EMULSION INDEX				
			Status (√,	
Test Method	AASHTO/ASTM	Pages	NP)	Technician
FREEZING TEST	T59 / D6929	2		
SETTLEMENT AND STORAGE STABILITY	T59 / D6930	3		
SIEVE TEST	T59 / D6933	4		
RESIDUE BY EVAPORATION	T59 / D6934	5		
CEMENT MIXING	T59 / D6935	6		
DEMULSIBILITY	T59 / D6936	7		
DETERMINING DENSITY	T59 / D6937	8		
RESIDUE BY DISTILLATION	T59 / D6997	9-10		
EVALUATING AGGREGATE COATING	T59 / D6998	11		
PARTICLE CHARGE	T59 / D7402	12		
SAYBOLT FUROL VISCOSITY @ 25°C	T59 / D7496	13-15		
SAYBOLT FUROL VISCOSITY @ 50°C	T59 / D7496	13-15		
SWEEP TEST	/ D7000	16-18		

^{*} NP for "Not Presented"

AASHTO Materials Reference Laboratory

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

FREEZING OF EMULISFIED ASPHALTS

(139)	
(D6929)	

	APPARATUS Date:
1.	Freezer capable of maintaining test temperature of -18 ± 5°C?
2.	Metal container with close fitting lid (such as a 500-mL press-top can)?
3.	Glass stirring rod?
	<u>PROCEDURE</u>
1.	Approximately 400g of emulsion poured into a clean metal container?
2.	Closed metal container with emulsion placed in freezer for 12 to 18 consecutive hours?
3.	Container removed from freezer and allowed to thaw by exposure to ambient temperature?
4.	Freezing and thawing periods repeated an additional two times (total of three cycles)?
5.	After third cycle, emulsion stirred?
6.	Asphalt examined and reported as either <i>Homogeneous</i> or <i>Broken</i> based on whether or not stirring at room temperature brought any separated emulsion back into a homogeneous state?



SETTLEMENT AND STORAGE STABILITY

(139)	
(D6930)	

	APPARATUS Date:
1.	Two 500 mL glass cylinders [ASTM: Only one required]?
	(a) $50 \pm 5 \text{ mm O.D.}$?
	(b) AASHTO Only: 5 mL graduations?
	(c) Cork or glass stoppers [ASTM: rubber stoppers are acceptable]?
	Note to Assessor: Cylinders equipped with side-arms are acceptable – AMRL
2.	A glass tube pipette or siphon assembly capable of removing and delivering 55 mL of sample
	[ASTM: 50 ml glass tube pipette]?
2	Note to Assessor: Not necessary if cylinder with side-arms is used
3. 4.	Four [ASTM: Only two required] 1000 mL glass, metal beakers or containers of similar dimensions?
4.	for residue by evaporation determination?
5.	Additional stir rods; glass or stainless steel, with rounded ends
5.	for stirring settlement and storage stability sample?
6.	Oven: maintained at 325 ± 5 °F (163 ± 3 °C)?
0. 7.	Class G2 balance conforming to the requirements of M231?
7.	Class G2 balance combining to the requirements of Wi231?
	<u>PROCEDURE</u>
1.	Sample brought to room temperature (Storage Stability: 70 to 80 °F (21 to 27 °C)
	[ASTM: 22 to 28 °C (72 to 83 °F)]?
2.	500 mL representative sample placed in each of 2 cylinders [ASTM: Number of cylinders optional]?
3.	Cylinders allowed to stand undisturbed at room temperature
	(Storage Stability: 70 to 80°F (21 to 27°C) [ASTM: 22 to 28 °C (72 to 83 °F)]?
	(a) Cylinders sealed airtight?
	(b) 24 hours for storage stability?
	(c) 5 days for settlement?
4.	Approximately 55 mL of emulsion pipetted or siphoned from the top of each cylinder
	(drained if using cylinders with side arms) without disturbing remainder of sample?
	(a) Each 55 ml portion thoroughly stirred?
	(b) 50.0 ± 0.1 g of each sample weighed into a separate beaker
	or container that has been previously weighed with a stirring rod?
5.	Content of each beaker or container evaporated by procedure specified under
	Residue by Evaporation and percent residue calculated? (A = top)
6.	Approximately next 390 mL siphoned (or drained if using cylinders with side arms) from each cylinder?
7.	Thoroughly mix the remaining emulsion in the cylinders?
8.	50.0 ± 0.1 g of each sample weighed into a separate beaker or container that
	has been previously weighed with a stirring rod?
9.	Content of each beaker or container evaporated by procedure specified under
	Residue by Evaporation, and percent residue calculated? (B = bottom)
10.	Storage stability or settlement for the cylinder calculated as follows:
	(a) Storage Stability, $\%$ (24 hrs) = B – A
	(b) Settlement, $\%$ (5 days) = B – A
11.	Report the storage stability as the average of the two individual
	cylinder results? [ASTM: Averaging Optional]?

CI	FV	F	TI	TP	1

(T59)	
(D6933)	

		<u>APPARATUS</u>	Date:
1.	2 in .	liameter, 850-μm (No. 20) sieve?	
2.			
2. 3.		ow metal pan or container to fit bottom of sieve?iner suitable for l kg of emulsion?	
3. 4.		TO: Solution of 2 percent sodium oleate dissolved in distilled water fo	
4. 5.		TO: Solution of 2 percent solution ofeate dissolved in distitled water for testing cationic emulsions?	
<i>5</i> . 6.		1: 1% solution of nonionic surfactant solution, (ethoxylated nonylph	
0.		f nonionic surfactant dissolved in distilled water and diluted to 100 m	
7.		cator equipped with desiccant?	
8.		maintained at 220°F (105°C) [AASHTO: 105 ± 5 °C (220 ± 9 °F)]?	
9.		Thatmanded at 220 T (103 C) [WishT10. 103 ± 3 C (220 ± 5 T)] TO: Oven or hot water bath at 50 ± 3 °C (122 ± 5 °F) if it is necessary	
<i>)</i> .		It prior to testing?	· ·
10.	Balan	1	
10.	(a)	Class G5 balance?	
	(u)	(Note to Assessors: More precise balances with capacity to weigh t	
	(b)	Class G2 balance available? (For weighing the sieve and residue)	
11.	. ,	nometers	
	(a)	ASTM 17C thermometer for tests at 25 °C?	
	(b)	ASTM 19C thermometer for tests at 50 °C?	
	(c)	Or any other thermometric device of equal accuracy?	
	(-)		
1.	(a) (b)	Temperature Room temperature for samples whose viscosity is 100 s or less where Test temperature at $122 \pm 5^{\circ}F$ ($50 \pm 3^{\circ}C$) for samples whose viscosity is $100 \times 100^{\circ}$ for	ty is greater
		than 100 s or whose viscosity is specified at 122°F (50°C)?le stirred to achieve homogeneity?	
2.	Samp	le stirred to achieve homogeneity?	rice Laboratory
3.	Weig	ht of 850-μm (No. 20) sieve and pan determined (= A)?	
4.		ASTM: 800-1000 g] (1000 g) of emulsion weighed into suitable contains	
_		determined (=C)]?	
5.		cloth wet with appropriate liquid?	
	(a)	AASHTO: With 2 percent sodium oleate solution for anionic emulsi	
	(b)	AASHTO: With distilled water for cationic emulsions?	
6.	(c)	le poured through sieve?	
0. 7.		ue on sieve [AASHTO: and container] washed with appropriate liquid washed washed with appropriate liquid washed was	
7.	(a)	With 2 percent sodium oleate solution for anionic emulsions?	
	(a) (b)	With distilled water for cationic emulsions?	
	(c)	ASTM: Distilled or deionized water for both anionic and cationic	
8.	\ /	As Mass of empty container determined (=D)?	
9.		laced under sieve?	
10.		nd sieve heated for 2 hr. in 220°F (105°C) drying oven?	
11.		nd sieve cooled in a desiccator?	
12.		ht of 850-μm (No. 20) sieve, pan, and residue weighed (= B)?	
13.		TO: Percentage sample retained on sieve calculated as: $\{(B - A)/10\}$	
14.		1: Percentage sample retained on the sieve calculated as: $\{(B-A)/(B-A)\}$	
			=

		(59)
	(D69))34)
	APPARATUS Date:	
1.	Beakers, low form, 1000 mL capacity, made of glass or metal (quantity as required for the test)?	
2.	Glass rods, flame polished ends, approximately 7 in. long x 1/4 in. in diameter (one rod per beaker)?	
3.	300-μm (No. 50) sieve (only required when the residue is to be further tested)?	
4.	Oven?	
	(a) Thermostatically controlled?	
	(b) Maintained at $325 \pm 5^{\circ}F$ ($163 \pm 3^{\circ}C$)?	
5.	Class G2 [ASTM: GP2] balance conforming to the requirements of M231?	
	<u>PROCEDURE</u>	
1.	Three beakers used if only the amount of residue is to be determined, four beakers if further testing is to	
1.	be performed on the residue?	
2.	Each beaker and stirring rod weighed to 0.1 g?	
3.	50 ± 0.1 g of thoroughly mixed emulsion weighed into each beaker?	
3. 4.	Water initially evaporated from the beakers or containers by one of the following methods:	
4.	(a) Heating in an oven at $325 \pm 5^{\circ}$ F ($163 \pm 3^{\circ}$ C) for 2 hrs?	
	(a) Heating in an oven at 323 ± 3 F (103 ± 3 C) for 2 lifs?	
	(c) Heating in a cold or warm oven and gradually bringing oven and sample up to	
	a temperature of 325°F (163°C)?	
5.	Initial evaporation performed in a manner that prevents loss of asphalt from the beaker	
<i>J</i> .	through foaming and splattering?	
6.	After initial evaporation, residue thoroughly stirred with pre-weighed glass rod?	
7.	Beakers, with rod and residue, replaced in oven at $325 \pm 5^{\circ}$ F ($163 \pm 3^{\circ}$ C) for one hour?	
8.	Beakers, with rod and residue, allowed to cool to room temperature and weighed to 0.1 g?	
9.	Percent residue calculated for each beaker?	
10.	Percentage of residue by evaporation reported as the average of the results?	
11.	Percentage of residue by evaporation reported as the average of the results? If further testing is to be performed on the material, are the following steps taken?	Ty_

COMMENTS:

(b)

CEMENT MIXING

(T59)_	
(D6935) _	

	APPARATUS Date:
1.	Sieves:
	(a) A 180-µm (No. 80) sieve?
	(b) A 3 in. diameter 1.40-mm (No. 14) sieve?
	(c) A shallow pan for 3 in. diameter sieve?
2.	A graduated cylinder with 100 mL capacity?
3.	Mixing bowl made of glass or metal, capacity approximately 500 mL?
4.	Stirring rod made of steel, approximately ½ in. (13 mm) [ASTM: approximately 10 mm] diameter, with rounded ends?
5.	Type III Portland Cement conforming to ASTM C150 (AASHTO M85)?
6.	Class G2 [ASTM: GP2] balance?
7.	Oven?
	(a) Thermostatically controlled?
	(b) Maintained at 325 ± 5 °F (163 ± 3 °C)?
	(c) AASHTO: Conforming to ASTM E145, Type 1B
8.	ASTM: Thermometer: a thermometric device capable of measuring the temperature of the oven and the
	emulsified asphalt to the nearest 1 ${f C}$?
	PROCEDURE
1.	Emulsion diluted to 55 percent residue?
	(a) Calculation of percent residue based on determination by:
	(1) Distillation?
	(2) Residue by Evaporation?
	(b) Diluted with calculated amount of distilled [ASTM: or deionized] water?
	(c) Calculation determined by %Water = 100 – [(55 / %Residue) x 100]
2.	Cement sieved through 180- μ m (No. 80) sieve?
3.	50.0 ± 0.1 g cement passing 180-µm (No. 80) sieve weighed into mixing bowl?
4.	Ingredients and apparatus brought to approximately 77°F (25°C) [AMRL: ± 9°F (±5°C)] before mixing?
5.	100 mL of diluted emulsion added to cement in mixing bowl?
6.	Mixture immediately stirred with steel rod, in a circular motion at approximately 60 rpm for one minute?
7.	After stirring 1 min., 150 mL distilled [ASTM: or deionized] water added?
8.	Stirring continued for additional 3 min. period?
9.	Tare weight of 1.40-mm (No. 14) sieve and shallow pan determined?
10.	Mixture poured through 1.40-mm (No. 14) sieve?
11.	Bowl rinsed over sieve with distilled [ASTM: or deionized] water until clean?
12.	Sieve rinsed with distilled water poured from a height of approximately
	150 mm (6 in.) until water is clear?
13.	Sieve placed in shallow pan and heated at 325°F (163°C) in oven?
14.	Heating and weighing repeated until successive weights differ by no more than 0.1 g?
15.	Weight of material retained on sieve and in pan reported as the percentage
	of break in the cement mixing test?

DEMULSIBILITY

(T59)	
(D6936)	

		<u>APPARATUS</u>	Date:
1.	Three pieces [ASTM: one] of 1	.40-mm (No. 14) wire cloth, unframed,	
	*	RL: sufficient to completely cover beaker]?	?
2.		ity metal beakers?	
<i>3</i> .		ther suitable metal container with a minin	
4.		5/16 in. (7.9 mm) [AMRL: $\pm 1/8$ in. (± 3.2	
		ely 10 mm diameter, with rounded ends? .	
5.		n 0.1 mL intervals?	
6.		nic emulsions (as appropriate to the materia	
		L), prepared with distilled water?	
	(b) CaCl ₂ solution (5.55 g/s)	L), prepared with distilled water?	
		ntainer?	
7.	Demulsifying solutions for catio		
	(a) Dioctyl sodium sulfosu	ccinate solution (8g/L), in distilled water?.	······
		rk glass [ASTM: or impermeable plastic]	
	a cool dark lo	cation] and less than 90 days old?	······· <u> </u>
8.	Class G2 [ASTM: GP2] balance	e conforming to the requirements of M231	?
9.	Oven capable of maintaining 16	3 ± 3 °C (325 ± 5°F)?	······· <u> </u>
10.	AASHTO: Timer graduated in 0	0.1 s and accurate within 0.1 percent when	tested over a 15 minute interval?
		PROCEDURE	
1.		etermined?	
2.		beaker, a stirring rod, and a wire cloth we	
		uired]?	
3.	100.0 ± 0.1 g of sample weighed	I into each of three assemblies?d reagent] brought to 77.0 ± 1.0 °F (25.0 \pm	
4.	Weighed samples [AASHTO: and	d reagent] brought to 77.0 ± 1.0 °F ($25.0 \pm$	0.5°C) [ASTM: 25 ±1.0 °C]?
5.	ASTM: Material kept covered of	during conditioning to avoid evaporation?	?
6.		r approximately 2 minute period [AMRL: ±	
		1.11 g/L) for anionic (Rapid Set) emulsion	
		g/L) for anionic (Medium Set or Mixing –	
		sulfosuccinate solution (8g/L) for cationic I	
		and for rapid-setting and medium-setting e	emulsions, no provision is given
-	for slow-setting type en		.0
7.		nuously and vigorously during addition of r	
8.		peaker?	
9.		es after addition of reagent?	
10.		h?	
11.		cloth with distilled water?	
12.		, and wire cloth rinsed until water runs clea	
13.		ced in beaker with rod?	
14.		3°C) oven?	
		IO: Pretiminary neating at lower temperature [ASTM: Pre-drying sample is not acceptable],	
15.		until change is less than 0.1 g?	
16.		$_{\rm r}/{\rm M}_{\rm dir}$) x 100?	
10.	Demaisionity calculated as (Mide	m114 A 100:	
w	here: M_{der} = average weight of	of demulsibility residues from the 3 tests [A	ASTM: weight from single test]

 M_{dir} = weight of residue by distillation in 100 g of sample

DETERMINING DENSITY OF EMULSIFIED ASPHALTS (Weight Per Gallon)

(159)	
(D6937)	

		<u>APPARATUS</u>	Date:		
1.					
	` '		······		
_			······		
2.			·····		
3.	Constant temperature water bath at 25	± 0.5°C?	······		
		DDOCEDUDE			
		<u>PROCEDURE</u>			
1.	Emulsions with a viscosity requiremen vented to relieve pressure, then stirred		ater bath or oven with the sample		
2.	Emulsions with a viscosity requiremen				
	Note: Emulsions specified at 25°C ma				
	to the material.				
3.	Emulsion stirred and placed in a 25 ± 0.5 °C water bath for approximately 1 hour?				
4.			······		
5.			e sample?		
6.			or film?		
7.					
8.	Cap placed tightly onto measure and ex				
	with a clean, dry rag or paper?		* <u> </u>		
9.					
10.	Density of emulsion calculated as follo	ows:			
	AACLITO	MILDE	1 1		
	W = (G) (11.98);	Materials Refer	of emulsion in the measure		
V	where: W is the unit density (g/L) of	the emulsion and G is the mass (g)	of emulsion in the measure		
		is exist for different units, please ve	rify with the laboratory if they are using a		
	different method.				

RESIDUE BY DISTILLATION

(T59)	_
(D6997)	

	<u>APPARATUS</u>	Date:
Still a	and Burner Assembly:	
(a)	Still:	
(4)	(1) AASHTO: Made of aluminum alloy?	
	(2) ASTM: Made of aluminum alloy or iron?	
	(3) Approximately 240 mm (9 1/2 in.) by 95 mm (3 3/4 in.) [AMRL: \pm	
(b)	Still head:	s many miner diameter
(0)	(1) AASHTO: Made of aluminum alloy?	
	(2) ASTM: Made of aluminum alloy or iron?	
	(3) One 1 inch hole for connecting tube?	
	(4) Two ½ in. holes for thermometers?	
(c)	Clamp for still head?	
(d)	Seal for still, either of the following:	•••••
(u)	Note for ASTM: this is only required if further analysis of the oil and water	distillate is required
	(1) Joint ground to a tight fit?	
	(2) Gasket made of asbestos or oiled paper [ASTM: any material prov.	
	maximum temperature reached during distillation]?	
(a)	Burner for still:	•••••
(e)		
	(1) Approximately 4 ¾ in. inner diameter ring burner?	
	(2) Forts on finite periphery:	•••••
Conne	ecting Apparatus:	
(a)	Glass connecting tube, approximately 12 mm outer diameter:	
(b)	Bunsen burner for connecting tube (wing tip optional) [ASTM: entire burner	
(c)	Metal flame shield?	
(d)	Suitable adapter between condenser and graduate?	
Conde		
(a)	One of the following: (1) West or Liebig type glass condenser?	Laborator
	(1) West or Liebig type glass condenser?	
	C .1 C	
	(2) Metal-jacketed condenser?	
	(2) Metal-jacketed condenser?	
(b)	(2) Metal-jacketed condenser?	
, ,	(2) Metal-jacketed condenser?	
Receiv	(2) Metal-jacketed condenser?	
Receiv (a)	(2) Metal-jacketed condenser?	
Receiv	(2) Metal-jacketed condenser?	
Receiv (a) (b)	(2) Metal-jacketed condenser? (3) ASTM: Other condensers with wetted length of 400 to 550 mm? Adapter end to accommodate cork connection?	
Receiv (a) (b)	(2) Metal-jacketed condenser? (3) ASTM: Other condensers with wetted length of 400 to 550 mm? Adapter end to accommodate cork connection?	
Receive (a) (b)	(2) Metal-jacketed condenser? (3) ASTM: Other condensers with wetted length of 400 to 550 mm? Adapter end to accommodate cork connection?	
Receive (a) (b) Therm (a)	(2) Metal-jacketed condenser? (3) ASTM: Other condensers with wetted length of 400 to 550 mm? Adapter end to accommodate cork connection?	l accuracy?
Receiv (a) (b) Therm (a) (b) (c)	(2) Metal-jacketed condenser? (3) ASTM: Other condensers with wetted length of 400 to 550 mm? Adapter end to accommodate cork connection?	l accuracy?
Receive (a) (b) Therm (a) (b) (c) Misce	(2) Metal-jacketed condenser? (3) ASTM: Other condensers with wetted length of 400 to 550 mm? Adapter end to accommodate cork connection?	l accuracy?
Receiv (a) (b) Therm (a) (b) (c) Misce (a)	(2) Metal-jacketed condenser? (3) ASTM: Other condensers with wetted length of 400 to 550 mm? Adapter end to accommodate cork connection?	l accuracy?
Receiv (a) (b) Therm (a) (b) (c) Misce (a) (b)	(2) Metal-jacketed condenser? (3) ASTM: Other condensers with wetted length of 400 to 550 mm? Adapter end to accommodate cork connection?	l accuracy?
Receiv (a) (b) Therm (a) (b) (c) Misce (a)	(2) Metal-jacketed condenser? (3) ASTM: Other condensers with wetted length of 400 to 550 mm? Adapter end to accommodate cork connection?	l accuracy?
Receive (a) (b) Therm (a) (b) (c) Misce (a) (b) (c)	(2) Metal-jacketed condenser? (3) ASTM: Other condensers with wetted length of 400 to 550 mm? Adapter end to accommodate cork connection?	l accuracy?
Receiv (a) (b) Therm (a) (b) (c) Misce (a) (b)	(2) Metal-jacketed condenser? (3) ASTM: Other condensers with wetted length of 400 to 550 mm? Adapter end to accommodate cork connection?	l accuracy?

RESIDUE BY DISTILLATION

(139)
(D6997)

	PROCEDURE Date:
1.	ASTM: Sample conditioned at the viscosity testing temperature \pm 3°C unless freshly obtained from a
	storage tank? Samples from a storage tank are exempt from the temperature conditioning requirement
2.	ASTM: Samples stirred to ensure homogeneity prior to testing?
3.	Still with lid; clamp; thermometers; and gasket (if used) weighed to 0.1 g?
4.	200.0 ± 0.1 g [ASTM: ± 1.0 g] representative sample weighed into still assembly?
5.	One thermometer positioned approximately 1/4 in. from bottom of still?
6.	Other thermometer positioned approx. 6 1/2 in. [AMRL: thermometer not immersed] from bottom?
7.	ASTM: (second thermometer is not required) Is hole in still head sealed if no second thermometer used?
8.	Ring burner placed around still about 6 in. from bottom?
	Note: the positioning of the ring burner is flexible depending on the behavior of the material. Starting the
	burner higher or lower or lowering the burner progressively during testing are both acceptable provided that
	the other requirements of the test are met.
9.	Ring burner lit: Time: ()
10.	Connecting tube heated by Bunsen burner to prevent condensation [★ASTM: optional]?
11.	Ring burner moved to bottom of still when lower thermometer is approximately 420°F (215°C)
12.	Temperature of lower thermometer increased to $500 \pm 10^{\circ} F (260 \pm 5^{\circ} C)$?
13.	Temperature maintained at $500 \pm 10^{\circ}$ F ($260 \pm 5^{\circ}$ C) for 15 minutes?
14.	Ring burner shut off: Time: ()
15.	Elapsed time from the application of the first heat to shutting off the ring burner is 60 ± 15 minutes?
16.	Hot still assembly containing residue immediately weighed to 0.1 g?
17.	Residue stirred [ASTM: or agitated by swirling]?
18.	Suitable portions immediately poured into an 8 oz. tin or suitable molds using a 300-µm (No. 50) sieve if foreign matter is present?
19.	Volume of oil distillate, if present, recorded to nearest 1/2-mL?
20.	Volume percentage of oil distillate calculated & reported?
21.	Thermal buoyancy correction of 1.5 g added to gross still weight?
22.	Percentage residue by distillation calculated and reported to the nearest 0.1%?
•	AASH I O Materials Reference Laboratory
~~	

EVALUATING AGGREGATE COATING USING EMULSIFIED ASPHALTS (T59)(D6998) _____ **APPARATUS** Date: 1. Sieves: 3/4 in. (19.0-mm)____ (a) 1/4 in. (6.3-mm).....______ (b) Steel spatula, or equivalent, having a blade approximately 200-mm in length?..... 2. 3. 4. Supply of washed and dried reference stone?.... All passing a 3/4 in. (19.0-mm) sieve and not more than 5% passing the 1/4 in. (6.3-mm) sieve?..... (a) Class G2 [ASTM: GP2] balance with a capacity of at least 1000 g?..... 5. **PROCEDURE** 1. Emulsion conditioned in the original container [ASTM: in a sample container] to the same temperature required for viscosity testing ± 3°C?....._____ 2. 35.0 ± 0.1 g of emulsion added to the stone in the pan and mixed vigorously with spatula for 3 minutes?...... 3. Note: ASTM allows an equivalent ratio of 93% stone to 7% emulsified asphalt to be used. Larger or smaller total masses are acceptable. 4. Reported Information: Any appreciable separation of asphaltic base from the water of the emulsified asphalt?..... (a) Is stone thoroughly coated with the emulsified asphalt?.....______ (b) **COMMENTS:**

AASHTO Materials Reference Laboratory

	PARTICLE CHARGE (T59) (D7402)
	APPARATUS Date:
1. 2. 3. 4.	12-Volt dc current source, milliammeter, and a variable resistor? Electrodes: two 1 in. x 4 in. stainless steel plates? (a) Held rigidly parallel, 1/2 in. apart? 250 mL capacity beaker? Insulator (a) polytetraflouroethylene resin square rod? (1) virgin electrical grade?
5. 6. 7. 8.	(2) 1/2 in. [ASTM: 12.5 ± 0.5 mm] thick?
	PROCEDURE (ASTM: Method A)
1.	AASHTO: Emulsion heated to 122 ± 5 F (50 ± 3 C) in a 160 ± 5 F (71 ± 3 C) water bath, while stirring thoroughly?
 3. 	ASTM: Emulsion either stirred at 25 ± 3 °C or heated to 50 ± 3 °C in a water bath or oven and stirred, in the original container, based on the temperature used for viscosity testing? Emulsion poured into 250 mL beaker inserting glass rod between the two electrodes
4.	with ends of glass rod on the two opposite top edges of beaker?
4 . 5.	Clean dry electrodes connected to current source and inserted approximately 1 in. [AMRL: 3/4 to 1 1/2] into
6.	the emulsion?
7.	After 30 minutes or at 2 mA, whichever occurs first, electrodes disconnected
8.	and gently washed with a smooth, thin stream of distilled water?
	ELECTRODE CLEANING PROCEDURE
1.	New electrodes and electrodes being reused cleaned in the following sequence? (a) Wash with distilled water

Note to Assessors: Deviations from the specified electrode cleaning procedure will be marked as

COMMENTS:

permanent observations.

SAYBOLT FUROL VISCOSITY

(T59, T72)
(D7496, D88)

<u>APPAR</u>	<u>ATUS</u>			Date:		
VISCOSITY TUBES	1	2	3	4	5	6
Correction 1% or less (0.990 to 1.010) for referee testing?						
Month and year of calibration reported (3 year interval)						
Tube has furol tip?						
Tube corrosion resistant?						
Inner surface smooth & clean?						
Good cork?						
Bottom of tube 10 – 13 cm from flasks graduated mark?						

No.	Manufacturer	Timers Electric Spring		Quartz	0.1 sec. graduations?	Accurate to 0.1% in 60 min.? (3.6 seconds in 60 min.)
1						
2						

1.	Testi	ng Bath
	(a)	Viscometer and bath in draft-free location?
	(b)	Bath must be capable of being filled to at least 6 mm (1/4 in.) above overflow rim of viscometer?
	(c)	Bath has stirrer?
	(d)	Rath has thermosteric temperature control?
	1	(1) Control regulating the temperature of the bath so that it
		does not fluctuate by more than ± 0.05 °F (± 0.03 °C)?
2.	Ther	mometers
	(a)	ASTM 17F or 17C for tests at 77°F (25°C)?
	(b)	ASTM 19F or 19C for tests at 122°F (50°C)?
	(c)	ASTM: Or any other device of equal accuracy?
	(0)	
3.	Wate	<u>r Baths</u>
	(a)	Temperature within range of 77.0 \pm 0.2°F (25.00 \pm 0.10°C) for 25°C testing?
	(b)	AASHTO and ASTM A: Temperature within range of 160 ± 5°F (71 ± 3°C) for 50°C testing?
	(c)	ASTM B: Temperature within range of 51.4 ± 0.3 °C (124.5 ± 0.5 °F)
4.	Misco	ellaneous
₹.	(a)	Withdrawal device?
	(b)	Thermometer support?
	(-)	
	(c)	Proper receiving flasks?
	(d)	850-μm (No. 20) sieve or a 20-mesh strainer of wire cloth (framed or unframed)?
	(e)	4 oz. bottle with stopper for tests at 77°F (25°C)?
	(f)	400 mL heaker for tests at 122 H $(50%)$

	SAYBOLT FUROL VISCOSITY	(T59, T72)
	PROCEDURE FOR TESTS AT 77 °F (25 °C)	Date:
1.	Viscometer bath thermostat adjusted to maintain the bath at a temperature of 77.0	
2.	Sample thoroughly stirred, without incorporating bubbles?	
3.	100 to 110 mL of sample poured into 4 oz. (118 mL) bottle?	
<i>4</i> .	Closed bottle placed for 30 minutes in a water bath maintained at 77.0 \pm 0.2 °F (25)	
5.	Bottle slowly inverted several times to mix sample?	
6.	Sample poured into viscometer through No. 20 (850-µm) sieve?	
	(a) Small portion allowed to flow through outlet to waste?	<u></u>
	(b) Tube corked and viscometer filled until liquid begins to overflow the overf	
<i>7</i> .	Viscosity determined without any further disturbance of sample (without clearing g	allery or stirring)?
	(a) Cork snapped from tube and timer started at same instant?	······
	(b) Flask located so stream just touches neck of flask?	
	(c) Timer stopped when bottom of meniscus reaches grad. mark?	<u>-</u>
8.	Time of flow exceeds at least 20 seconds?	
9.	Calculation: Viscosity = Efflux Time X Calibration Factor of Viscometer Used	
	PROCEDURE FOR TESTS AT 122 °F (50 °C)	
1.	Emulsion sample heated in the original container to 122 \pm 5 F (50 \pm 3 °C) in	
	a 160 \pm 5 °F (71 \pm 3 °C) water bath or oven?	
2.	Sample thoroughly stirred without incorporating bubbles?	
3.	Approximately 100 mL of sample poured into 400 mL beaker?	
4.	Beaker immersed approximately 2 inches in water bath at 160 \pm 5 °F (71 \pm 3 °C) an	d sample heated until
	temperature is usable for the test but not greater than 53%)?	·····
5.	Cleaned and dried viscometer pre-corked?	·····
6.	Sample poured into viscometer through No. 20 (850-µm) sieve above overflow rim	of viscometer?
	(a) Sample stirred at approximately 60 rpm with thermometer and avoiding a	ir bubbles?
	(b) Temperature of emulsion adjusted until it remains constant for one	
	minute at $122.0 \pm 0.1 \text{F} (50.00 \pm 0.05 \text{C})$?	
	minute at 122.0 ± 0.1 °F $(50.00 \pm 0.05$ °C)?	erature at start of test?
7.	Viscosity determined after thermometer withdrawn from viscometer tube?	
	(a) Excess emulsion quickly removed from the gallery?	
	(b) Cork snapped from tube and timer started at same instant?	

Flask located so stream just touches neck of flask?.....

Timer stopped when bottom of meniscus reaches grad. mark?.....

COMMENTS:

8.

(c)

		SAYBOLT FUROL VISCOSITY	(D7496, D88) _
		PROCEDURE FOR TESTS AT 77 °F (25 °C)	Date:
Visco	meter cle	an, dry, and stoppered or corked?	······
Mater	rial prepa	red according to either Procedure A or Procedure B?	
(a)		dure A (using an open beaker)	
	(1)	Approximately 100 mL of emulsion poured into a 400 mL glass bed	aker?
	(2)	Bottom of the beaker immersed 50 mm below the level of a 25°C (7	
	(3)	Beaker held upright and stirred with a thermometer at approximate	
	(4)	Incorporation of air bubbles avoided?	
(b)	Proce	dure B (using a sealed bottle)	
	(1)	Sample poured into an approximately 120 mL bottle?	
	(2)	Sealed bottle placed into a water bath maintained at 25°C (77°F) for	
	(3)	Bottle removed from the bath and mixed by inverting the bottle sev	
	(4)	Bubble formation avoided?	
Samn	\ /	l into the viscometer through an 850-µm sieve or 20-mesh strainer?	
		ne overflow rim?	
Fmul	sified asr	ohalt stirred with the thermometer?	
		ion avoided?	
		th temperature adjusted until the emulsion temperature remains const	
		2°F)?	
		withdrawn and excess asphalt removed from the gallery by suction?	
		from the tube and timer started at the same instant?	
		so that the stream just touches the neck of the flask?	
		when the bottom of the meniscus touches the graduation mark?	
1 ime	oj jiow e.	xceeds 20 seconds?	
	6	DDOGEDINE FOR TEGTS AT 122 OF (50 OC)	
		PROCEDURE FOR TESTS AT 122 °F (50 °C)	
Visco	meter cle	an, dry, and stoppered or corked?	
		red according to either Procedure A or Procedure B?	
(a) 1			
	(1)	dure A If the sample is cooler than $50^{\circ}C$ (122°F), heated to $50 \pm 3^{\circ}C$ (122 (160 ± 5°F) water bath or oven?	
	(2)	Sample stirred without incorporating bubbles?	
	(3)	Approximately 100 mL poured into a 400 mL glass beaker?	
	(4)	Bottom of the beaker immersed approximately 50 mm (2 in.) below	
	()	$(160 \pm 5^{\circ}F)$ water bath?	
	(5)	Beaker held upright and stirred with the thermometer at 60 revolut	ions/minute?
	(6)	Sample heated to $51.4 \pm 0.3^{\circ}C$ ($124.5 \pm 0.5^{\circ}F$)?	
(b)	` /	dure B	
(0)	(1)	Sample poured into an approximately 120 mL bottle and sealed?	
	(2)	Sealed bottle placed in a water bath at 51.4 ± 0.3 °C (124.5 ± 0.5 °F)	
	(3)	Sample mixed by slowly inverting the bottle several times, avoiding	
C			
		l into the viscometer through an 850-µm sieve or 20-mesh strainer?	
		ne overflow rim?	
		ed at approximately 60 revolutions/min. until the sample reached 50 ±	
		withdrawn and excess emulsion removed from the gallery by suction?.	
		from the tube and timer started at the same instant?	
Flask	located s	so that the stream just touches the neck of the flask?	······ _

Time stopped when the bottom of the meniscus touches the graduation mark?.....

Time of flow exceeds 20 seconds?....._______

COMMENTS:

9.

10.

SWEEP TEST OF BITUMINOUS EMULSION SURFACE TREATMENT SAMPLES

(D7000)	
(\mathbf{D}_{I})	

	<u>APPARATUS</u>	Date:	
Mixer, Capable of abrasion at a rate of 0.83 gyrations/sec (50 gyr/min) [AMRL: ± 2 gyr/min]?			
Quick-0 1. 2.	clamp Mounting Base: Level support for clamping sample in place? Does not move during abrasion test?		
Pan, Ca	apable of holding sample on mixer and dislodged aggregate?		
Oven: 1. 2. 3.	Constant temperature, force draft?	l floor?	
Balance 1. 2.	e: Must be capable of weighing 800g or more to within ± 0.1g? Has a platform length and width of 240 mm (9.5 in.)?		
Brush F 1. 2.	Holder: Removable? Free floating movement of 19 ± 1 mm? (a) Brush head length 128mm? (b) Overall trim 25.4 mm [AMRL: ± 2 mm]? Total weight of brush head and attached weight shall be 1500 ± 15 g (Note: The collar and nylon strip are not included in the weight)		
Nylon S 1.	Strip Brush: Shall conform to following specifications?	han brush is in poor condition.] (Note: check fiber diameter w/ calipers)?	
Strike-0 1. 2.	Flat, stainless steel metal plate?		
<u>Strike-o</u> 1. 2.	off Rod: Made of 17 to 20 mm diameter electrical conduit [AMRL: metal or] Length of 750 ± 100 mm?		
Sweep '1. 2.	Test Compactor: Suitable design? Mass of 7500 ± 500g?		
COMM	MENTS (D7000):	(D7000)	

SWEEP TEST OF BITUMINOUS EMULSION SURFACE TREATMENT SAMPLES

(D7000)	
(D7000)	

SAMPLE PREPARATION	Date:
--------------------	-------

Emulsion:

Aggregate:

- 1. Representative sample obtained?
- 2. Dried to constant weight (sample should not be washed) [AMRL: at $110 \pm 5^{\circ}$ C]?.....
- 4. The amount of aggregate to be used calculated from the following equation:

Note: A pre-calculated table with known masses and corresponding bulk specific gravities may be used to interpolate the correct weight.

$$Y = \frac{A(202.1X - 15.8)}{100} + \frac{B(146.4X - 4.7)}{100}$$

Where:

A = % of aggregate from 9.5 to 6.3 mm

B = % of aggregate from 6.3 to 4.75 mm

X = bulk specific gravity

Y = amount of aggregate needed for test, g

Asphalt felt disks:

COMMENTS (D7000):

1.	Disks must be made of 30 lb asphalt felt paper and free of breaks, cracks, and tears?	
	(a) Cut into 300 ± 10 mm diameter disks?	
	(b) Placed in a 50°C oven for 24 to 72 hours to flatten?	
	(c) Stored flat and at room temperature for a minimum of 3 days?	
Test Spe	AASHTO Materials Reference Laboratory	
1.	Weigh the asphalt felt disk to the nearest 0.1g?	
2.	Felt disk placed on a flat surface and manipulated until flat?	
3.	Disk not used if edges curl, bubble, or the disk contained foreign matter?	
4.	Aggregate weighed and the mass recorded to the nearest 1 g?	
5.	Strike off template placed over felt disk?	
6.	83 ± 5 g of emulsion at 60°C poured along top arc of the disk (Assessor: start timer, see Step 12)?	
7.	Excess emulsion removed with strike-off rod using a gentle side-to-side motion?	
8.	Excess emulsion struck off within a 3 ± 1 s [AMRL: < 10 s finding is an <i>observation</i>] period without stopping motion?	
9.	Pre-weighted aggregate (See table 1) immediately applied using a back and forth motion?	
10.	Aggregate compacted using three -½ cycles in one direction and three -½ cycles in the perpendicular direction?	
11.	Sample weighed (Assessor: stop timer) and placed in oven?	
12.	Time from beginning of sample production to weighing not longer than 4 minutes?	

(D7000)

SWEEP TEST OF BITUMINOUS EMULSION SURFACE TREATMENT SAMPLES

(D7000)	
(D 1000)	

	SAMPLE PREPARATION (Continued) Date:
Conditi	oning:
1.	Conditioning based on road parameters and field performance?
1.	Note: Conditioning times may vary greatly pending on emulsion type.
2.	Time and temperature shall be within 10% of desired value and relative humidity within 25%?
3.	Sample removed from the oven (Assessor: start timer, see Procedure Step 1)?
4.	After conditioning sample turned vertically and loose aggregate removed by gentle brushing with fingers?
5.	Specimen weighed to the nearest 0.1 g and recorded?
٥.	
	<u>PROCEDURE</u>
1. 2. 3. 4. 5. 6. 7.	The time from removing sample from oven to being placed in test apparatus not longer than 2 minutes?
	CALCULATION Loss = (A-B)/(A-C) x 100 x 1.33 A = initial weight B = final weight, and C = asphalt sample disk weight