBRATION AND VERIFICATION (Continued) Date:
	R Test Specimen Temperature
(a)	Recorded to the nearest 0.1°C
(b)	If the difference between portable thermometer and DSR controller is 0.1°C or more, is a temperature correction [ASTM: offset] determined by using the portable thermometer in a silicone wafer or an asphalt binder or other polymer dummy sample (silly putty is NOT recommended)?
(c)	Temperatures taken over a range with approximately 6°C increments?
(d)	Does the temperature vary by not more than 0.1°C over 5 minutes before a reading is taken?
(e)	Is the difference (temperature correction) either plotted vs. the temperature measured by the portable thermometer or incorporated into the instrument software?
DSI	R Torque Transducer
(a)	ASTM: verified each time the temperature offset is verified?
	Note: A newly installed or reconditioned DSR should be verified weekly until acceptable
	verification has been demonstrated. Data should be maintained in a control chart where
	the measurements are plotted versus date.
(b)	Reference fluid or manufacturer supplied fixtures used to verify the transducer at least every 6
	months and whenever the calibration is suspect?
	(1) Reference fluid 1. Complex viscosity measured with DSP within 2% of the carillery viscosity of the
	1. Complex viscosity measured with DSR within 3% of the capillary viscosity of the reference fluid as reported by the manufacturer?
	2. Complex viscosity calculated as G* divided by the angular frequency in rad/s?
	3. Reference fluid not used as a method of verifying the phase angle?
	4. When used at 10 rad/s, reference fluid only used at 64°C and above [ASTM: used between 58 and 64°C]?
	(2) Fixtures
	1. Used to verify the DSR angular displacement and torque transducers (if fixtures are available)?
	<u>PROCEDURE</u>
Drai	paration of Apparatus
(a)	Surfaces of the test plates inspected, and any plates with jagged or rounded edges or deep scratches
(u)	discarded?
(b)	Any asphalt binder residue cleaned from the plates with an organic solvent and wiped with a cotton
. ,	swab or a soft cloth dampened with acetone [ASTM: a reagent that does not leave residue]?
(c)	If necessary, a cotton swab or soft cloth used to ensure that no moisture condenses on the plates?
(d)	Test plates mounted on test fixtures and tightened firmly?
(e)	ASTM: test plates visually checked to ensure that they are parallel?
(f)	Temperature selected according to the grade of the asphalt binder or according to the preselected
	testing schedule?
(g)	DSR allowed to stabilize within ± 0.1°C of test temperature?
(h)	Zero gap level established through one of the following methods at test temperature or in the middle
	of the testing range?
	(1) Removable test plate manually spun and gap closed until plate stops spinning?
	(2) Can alogaed until the plates contact each other and the normal force is annually and the normal force is annually and the normal force in annual force is annually and the normal force in annual force is annually and the normal force in annual force is annually and the normal force in annual force is annually and the normal force in annual force is annually and the normal force in annual force is annually and the normal force is annually and the normal force is annually annual force is annually annually annually annual force is annual force in annual force is a
	 (2) Gap closed until the plates contact each other and the normal force is approximately zero? (3) Instrument zeroed automatically according to the manufacturer's procedures?

COMMENTS (T315/D7175):

(AASHTO T315)
(ASTM D7175)

PROCEDURE (Continued)

			PROCEDURE (Continued)	Date:		
2	Б					
2.	-	ration of Sample		- 1 f T214 :- 4b 1		
	(a)		ged material, if the sample is also being teste			
			not also being tested under T314, the degass			
	(b)		y fluid to pour while minimizing heating tim			
			s over 163°C [ASTM: 135°C] avoided (may			
	(c)		ed binders)?			
	(d)		irred occasionally?			
	(e)		following temperatures to promote adhesion			
	(C)		to test temperature or lowest test temperature			
			to between 34 and 46°C?			
	(f)	` '	d through one of the following methods?			
	(1)		inder into a mold that will form a pellet simil			
			wed to cool in the mold at room temperature			
			r or lower test plates?			
			ds covered during cooling to prevent contami			
		a.	Lower plate: specimen removed from the			
		b.	Upper plate: specimen in the mold press			
			upper plate, and then the mold removed			
			Note: Solvents should not be used to cle			
			specimen from the mold becomes difficu			
		3. Cool	ed on a flat laboratory bench surface at room	n temperature (not chilled)?		
		Note	: chilling is permitted for extremely soft grad	des that do not readily detach from		
		the n	nold (PG 46-34, 52-34, 58-34 for example).	These grades should not be chilled		
			nore than 10 minutes in a refrigerator or freez			
	1		ct transfer used if the binder does not easily of			
		△ △ Ctemp	erature?	mee laborator.		
			M: Testing completed within two hours of i			
			!?			
			eximately 15 mm [ASTM: 15 to 20 mm] abo			
			SHTO: only when using a DSR designed for			
			gero setting]?			
			coximately 2-mm [ASTM: a small portion] 1			
			neter?			
			HTO: Specimen allowed to harden before p			
			M: Plate mounted in the DSR immediately			
			an eye dropper or syringe may be used to to for the following of hot binder to plates using a glass or metal			
			e transferred, are the plates immediately mov			
			ble trimmed?			
	(g)		at trimming of the specimen is required?			
	(b)		flush with the plates using a heated trimmin			
	(i)		excessively hot (burns the sample) nor too co			
	(j)			b between plates equals the		
	J)	test gap plus enough to create a bulge?				
	(k)					
	` '		etermine the appropriate gap setting can be			
				-		

COMMENTS (T315/D7175):

(AASHTO T315)
(ASTM D7175)

PROCEDI	IRE ((Continued)	

- 3. Testing of Specimens
 - (a) Temperature controller set to desired test temperature including any offset?.....______
 - (1) AASHTO: If conducting a temperature sweep, testing started at the midrange test temperature?.....
 - (2) ASTM: If testing is to be performed at multiple temperatures, is testing started at the lowest test temperature for 25 mm plates, or the highest test temperature for 8 mm plates?......
 - - Note: The method for determining the correct thermal equilibration time is described in the test methods.
 - (c) Test performed in either strain controlled or stress controlled mode?....._____
 - (1) Strain controlled
 - 1. Strain value determined according to the value of the complex modulus (G^*) ?
 - 2. Strain controlled within 20% of the target value?_____
 - 3. When testing for compliance to Specification M320, appropriate strain value selected from Table 2?
 - (2) Stress controlled
 - 1. Stress value determined according to the value of the complex modulus (G*)?
 - 2. Stress controlled within 20% of the target value?_____
 - 3. When testing for compliance to Specification M320, appropriate strain value selected from Table 3?

	Table 2 – Target Strain Values			
	Material	Target, %	Range, %	
\	Original	ΔS^{12}	9 to 15	
	RTFO Residue	10	8 to 12	
	PAV Residue	1	0.8 to 1.2	

	Table 3 – Target Stress Levels			
	Material	Target, kPa	Range, kPa	
	Original	0.12	0.09 to 0.15	
	RTFO Residue	0.22	0.18 to 0.26	
	PAV Residue	50.0	40.0 to 60.0	

- (d) Specimen conditioned by applying the required strain for 8 to 16 cycles at a frequency of 10 rad/s?....___
- (e) Test measurement obtained by recording data for an additional 8 to 16 cycles?......
- (g) Data acquisition system automatically acquires and reduces the data when properly activated?.....
- (h) When conducting tests at more than one frequency, testing started at the lowest frequency and increased?.....
- (i) Testing initiated immediately after preparing and trimming the specimen?.....______
- (j) All testing completed within 4 hours of pouring the specimens?

COMMENTS (T315/D7175):

(AASHTO T315)
(ASTM D7175)

		REPORTING	Date:		
1.	Intom	prototion of Populta (Linguistry)			
1.		oretation of Results (Linearity) This section is not mandatory for the laboratory assessment of the test me	ath a d		
	(a)	Load or strain amplitude gradually increased (strain sweep)?			
	(b)	Results obtained and plotted to define linear region?			
	(c)	Graph of complex shear modulus, G*, (kPa) vs. strain (%) generated?			
	(d)	Linear region defined as 95% [ASTM: 90%] or more of zero-strain values.	ue?		
2.	AASH	ITO Report Required Information			
	(a)	Identification and description of the material tested including name cod	e, and source?		
	(b)	If the stress or strain levels in tables 2 and 3 are exceeded, is that noted			
	(c)	The test plate diameter (nearest 0.1 mm)?			
	(d)	The test gap (nearest 1 µm)?			
	(e)	The test temperature (nearest 0.1°C)?			
	(f)	The test frequency (nearest 0.1 rad/s)?			
	(g)	The strain amplitude (nearest 0.01%) or torque (nearest mN•m)?			
	(h)	The complex modulus (G*) for the 10 measurements (kPa to 3 significantly significant s			
	(i)	The phase angle (δ) for the second ten cycles (nearest 0.1 degrees)?			
	(j)	G* / sinδ to the nearest 0.01 kPa or G* sinδ to the nearest whole number			
3.	ASTM Report Required Information				
٥.	(a)	Sample identification information?			
	(b)	Operator's name?			
	(c)	Date of Test			
	(d)	Time of test?			
	(e)	Test temperature (nearest 0.1 $^{\circ}$ C)?			
	(f)	Temperature correction, if offset was applied (nearest 0.1 °C)?			
	(g)	The complex modulus (G*) (kPa to 3 significant figures)?	ce aborator	~/	
	(h)	The phase angle (nearest 0.1 degrees)?	ce raporator	y	
	(i)	For unaged and RTFO aged binders: G*/sin\delta?			
	(j)	For PAV aged binders: G*sino?			
	(k)	Indicates whether the specimen passes or fails the specification?			
	(1)	The strain amplitude (to 3 significant figures)?			
	(1)	The sham ampulate (to 3 significant figures):	•••••	····	

COMMENTS (T315/D7175):

VISCOSITY DETERMINATIONS OF ASPHALT BINDER USING ROTATIONAL VISCOMETER

(AASHTO T316)

		APPARATUS Date:		
Manufacturer:		Model:		
1.	Tester			
	(a)	Rotational viscometer?		
	(b)	Level (both torque transducer and thermal chamber if separate)?		
	(c)	Cylindrical spindles of various sizes?		
	(d)	Capable of measuring torque?		
2.	Tempe	rature Controller		
	(a)	Capable of maintaining test temperatures within 1.0°C?		
	(b)	Temperature range from 60 to 165°C or greater?		
3.	Calibra	tion and Standardization		
	(a)	Has accuracy of rotary transducer been checked using a certified reference fluid		
	()	(Newtonian fluid) of known viscosity over full range of expected test temperatures?		
		(1) Viscosity measured within ± 2 percent of the reference fluid or rotary		
		transducer recalibrated?		
	(b)	Has accuracy of temperature controller been checked with a NIST traceable measuring device?		
		<u>PROCEDURE</u>		
1.	Rotatio	nal viscometer and temperature controller turned on?		
2.		holder, sample chamber and spindle preheated according to manufacturer's recommendation?		
3.		mperature controller set to desired test temperature?		
4.		ed amount of asphalt binder as recommended by manufacturer heated until sufficiently fluid to pour?		
5.	When t	emperature controller reaches desired temperature, asphalt added to sample chamber?		
6.	Sample	chamber inserted into temperature controller unit?		
7.	Preheat	ted spindle inserted and attached using coupling?		
8.	Spindle	ted spindle inserted and attached using coupling?egently lowered into sample?		
9.		t covers the upper conical portion of the spindle?		
10.	Asphal	t brought to desired temperature within 30 minutes?		
11.	Viscon	neter speed set to 20 rpm, display set to read Pascal seconds (Pa•s)?		
12.		equilibrated at desired test temperature for at least 10 minutes?		
13.	Spindle	e rotation begun during the 10 minute equilibration period?		
14.	After th	ne asphalt has reached the specified temperature and equilibrated and once viscosity readings		
	have st	abilized, test started?		
15.	Viscosi	ty measured at one minute intervals for three minutes?		
16.	If obse	rved torque is out of range (check manufacturer's instructions for out of range torque) for		
		d spindle and speed, spindle or speed changed according to manufacturer's instructions?		
17.		ferent spindle is used, the test restarted with a new sample?		
18.	Viscosi	ty is reported as average of three readings?		
19.	Report	includes test temperature to nearest 1.0°C?		
COM	MENTS (Γ316): (T316)		
COM	TATESTAT 129 (:	(1310).		

VISCOSITY DETERMINATION OF ASPHALT AT ELEVATED TEMPERATURES USING A ROTATIONAL VISCOMETER

(ASTM D4402)

		<u>APPARATUS</u>	Date:
Manufacturer:		Model:	
1.	Rotatio	ional Viscometer	
	(a)	Capable of measuring torque required to rotate selected apparatus-measuring ge	
		at constant speed and temperature while submerged in asphalt.	
	(b)	Contains Platinum Resistance Thermometer with a probe of three- or four- wire E1137 and calibrated as a unit in accordance with E644, for calibrating the temp	
		E1137 and canorated as a unit in accordance with 2044, for canorating the temp	
2.	Appara	ratus-Measuring Geometry (Spindles) (may be of various shapes and sizes)	······
3.	Tempe	erature Controlled Thermal Chamber Heater	
	(a)	Maintains sample of asphalt at the test temperature?	······ <u></u>
4.	Sample	le Chamber (reusable or disposable)	
5.	Tampa	erature Controller	
<i>J</i> .	(a)	Capable of maintaining test temperatures within $\pm 1.0^{\circ}\text{C} (\pm 2.0^{\circ}\text{F})$?	
	(b)	Temperature range from 38 to 260°C (100 to 500°F)?	
6.	Balanc	ce readable to 0.1 g	
		o Assessors: The standard lists a PRT requirement which is confusing. The requirement is a of the thermosel itself. Check manufacturers specifications if unsure. CALIBRATION AND STANDARDIZATION	actually for the PRT which is
1.	Viscon	meter zeroed before use, or as needed, or both, according to manufacturer's instruc	tions?
2.		meter zeroed before use, or as needed, or both, according to manufacturer's instructory of viscometer checked at least annually using a certified reference fluid of knowns.	
		sity at various temperatures, using the method described by the supplier of the reference	ence fluid?
3.		ence fluid certified to be Newtonian in behavior over full range of expected	
4.		emperatures and shear rates?	
4.		g the test?	
5.		asured viscosity is not within $\pm 2\%$ of the certified value, is the viscometer recalibration	
6.		racy and stability of temperature controller checked at least every 6 months?	
7.		erature checked using asphalt sample or high flash point oil equilibrated to within 5	
8.		t temperature(s)?le temperature measured to within ± 0.1 °C (± 0.2 °F) using NIST traceable measuring	
0.		scribed in Method E644?	
9.		erature set point offset accordingly?	
COM	MENTS ((D4402):	(D4402)

VISCOSITY DETERMINATION OF ASPHALT AT ELEVATED TEMPERATURES USING A ROTATIONAL VISCOMETER

(ASTM D4402)

	PROCEDURE Date:	
1.	Manufacturer's instructions for the operation of the instrument followed?	
2.	Instrument warmed up for at least 5 minutes before any calibrations or analyses conducted?	
3.	Temperature controller set to desired temperature, taking into account any offset?	
4.	Apparatus-measuring geometry selected to produce torque between 10 and 98% of instrument capacity at selected speed?	
	Note: Measurements will generally be more accurate at higher torque readings.	
5.	Preferably, sample chamber and apparatus-measuring geometry preheated until temperature	
	has equilibrated for at least 15 min (Filled Asphalts: this is mandatory)?	
6.	Volume of sample added to sample chamber as specified by manufacturer for apparatus-	
	measuring geometry used (Filled Asphalts: and thoroughly stirred)?	
7.	Sufficient sample volume to ensure measuring portion of apparatus-measuring geometry completely immersed without overfilling sample chamber?	
8.	Manufacturer's instructions followed to ensure accurate sample volume?	
9.	Selected apparatus-measuring geometry inserted into sample and coupled to viscometer?	
10.	Manufacturer's instructions followed to ensure proper alignment?	
11.	Sample brought to desired temperature within 30 min and allowed to equilibrate at the desired test	
	temperature for at least 10 min before beginning the measurements (Filled Asphalts: and rotation	
10	started immediately)?	
12.	scale instrument capacity?	
13.	Speed maintained and sample allowed to equilibrate for an additional 5 min?	
14.	Temperature does not deviate by more than $\pm 1.0^{\circ}$ C ($\pm 2.0^{\circ}$ F) during this conditioning period?	
15.	Viscosity or torque measured at 1-min intervals for a total of three min?	
16.	Steps 11 through 15 repeated for each test temperature required (Filled Asphalts: A new, freshly	
	stirred sample is required for each test temperature)?	
17.	If torque readings are above 98% of instrument capacity at the lowest temperature either of	
	the following performed?	
	(a) Speed of rotation increased and the test continued?	
	(b) Steps 5 through 15 repeated with a smaller diameter geometry and appropriate sample volume?	
18.	If torque reading is below 10% of instrument capacity at the highest temperature either of the	
	following performed?	
	(a) Speed of rotation increased and the test continued?	
10	(b) Steps 5 through 15 repeated with a larger diameter geometry and appropriate sample volume?	
19.	If viscosity units not read out directly by instrument, torque readings multiplied by the appropriate factor to obtain viscosity values?	
20.	Arithmetic average of three readings calculated and rounded to three significant figures?	
21.	If digital output displayed in centipoises (cP), result multiplied by 0.001 to obtain value in	
	Pascal seconds (Pa·s)?	
22.	For instruments that offer automation, results of a 3-min integration accepted?	
	Note: Filled asphalts refer to asphalt with a powdered limestone additive and are used	
~~-		
COMN	(D4:02):	402)

COMMENTS (D3289):

RESIDUE OF SPECIFIED PENETRATION

(D243)

		<u>APPARATUS</u>	Date:	
1.	Heatir	ng (air) bath		
1.	(a)	Cast iron (or equivalent)?		
	(b)	Inner diameter or top opening 71.4 mm (2 3/16 in.)?		
	(c)	Outer diameter of top opening 90.5 mm (3 9/16 in.)?		
	(d)	Inner diameter of bottom opening: 84.1 mm (3 5/16 in.)?		
	(e)	Total height: 38.1 mm (1.5 in.)?		
	(f)	Height from top bolt shank to the bottom of the bath at least 6.4 mm (.25		
	(g)	Flange thickness 3.2 mm (1/8 in.)?		
	(h)	Opening 1.6 mm (1/16 in.) larger than container?		
2.	Hot Pl			
	(a)	Bunsen burner under plate on ring stand with pressure regulator?		
or	(b)	Conventional type hot plate with rheostat?		
3.	Therm	nometer and Support		
	(a)	ASTM 11F or 11C thermometer used?		
	(b)	Suitable thermometer support?		
	(c)	Thermometer equidistant from container sides?		
	(d)	Bottom of bulb less than 6.4 mm (1/4 in.) above can bottom, but not touc		
4.	Contai			
	(a)	70 mm diameter, 45 mm deep (6 oz. ointment box)?		
		<u>PROCEDURE</u>		
1.	Sampl	e as received thoroughly stirred and agitated before removing test sample?.		
2.		iner (6 oz. ointment box) tared?		
3.	Test sa	ample of 100.00 ± 0.10 g weighed into container?		
4.	Contai	iner placed in air bath?		
5.		nometer supported centered less than 1/4 inch from bottom?		
6.	Test sa	ample heated rapidly without foaming to 249°C (480°F)?		
7.	During	g evaporation, temperature maintained between 249 and 260°C (480 and 50	00°F)?	
8.	Sampl	e stirred occasionally with thermometer to avoid local overheating?		
9.		rdened bitumen fluxed during heating and stirring?		
10.		thought that residue has reached required penetration, bitumen on thermom		
		y scraped off returned to container?		
11.		iner cooled and weighed?		
12.		ration of residue determined by D5 (AASHTO T49) using evaporation conta		
13.		etration of residue is greater than required, all water removed from the conta		
		e surface of the test sample?		
14.		4 through 13 repeated until penetration of residue within 15 of 100?		
15.		ntage of residue by weight calculated?		
16.		ted as percentage of residue of penetration (determined), stating first the spe		
		ation, and second the penetration actually determined or calculated by inter-		
		e sample tested?		
		1		

Revised 2014-06-02

(D3289)

COMMENTS (D3289):

SPECIFIC GRAVITY OF SEMI-SOLID AND SOLID BITUMINOUS MATERIALS BY NICKEL CRUCIBLE

(D3289)

		<u>APPARATUS</u>	Date:
Crucik	ole made	of nickel, high-form, 30 mL capacity, approximately 43 mm hi	oh x 41 mm diameter
		m]?	
		rature bath: ± 0.1 °C (± 0.2 °F)?	
		distilled water?	
		rith graduations of at least 0.1°C (0.2°F) [such as ASTM 63C of	
		evice of equal accuracy?	
		alibrated?	
		low-form beaker?	
		B (0.001 g)?	
		ending crucible from balance?	
		chang cracios from caranee	
T dir st	radare		
		<u>PROCEDURE</u>	
Baske	t Preparat		
(a)		ole placed in wire basket?	
(b)		t suspended from arm of balance?	
(c)	Weigh	at of empty crucible + basket in air (W_1) determined to 0.001 g	?
(d)		r filled with distilled water and placed on pan straddle?	
	(1)	Freshly boiled distilled water?	
		Water temperature 25 ± 1 °C, 15.0 ± 1 °C, or 15.6 ± 1 °C?	
	(3)	600 mL Griffin low-form beaker?	
(e)		t containing crucible suspended from balance arm so that cruci	
(f)	Weigh	at of empty crucible + basket immersed in water determined (W	(2)?
(g)		ole removed from basket and dried?	
	nen Prepa	ala malead.	
(a)		als melted: Temperature no more than 55°C or 100°F above the softeni	nea Laboratory
	(1)		
	(2)	Temperature no more than 110°C or 200°F above the soften Heated no more than 30 minutes?	
(b)	(3)	Heated no more than 30 minutes?	
(b)	(1)	In oven for 10 minutes?	
	(2)	Oven at 120°C (250°F)?	
(c)		oven at 120 C (250 F)?	
(d)		e cooled to ambient temperature for at least 40 minutes?	
Weigh		e cooled to amolent temperature for at least 40 minutes?	
(a)		1(b) - 1(d) performed?	
(b)		tt of sample + crucible + basket in air determined (W)?	
(c)		at of sample + crucible + basket immersed in water determined	
(-)	(1)	Sample in crucible tempered for at least 30 minutes in water	
	(-)	at $25 \pm 1^{\circ}$ C, $15.0 \pm 1^{\circ}$ C, or $15.6 \pm 1^{\circ}$ C?	
	(2)	Weighed in bath in 600 mL Griffin low-form beaker?	
	(3)	Freshly boiled distilled water?	
	(4)	Water temperature 25 ± 0.1 °C, 15.0 ± 0.1 °C, or 15.6 ± 0.1 °C	
Calcul		5, 2010 = 011 G, 01 1010 = 011	
(a)		$= (W-W_1)/[(W-W_1)-(W_3-W_2)]?$	

(D3289)

TOUGHNESS AND TENACITY OF BITUMINOUS MATERIALS

(D5801)

A DDA D ATLIC	D .
<u>APPARATUS</u>	Date:

Tension Head	1	2	3	4	5	6
Polished metal, hemispherical head?						
Radius 11 mm (7/16 in.)?						
Stem diameter 6.4 mm (1/4 in.)?						
Stem length approximately 33 mm (1 5/16 in.) [AMRL: ± 3 mm]						
Threaded and fitted with a lowering screw?						
Fitted with a small pin?						
Spider						
Stem moves freely and parallel to the axis?						
Groove for the pin on the stem?						
Three equidistant and notched arms?						

1.	Contain	er
1.	(a)	55 mm diameter, 35 mm deep (3-oz ointment box)?
2.	Testing	Machine
	(a)	Maker:
	(b)	Serial Number (or other ID):
	(c)	Capable of maintaining a rate of 50 cm/min. (20in./min) to within 2%?
	(d)	Capable of recording the force vs. elongation curve?
	(e)	Maximum capacity of at least 45 kg (100 lb) (Polymer modified asphalts: 90 kg (200 lb))?
	(f)	Holds the sample container firmly in place?
	(g)	Minimum effective pull length of 61 cm (24 in.) after installing the sample holder?
3.	Water E	AASHTO Materials Reference Laboratory
٥.	(a)	Maker:
	(b)	Capable of maintaining a temperature at 25 ± 0.1 °C (77 ± 0.18 °F)?
	(c)	Perforated shelf not less than 50 mm (2 in.) from the bottom of the bath?
	(d)	Perforated shelf not less than 100 mm (4 in.) below the liquid level?
4.	Oven ca	spable of maintaining temperature at 163 ± 5.5 °C (325 ± 10 °F)?
5.	Thermo	meter
	(a)	ASTM 63C or 63F?
		Note: If a 77°F (25°C) penetration bath is used, any thermometer or thermometric device with 0.1°C
		(0.2°F) subdivisions may be used
	(b)	Calibrated?
COMN	MENTS (E	(D5801):

TOUGHNESS AND TENACITY OF BITUMINOUS MATERIALS

(D5801)

		PROCEDURE Date:
1.	Sample	Preparation
1.	(a)	Heated in loosely covered container in an oven at 163° C (325° F)?
	(u)	(1) Uniform temperature and sufficiently fluid to pour?
		(2) Local overheating prevented?
	(b)	Sample stirred without incorporating air bubbles?
	(c)	36 ± 0.5 g poured into each of three sample containers?
	(d)	Tension head mounted in spider and immediately placed in each container?
	(e)	Tension head lowered until asphalt level is approximately 1 mm below diameter of the head?
	(f)	Container, tension head, and spider assemblies placed in a 163° C (325°F) oven for 15 minutes?
	(g)	Containers removed and tension heads lowered until the asphalt is level with
	ν	the diameter of the head?
	(h)	Cooled at room temperature for 75 ± 5 minutes?
	(i)	Placed in a 25° C (77° F) water bath for 75 ± 5 minutes?
2.	Tought	ness and Tenacity Test
	(a)	Chart pen zeroed and testing machine prepared?
	(b)	Sample container removed from water bath and immediately placed in testing machine?
	. ,	Note: The centering spider may be removed before the sample is placed in the machine
	(c)	Room temperature at $25 \pm 3^{\circ}$ C $(77 \pm 5^{\circ}$ F)?
	(d)	Transfer time less than 3 minutes?
		Note: Water may be left in the container to prevent surface cooling
	(e)	Tension head pulled from the sample at 50 cm/min. (20 in/min.)?
	(f)	Force vs. elongation curve recorded?
	(g)	Test continued until one of the following?(1) Asphalt column breaks?
	1 -	(2) Force returns to zero?
	1	(3) Extension limit of the machine is reached?
		AASHTO Materials Reference Laboratory
3.	Report	
	(a)	Three values averaged (Newton-meters or inch-pounds) for toughness and tenacity?
	(b)	If one breaks prematurely, that sample is considered invalid and the two valid
		values are averaged?
		Note: Toughness is calculated as the complete area under the curve before the end of the test
		(total work required to separate the specimen). Tenacity is calculated as the area under the
		curve excluding the initial resistance (work required to stretch the specimen after the initial
		resistance is overcome.)
COM	MENTS (1	D5801): (D5801)

MULTIPLE STRESS CREEP AND RECOVERY (MSCR) OF ASPHALT BINDER USING A DYNAMIC SHEAR RHEOMETER

(D7405)

	APPARATUS Date:
1.	Apparatus and materials for running ASTM test method D7175?
2.	Manufacturer-supplied certificate stating that full torque is achieved within 0.003 seconds from
۷.	the start of the loading cycle?
	the start of the foating cycle.
	<u>PROCEDURE</u>
Sample	Preparation Preparation Preparation
1.	Samples prepared in accordance with ASTMD7175?
	Note: This test may be run on a specimen previously used for ASTM D7175.
Procedu	
1.	Specimen allowed to reach thermal equilibrium at the desired test temperature as in ASTM D7175?
2.	If the specimen was previously used for ASTM D7175, allowed to remain unloaded for at least 1 minute
2	before starting the creep recovery test?
3. 4.	Specimen loaded at a constant creep stress of 0.100 kPa for 1.00 seconds?
4. 5.	After creep stress is applied, specimen is allowed to recover (zero stress) for 9.00 seconds?
<i>5</i> . 6.	Stress and strain recorded at least every 0.10 seconds during the creep cycle?
7.	Stress and strain recorded at least every 0.45 seconds during the ercovery cycle?
8.	One of the following:
0.	(a) Data points at 1.00 and 10.00 seconds explicitly recorded?
	(b) DSR software extrapolates prior data to determine the strain value at the required time?
9.	If data points are extrapolated, data shall include a measured data point no more than 0.05 s prior to
	1.00 seconds, and no more than 0.30 seconds prior to 10.00 seconds?
10.	Creep and recovery cycle described above repeated nine more times?
11.	No rest period allowed between cycles?
12.	Ten more creep and recovery cycles repeated at a load of 3.200 kPa?
13.	No rest period allowed between 0.100 kPa and 3.200 kPa loading cycles?
14.	No rest period allowed between 0.100 kPa and 3.200 kPa loading cycles?
	(a) Initial strain value at the beginning of the creep portion of each cycle (denoted ε_0)?
	(b) Strain value at the end of the creep portion (after 1.0 s) of each cycle (denoted ε_c)?
	(c) The adjusted strain value at the end of the creep portion (after 1.0 s) of each cycle ($\varepsilon_1 = \varepsilon_c - \varepsilon_0$)?
	(d) The strain value at the end of the recovery portion (after 10.0 s) of each cycle? (denoted ε_r)?
	(e) The adjusted strain value at the end of the recovery (after 10.0 s) of each cycle ($\epsilon_{10} = \epsilon_r - \epsilon_0$)?

COMMENTS (D7405): (D7405)

MULTIPLE STRESS CREEP AND RECOVERY (MSCR) OF ASPHALT BINDER USING A DYNAMIC SHEAR RHEOMETER

(D7405)

	PROCEDURE (Continued) Date:
Calcula	ions - are the following calculations done according to the test method?
1.	Percent recovery for each of the ten cycles at 0.100 kPa ($\varepsilon_r(100, N)$)?
2.	Percent recovery for each of the ten cycles at 3.200 kPa ($\varepsilon_{\rm r}(3200,{\rm N})$)?
3.	Average percent recovery at 0.100 kPa (R100)?
4.	Average percent recovery at 3.200 kPa (R3200)?
5.	Percent difference is recovery between 0.100 kPa and 3.200 kPa (R _{diff})?
6.	Non-recoverable creep compliance $(J_{nr}(100, N))$ for each of the ten cycles at 0.100 kPa?
7.	Non-recoverable creep compliance ($J_{nr}(3200, N)$) for each of the ten cycles at 3.200 kPa?
8.	Average non-recoverable creep compliance at 0.100 kPa (J _{nr} 100)?
9.	Average non-recoverable creep compliance at 3.200 kPa (J _{nr} 3200)?
10.	Percent difference in non-recoverable creep compliance between 0.100 kPa and 3.200 kPa (J _{nr-diff})?
Report -	does report include the following information?
1.	Sample identification information?
2.	Test temperature, to the nearest 0.1°C?
3.	Average percent recovery at 0.100 kPa, R100, to the nearest 0.1%?
4.	Average percent recovery at 3.200 kPa, R3200, to the nearest 0.1%?
5.	Percent difference between average recovery at 0.100 kPa and 3.200 kPa, R _{diff} , to the nearest 0.1%?
6.	Non-recoverable creep compliance at 0.100 kPa, J _{nr} 100, to three significant figures?
7.	Non-recoverable creep compliance at 3.200 kPa, J _{nr} 3200, to three significant figures?
8.	Percent difference between non-recoverable creep compliance at 0.100 kPa and
	3.200 kPa, J _{nr-diff} , to nearest 0.1%?
COMM	ENTS (D7405): (D7405

AASHTO Materials Reference Laboratory

(D7553)

SOLUBILITY OF ASPHALT MATERIALS IN N-PROPYL BROMIDE

		<u>APPARATUS</u>	Date:		
1.	Gooch	n Crucibles			
	(a)	Glazed surface throughout except bottom exterior unfinished?			
	(b)	Approximately 44 mm at the top tapering to 36 mm at the bottom [AMRL:			
	(c)	Approximate depth of 20 to 30 mm?			
2.	Filtrat	ion Assembly			
	(a)	Heavy walled filter flask with side tube, capacity 250 to 500 mL?			
	. ,	Note : any other assemblies permitting vacuum filtration with a Gooch cruc			
	(b)	Glass fiber pads with a diameter of 32 to 34 mm, fine porosity, fast flow rate			
		1.5 µm particle retention?			
3.	Suctio	on Assembly			
	(a)	Satisfactory assembly?			
4.	N-Pro	pyl Bromide Solvent			
	(a)	N-Propyl Bromide, Technical grade (conforming to ASTM D6368)?			
5.	Desiccator				
	(a)	Satisfactory design and charged with effective desiccant?			
6.	Drying	g Oven			
	(a)	Maintains temperature at 110 ± 5 °C (230 ± 9 °F)?			
7.	Misce	llaneous Items			
	(a)	Suitable container for weighing and dissolving sample?			
	(b)	Class A balance (readable to 0.0001 g) available?			
	(c)	Policeman (optional)?	<u></u>		
		PREPARATION OF THE GOOCH CRUCIBLE			
1.	New f	filter pad placed in crucible and dried in oven at 110 ± 5 °C for 15 min			
	(no we	etting and seating required)?			
2.		d in a desiccator for 30 ± 5 and weighed to the nearest 0.0001 g?			
3.		l in a desiccator until ready to use?			
		AASHTO Materials Reference	Laboratory		
COM	MENTS ((D7553):	(D7553)		

(D7553)

SOLUBILITY OF ASPHALT MATERIALS IN N-PROPYL BROMIDE

PROCEDURE Date: 1. If the sample is not fluid is it heated with care to prevent local overheating?..... 2. Sample stirred occasionally and the entrapment of air avoided?..... Sample heated at any temperature not more than 100°C above softening point?..... 3. Approximately 2 g [AMRL: ± 0.5 g] of sample placed in tared (nearest 0.001 g) container? 4. 5. Container with sample allowed to cool and then weighed to nearest 1 mg (0.001 g)?..... 100 mL of solvent added to container, flask stoppered, and then container agitated 6. as necessary until the sample is dissolved? 7. Solvent added in small portions with constant agitation?.... Lumps completely digested and container sides free of undissolved sample? 8. 9. Container stoppered and set aside for at least 15 min?..... **Note**: for referee testing, flask and solution shall be placed in a water bath at 38.0 ± 0.3 °C for 1 hour before filtering. Please discuss with laboratory if necessary. Crucible placed in filter tube and wetted? 10. Asphalt solution decanted through filter with light suction? 11. If insoluble matter is visible: 12. Retained in container until solution has drained through filter?..... (a) Container washed with solvent and insoluble matter transferred to crucible?..... (b) (c) Container and policeman (if used) rinsed?..... Insoluble matter washed until the filtrate is substantially colorless?......______ (d) Strong suction applied to remove remaining solvent?..... (e) Crucible removed and bottom washed free of dissolved matter? 13. Placed in oven at 110 ± 5°C for at least 20 min? 14. Cooled in desiccator for 30 ± 5 min and then weighed to nearest 0.0001g?.... 15. Steps (14) and (15) repeated until constant mass of ± 0.0003 g obtained?..... 16. 17. Percent insoluble reported to nearest 0.1%? If percent insoluble is less than 1%, reported to nearest 0.01%?..... 18. 19. If the crucible in desiccator must be left overnight, is it placed in oven for 30 min. and then cooled again in desiccator?..... ASH IO Materials Reference Laborator

COMMENTS (D7553): (D7553)

DETERMINING THE CONTINUOUS GRADING TEMPERATURES AND CONTINUOUS GRADES FOR PG GRADED ASPHALT BINDERS

(D7643)

		<u>APPARATUS</u>	Date:		
1. o	(a) Method D71	75 (Dynamic Shear Rheometer) 48 (Bending Beam Rheometer)			
2.	(a) Method D28	equired:			
		<u>PROCEDURE</u>			
1.	(a) The difference (b) The two tem Note: For example: A P	ermined (T ₁ and T ₂) for each specification property ce between the two temperatures is 6°C? peratures bracket the specification requirements? G 64-XX tested for G*/sinδ at 64°C and 70°C may giv pectively. These results bracket the specification require	/e test results		
2.	If the result of the applicable specification provides a temperature rather than a limiting property value, the temperature used as the continuous grading temperature?				
<u>Perfo</u> 3.	The interpolation for	termine Continuous Grading Temperatures (all process of the properties calculated using linear relation $-\log_{10}\left(P_{1}\right)) / \left(\log_{10}\left(P_{2}\right) - \log_{10}\left(P_{1}\right)\right)] * (T_{2} - T_{1})$	onship using formula below?		
<u>Perfo</u>	rming Interpolation to De	termine Continuous Grading Temperatures (m-valu	ue only)		
4.	The interpolation for each of the properties calculated using arithmetic scale using formula below?				
	Note: Because the proper linear interpolation resulting T_c = Continuous grading T_1 = Lower of the two to P_S = Specification requirespective PG grad	gns for temperatures below 0°C. erties are a non-linear function of temperature, lts in a slight error in the estimated values of $T_{\rm C.}$.			
		P_2 = Test result for the specification property in question at T_2			

COMMENTS (D7643):

 T_2 = Higher of the two test temperatures

(D7643)