



# Standard Test Method for Centrifuge Kerosine Equivalent<sup>1</sup>

This standard is issued under the fixed designation D 5148; the number immediately following the designation indicates the year of original adoption or, in the case of revision, the year of last revision. A number in parentheses indicates the year of last reappraisal. A superscript epsilon ( $\epsilon$ ) indicates an editorial change since the last revision or reappraisal.

## 1. Scope

1.1 This test method determines the centrifuge kerosine equivalent (CKE) of aggregate used in bituminous mixtures.

### 1.2 Units of Measure:

1.2.1 With regard to sieve sizes and size of aggregate as determined by the use of testing sieves, the values in inch-pound units are shown for the convenience of the user, but the standard sieve designation shown in parentheses is the standard value as stated in Specification E 11.

1.2.2 With regard to other units of measure, the values shown in parentheses are for information purposes.

1.3 *This standard does not purport to address all of the safety concerns, if any, associated with its use. It is the responsibility of the user of this standard to establish appropriate safety and health practices and determine the applicability of regulatory limitations prior to use.* For specific hazard statements, see 7.1.

## 2. Referenced Documents

### 2.1 ASTM Standards:

C 127 Test Method for Specific Gravity and Absorption of Coarse Aggregate<sup>2</sup>

C 128 Test Method for Specific Gravity and Absorption of Fine Aggregate<sup>2</sup>

C 702 Practice for Reducing Samples of Aggregate to Testing Size<sup>2</sup>

D 75 Practice for Sampling Aggregates<sup>3</sup>

D 4753 Specification for Evaluating, Selecting, and Specifying Balances and Scales for Use in Testing Soil, Rock, and Related Construction Materials<sup>4</sup>

E 11 Specification for Wire-Cloth Sieves for Testing Purposes<sup>5</sup>

E 832 Specification for Laboratory Filter Papers<sup>5</sup>

## 3. Terminology

### 3.1 Symbols: Symbols:

3.1.1  $C$ —coarse aggregate fraction, that portion of the

sample which passes the  $\frac{3}{8}$ -in. (9.5-mm) sieve and is retained on the No. 4 (4.75-mm) sieve.

3.1.2  $F$ —fine aggregate fraction, that portion of the sample which passes the No. 4 (4.75-mm) sieve.

3.1.3  $SA$ —surface area. The sum,  $m^2/kg$  ( $ft^2/lb$ ), obtained by adding the products of the percent passing each sieve and its corresponding factor, (see 11.1) and dividing by 100.

3.1.4  $K$  factors—values determined as described in 3.1.5 through 3.1.8 and identified as  $K_c$ ,  $K_f$ , or  $K_m$ .

3.1.5  $K_c$ —determined from the percent of SAE No. 10 oil retained, which represents the total effect of the aggregate's absorptive properties and surface roughness of the aggregates coarse fraction.

NOTE 1—Based on comparative testing in California, the same results can be obtained substituting Shell Tellus No. 100 oil for SAE No. 10 oil.

3.1.6  $K_f$ —determined from the following factors:

3.1.7 Percent of kerosine retained, which represents the total effect of superficial area, the aggregate's absorptive properties and surface roughness of the aggregate's fine fraction.

3.1.7.1 Computed surface area, based on particle size.

3.1.7.2 Percent of aggregate passing No. 4 (4.75-mm) sieve.

3.1.8  $K_m$ —the "mean" or composite value of  $K$  for a given combination of coarse and fine materials on which  $K_c$  and  $K_f$  have already been determined independently.

## 4. Significance and Use

4.1 The CKE furnishes an index, designated as the  $K$  factor, that indicates the aggregate particle roughness and surface capacity based on porosity.

4.2 The CKE is used as part of the Hveem mix design procedure to determine the approximate bitumen ratio (ABR), as shown in Appendix X1. However, there are other applications such as determining the coarse aggregate fraction constant ( $K_c$  for use as an aid in selecting a bitumen content for open-graded friction courses.

## 5. Apparatus

5.1 *Centrifuge*, power driven, capable of exerting a force of  $400 \pm 8$  times gravity (400 G) on a 100-g sample.

The required  $r/min$  ( $\pm 10$ ) of the centrifuge head =

$$\sqrt{(25.4(14\,000\,000/r))}$$

where  $r$  = radius to center of gravity of sample, mm.

5.2 *Centrifuge Cups*,  $71.4 \pm 1.6$  mm ( $2\frac{13}{16} \pm \frac{1}{16}$  in.) in height and  $52.4 \pm 1.6$  mm ( $2\frac{1}{16} \pm \frac{1}{16}$  in.) inside diameter (see

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<sup>2</sup> Annual Book of ASTM Standards, Vol 04.02.

<sup>3</sup> Annual Book of ASTM Standards, Vol 04.03.

<sup>4</sup> Annual Book of ASTM Standards, Vol 04.08.

<sup>5</sup> Annual Book of ASTM Standards, Vol 14.02.

Fig. 1) complete with perforated brass plate  $0.787 \pm 0.03$  mm ( $0.031 \pm 0.001$  in.) thick with a minimum of 15 holes,  $1.575$  mm  $\pm 0.03$  mm ( $0.062 \pm 0.001$  in.) in diameter, per square centimetre (100 holes/in.<sup>2</sup>).

5.3 *Balance*—A balance having a minimum capacity of 500 g and meeting the requirements of Specification D 4753, Class GP2.

5.4 *Metal Funnel*, top diameter  $98.4 \pm 1.6$  mm ( $3\frac{7}{8} \pm \frac{1}{16}$  in.), height  $109.5 \pm 1.6$  mm ( $4\frac{5}{16} \pm \frac{1}{16}$  in.), orifice  $12.7 \pm 1.6$  mm ( $\frac{1}{2} \pm \frac{1}{16}$  in.), with a piece of No. 10 (2.0-mm) sieve soldered slightly above the orifice (Fig. 2).

5.5 *Tin Pan*, round,  $114.3 \pm 1.6$  mm ( $4\frac{1}{2} \pm \frac{1}{16}$ -in.) diameter,  $25.4 \pm 1.6$  mm ( $1 \pm \frac{1}{16}$  in.) deep.

**6. Materials**

6.1 *Kerosine*.

6.2 *Lubricating Oil*, SAE No. 10 (see Note 1).

6.3 *Filter Paper*, size 5½-cm diameter, Type 1, Class B.

NOTE 2—VWR Guide No. 613 satisfies ASTM grade Type 1, Class B, Specification E 832.

**7. Hazards**

7.1 **Warning**—Kerosine is flammable, and therefore caution should be used in storage and use.

**8. Sampling**

8.1 Sampling is done in accordance with Practice D 75.

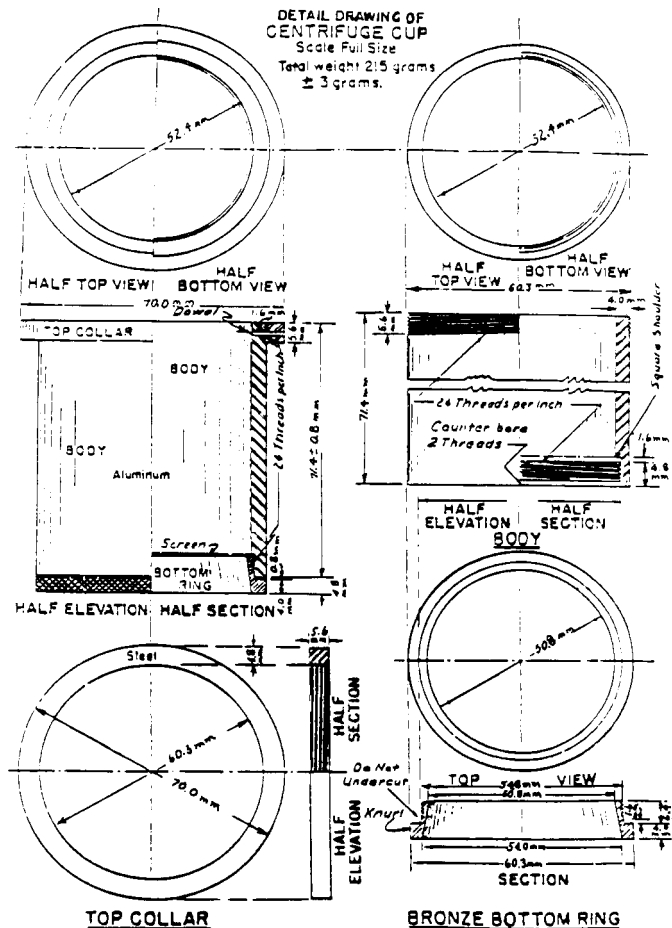


FIG. 1 Detailed Drawing of a Centrifuge Cup

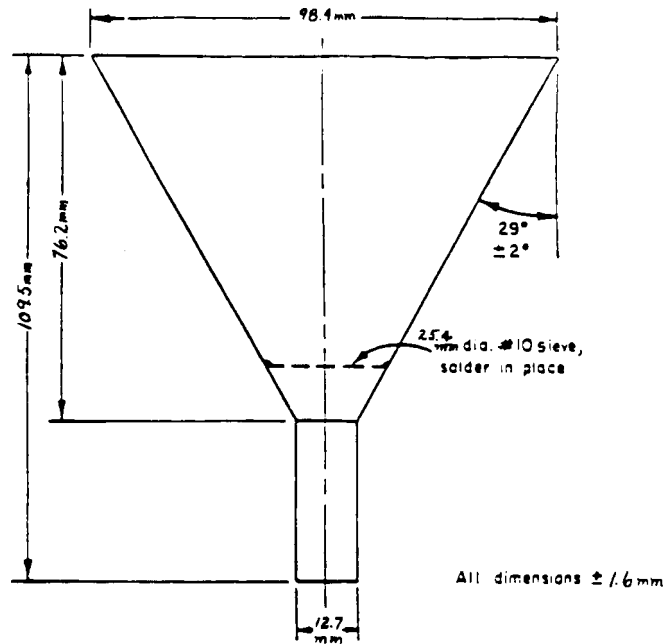


FIG. 2 Detailed Drawing for Metal Funnel

8.2 Reduce the sample in accordance with Test Method C 702.

**9. Preparation of Sample**

9.1 Determine the bulk specific gravity of the coarse aggregate (4.1) and apparent specific gravity of the fine aggregate (4.2), using Test Methods C 127 and C 128, respectively.

NOTE 3—Apparent specific gravity is used for the fine aggregate because it is easier to determine than the bulk specific gravity, and its use does not affect the CKE results.

9.2 *Specific Gravity*—Calculate the average specific gravity for the aggregate based upon the design grading by the following formula:

$$G = \frac{1}{\frac{P_c}{100G_c} + \frac{P_f}{100G_f}} \quad (1)$$

where:

$G$  = average specific gravity,

$P_c$  = coarse aggregate present in the original sample, weight %,

$P_f$  = fine aggregate present in the original sample, weight %,

$G_c$  = bulk (oven dry) specific gravity of the coarse aggregate, and

$G_f$  = apparent specific gravity of the fine aggregate.

9.3 Separate the aggregate into two size groups, “C” material (used for  $K_c$  determinations) passing the ¾-in. (9.5-mm) sieve and retained on the No. 4 (4.75-mm) sieve, and “F” material (for  $K_f$  determination) all passing the No. 4 (4.75-mm) sieve.

**10. Procedures**

10.1 *Procedure for Fine F:*

10.1.1 Quarter or split out approximately 105 g for each sample, representative of the material passing No. 4 (4.75-mm) sieve.

10.1.2 Place on hot plate or in  $110 \pm 5^\circ\text{C}$  ( $230 \pm 9^\circ\text{F}$ ) oven and dry to constant weight.

10.1.3 Allow to cool.

10.1.4 Place  $100.0 \pm 0.1$  g in each of the tared centrifuge cups fitted with the perforated metal disk underlying a disk of filter paper.

10.1.5 Place centrifuge cups containing samples in pan with sufficient kerosine  $12.7 \pm 3.2$  mm ( $\frac{1}{2} \pm \frac{1}{8}$ in.) deep to saturate the sample. When specimens are thoroughly saturated (by capillary action), place the cups with samples in centrifuge. Samples should be tested in pairs, placed opposite of each other to avoid damage to the centrifuge.

10.1.6 Spin in centrifuge for 2 min at a force of 400 G.

10.1.7 Reweigh each cup, containing samples, to nearest 0.1 g and subtract original weight. The difference is the percent of kerosine retained (based on 100 g of dry aggregate). The percent of kerosine retained is the CKE value. Record the average of the two values for duplicate samples.

10.2 Procedure for Coarse C:

10.2.1 Quarter or split out approximately 105 g for each sample, representative of the material passing  $\frac{3}{8}$ -in. (9.5-mm) and retained on No. 4 (4.75-mm) sieve material.

10.2.2 Dry sample on hot plate or in  $110 \pm 5^\circ\text{C}$  ( $230 \pm 9^\circ\text{F}$ ) oven to constant weight and allow to cool to room temperature.

10.2.3 Weigh out  $100.0$  g  $\pm 0.1$  g and place in funnel (see 5.4).

10.2.4 Completely immerse specimen in SAE No. 10 lubricating oil for 5 min (see Note 1).

10.2.5 Place the funnel in a container, maintaining the axis in a vertical position and allow to drain for 2 min.

10.2.6 Place funnel containing sample in  $60^\circ\text{C}$  ( $140^\circ\text{F}$ ) oven for 15 min of additional draining, remembering to keep the funnel axis in a vertical position.

10.2.7 Pour sample from funnel into tared pan, cool to room temperature, and reweigh sample to nearest 0.1 g. Subtract original weight and record difference as percent of oil retained (based on 100 g of dry aggregate).

11. Determination of K Factors

11.1 Use the following surface area factors to calculate surface area based upon design grading as follows:

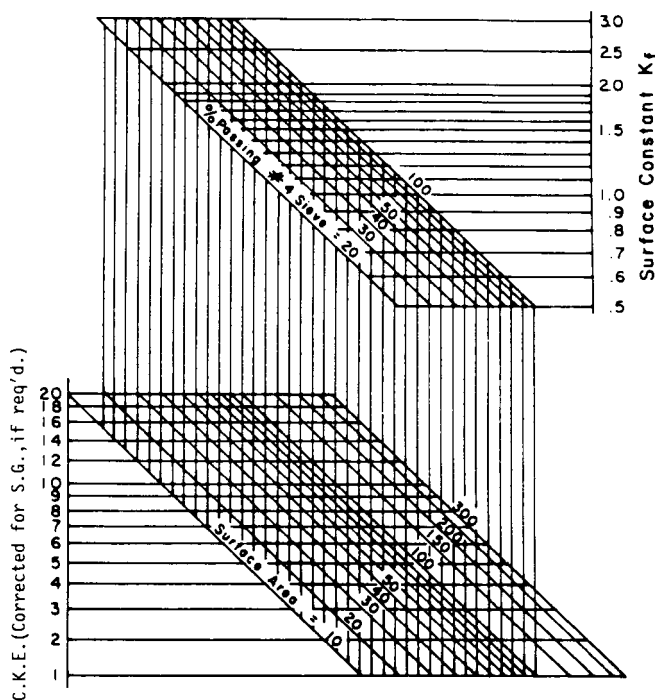
Sieve Size Passed	m <sup>2</sup> /Kg	ft <sup>2</sup> /lb
Maximum size	0.41	2
No. 4 (4.75 mm)	0.41	2
No. 8 (2.36 mm)	0.82	4
No. 16 (1.18 mm)	1.6	8
No. 30 (600 μm)	2.9	14
No. 50 (300 μm)	6.1	30
No. 100 (150 μm)	12.3	60
No. 200 (75 μm)	32.8	160

11.1.1 All surface area factors must be used in calculations; thus, if a sample passes No. 4 (4.75-mm) sieve 100 %, include in calculations  $100 \times 0.41$  m<sup>2</sup>/kg (2 ft<sup>2</sup>/lb), for passing maximum size as well as  $100 \times 0.41$  m<sup>2</sup>/kg (2 ft<sup>2</sup>/lb) for passing No. 4 (4.75 mm) sieve.

11.2 Use the chart shown in Fig. 3 for determination of K<sub>f</sub>.

11.2.1 If the apparent specific gravity for F is greater than 2.70 or less than 2.60, make correction to oil retained, using the following formula:

CHART FOR DETERMINING K<sub>f</sub> FROM C.K.E.



NOTE: C.K.E. Corrected for S.G. = % ker. ret. (C.K.E.)  $\times \frac{\text{app. S.G. (f)}}{2.65}$

FIG. 3 Chart for Determining K<sub>f</sub> from CKE

$$\begin{aligned} &\text{Percent of kerosine retained} \\ &\times (\text{apparent specific gravity } F/2.65) \\ &= \text{CKE corrected for specific gravity} \end{aligned} \tag{2}$$

11.2.2 Start in lower left hand corner of chart in Fig. 3 with value for CKE corrected for specific gravity, following straightedge horizontally to right to the intersection with calculated surface area, hold point, move vertically upward to the intersection with the percent passing the No. 4 (4.75-mm) sieve, hold point, and follow straightedge horizontally to right. The value obtained will be the surface constant for the passing No. 4 (4.75-mm) fraction F and is known as K<sub>f</sub>.

11.3 Use chart shown in Fig. 4 for determination of K<sub>c</sub>.

11.3.1 If the bulk (oven dry) specific gravity for C is greater than 2.70 or less than 2.60, apply correction to oil retained, using formula at top of chart in Fig. 4.

11.3.2 Start at the bottom of chart in Fig. 4 with the corrected percent of oil retained, follow straightedge vertically upward to intersection with the diagonal line, hold point, and follow the straightedge horizontally to the left. The value obtained will be the surface constant for the retained fraction C and is known as K<sub>c</sub>.

11.4 Use the chart shown in Fig. 5 to combine K<sub>f</sub> and K<sub>c</sub> for determination of K<sub>m</sub>.

$$K_m = K_f + \text{correction to } K_f \tag{3}$$

MATERIALS USED: aggregate passing 3/8in(7.5mm) sieve, ret. #4(4.75mm) sieve.  
Oil - SAE 10 or Shell Tallus # 100.

$$\% \text{ oil ret. corrected} = \% \text{ oil ret.} \times \frac{\text{bulk s.g. of aggregate}}{2.65}$$

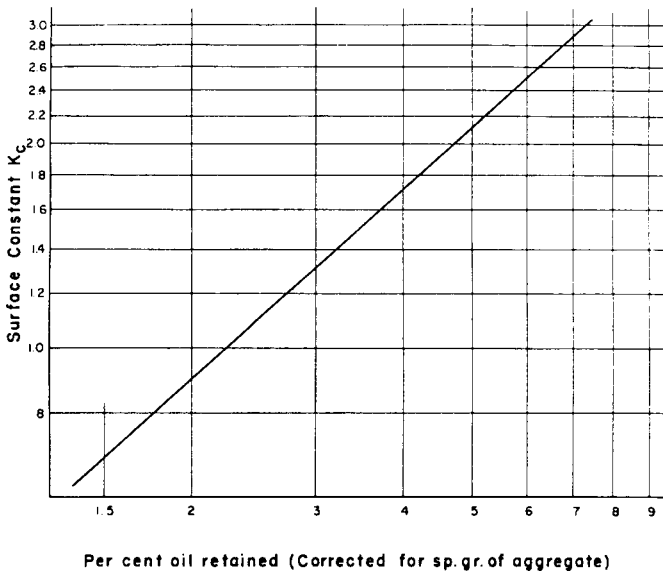


FIG. 4 Chart for Determining  $K_c$  from Coarse Aggregate Percent Oil Retained

11.4.1 The “correction to  $K_f$ ” value obtained from Fig. 5 is positive if  $(K_c - K_f)$  is positive and is negative if  $(K_c - K_f)$  is negative.

11.4.2 No correction needs to be applied for asphalt viscosity.

NOTE 4—When there is 20 % or less coarse material in a sample, the  $K_c$  is not used; therefore, the  $K_f$  and  $K_m$  are the same.

11.4.3 The determination of  $K_m$  is shown in the following example:

$$K_c = 1.0, K_f = 1.8, SA = 5.12 \text{ m}^2/\text{kg} \text{ (25 ft}^2/\text{lb), passing No. 4} = 60 \% \quad (4)$$

11.4.3.1 Using the chart in Fig. 5 start in lower left corner with  $SA = 5.12 \text{ m}^2/\text{kg} \text{ (25 ft}^2/\text{lb)}$ , follow straightedge horizontally to percent of coarse aggregate (40 %), hold point, follow straightedge vertically upward to intersection with the difference between  $K_c$  and  $K_f$  (0.8), hold point, and follow straightedge horizontally to right to a “correction to  $K_f$ .” In this example, the correction is 0.2. Because  $K_c - K_f$  (1.0 - 1.8) is negative, the correction is negative; therefore,  $K_m = 1.8 - 0.2 = 1.6$ . If  $K_c$  had been 1.8, and  $K_f$  1.0,  $K_c - K_f$  would have been positive (+0.8), and the correction (0.2) would have been positive. In this case,  $K_m$  would be  $1.0 + 0.2 = 1.2$ .

## 12. Report

12.1 Report percent kerosine retained, percent oil retained,  $K_f$ ,  $K_m$ , and  $K_c$ .

If  $(K_c - K_f)$  is neg., corr. is neg.  
If  $(K_c - K_f)$  is pos., corr. is pos.  
 $K_m = K_f + \text{corr. to } K_f$

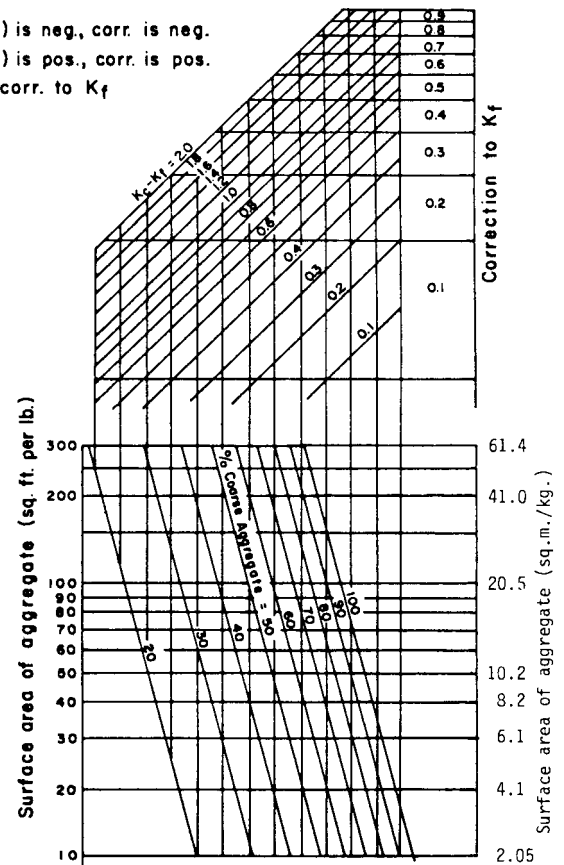


FIG. 5 Chart for Combining  $K_f$  and  $K_c$  to Determine  $K_m$

## 13. Precision and Bias

### 13.1 Precision:

13.1.1 Estimates of variations within a laboratory cannot be made with available data because in several cases the same operator did not conduct all the tests from which the data were generated. However, the following is an estimate of variation between laboratories based on a test result that is the average of two samples.

	Variation Between Laboratories	
	Standard Deviation	Acceptable Range of Two Results
Kerosine retained, %	0.34	0.962
Oil retained, %	0.416	1.206

13.1.2 The precision statement is based on an interlaboratory study of 19 laboratories that tested two aggregates twice with an interval of one week. The same operator conducted the first series of tests on both aggregates but did not necessarily conduct the second series of tests. More specifically, the results are based on using an SAE No. 10 oil.

13.2 *Bias*—The procedure in this test method has no bias because the values of the kerosine retained and the oil retained are defined in terms of this test method.

## 14. Keywords

14.1 aggregates; bitumin content; centrifuge kerosine equivalent; surface roughness

**APPENDIX****(Nonmandatory Information)****X1.****X1.1 Scope**

X1.1.1 The  $K_f$  and  $K_c$  constants for an aggregate are used in bituminous mix design procedures to determine an approximate bitumen ratio (ABR). When used in this manner in the Hveem mix design procedure for dense-graded bituminous

mixtures, other mix properties are also considered such as appearance (for flushing condition), voids, and Hveem stability and cohesion. The ABR calculated for open-graded friction courses from acceptable relations should also be verified by conducting an asphalt drainage test.

X1.1.2 The ABR for dense-graded bituminous mixtures is determined by use of Fig. X1.1.

X1.1.3 Fig. X1.2 is used for correcting the bitumen requirement for paving asphalts.

X1.1.4 The ABR for open-graded mixtures can be calculated from  $ABR = 2K_c + 4.0$  and correcting for aggregate specific gravity.<sup>6,7</sup>

<sup>6</sup> Federal Highway Administration, "Design of Open-Graded Asphalt Friction Courses," Report No. FHWA-RD-74-2, January 1974, Washington, DC, Suppl. No. 1, July 11, 1975.

<sup>7</sup> White, Thomas D., "Field Performance of Porous Friction Course," miscellaneous paper S-76-13, April 1976, US Army Engineer Waterways Experiment Station, CE, Vicksburg, MS; and Report No. FAA-RD-73-197, February 1975, Federal Aviation Administration, Washington, DC.

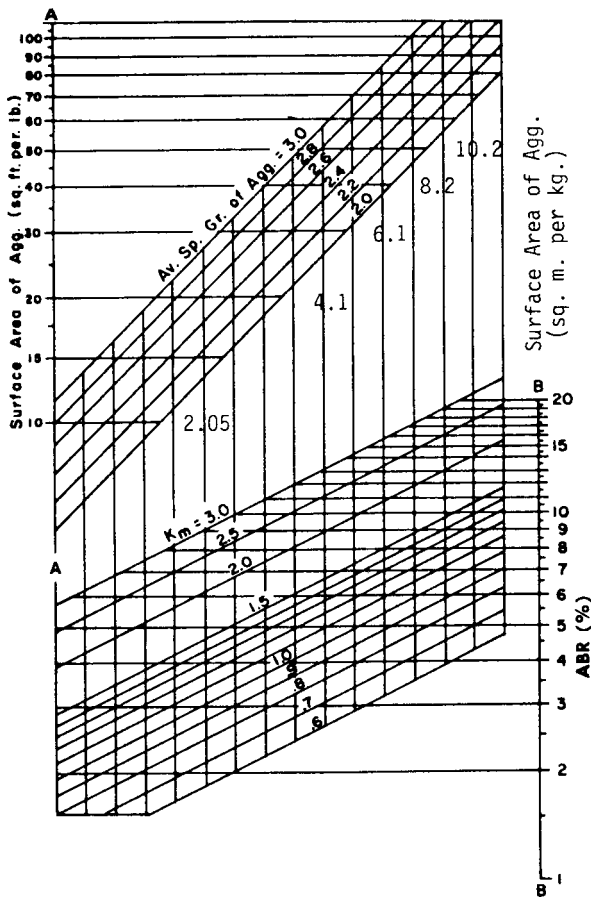


FIG. X1.1 Chart for Computing Approximate Bitumen Ratio (ABR) for Dense-Graded Bituminous Mixtures

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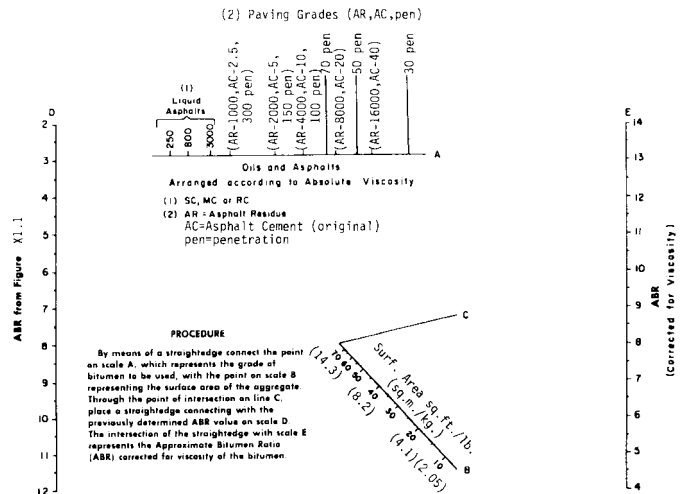


FIG. X1.2 Chart for Correcting ABR for Grade of Asphalt