



Designation: D 6144 – 9704

## Standard Test Method for Analysis of AMS ( $\alpha$ -Methylstyrene) by Capillary Gas Chromatography<sup>1</sup>

This standard is issued under the fixed designation D 6144; 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 reapproval. A superscript epsilon ( $\epsilon$ ) indicates an editorial change since the last revision or reapproval.

### 1. Scope\*

1.1 This test method covers the determination of the purity of AMS ( $\alpha$ -methylstyrene) by gas chromatography. Calibration of the gas chromatography system is done by the external standard calibration technique.

1.2 This test method has been found applicable to the measurement of impurities such as cumene, 3-methyl-2-cyclopentene-1-one, *n*-propylbenzene, *tert*-butylbenzene, *sec*-butylbenzene, *cis*-2-phenyl-2-butene, acetophenone, 1-phenyl-1-butene, 2-phenyl-2-propanol, and *trans*-2-phenyl-2-butene, *m*-cymene, *p*-cymene, and phenol, which are common to the manufacturing process of AMS. The method has also been found applicable for the determination of para-tertiary-butylcatechol typically added as a stabilizer to AMS. The impurities in AMS can be analyzed over a range of 5 to 800 mg/kg by this method. (See Table 1.) The limit of detection for these impurities is 0.01 wt %.

~~1.3 The following applies to all specified limits typically in this test method: for purposes the range of 5 to 10 mg/kg. (See Table 1.)~~

~~1.3 In determining the conformance with this of the test method, an observed value or a calculated value results using this method to applicable specifications, results shall be rounded off “to the nearest unit” in the last right-hand digit used in expressing the specification limit, in accordance with the rounding-off method of Practice E 29.~~

1.4 *This standard does not purport to address all 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 Section 8.

<sup>1</sup> This test method is under the jurisdiction of ASTM Committee D16 on Aromatic Hydrocarbons and Related Chemicals and is the direct responsibility of Subcommittee D16.07 on Styrene, Ethylbenzene, and C<sub>9</sub> and C<sub>10</sub> Aromatic Hydrocarbons.

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\*A Summary of Changes section appears at the end of this standard.

**2. Referenced Documents**

2.1 *ASTM Standards:* <sup>2</sup>

- D 3437 Practice for Sampling and Handling Liquid Cyclic Products
- D 4790 Terminology 4307 Practice for Preparation of Liquid Blends for Use as Analytical Standards
- D 4790 Terminology of Aromatic Hydrocarbons and Related Chemicals
- D 6809 Guide for Quality Control and Quality Assurance Procedures for Aromatic Hydrocarbons and Related Materials
- E 29 Practice for Using Significant Digits in Test Data to Determine Conformance with Specifications
- E 355 Practice for Gas Chromatography Terms and Relationships
- E 691510 Practice for Conducting an Interlaboratory Study to Determine the Precision of a Test Method
- E 1510 Practice for Installing Fused Silica Open Tubular Capillary Columns in Gas Chromatographs

<sup>2</sup> For referenced ASTM standards, visit the ASTM website, [www.astm.org](http://www.astm.org), or contact ASTM Customer Service at [service@astm.org](mailto:service@astm.org). For Annual Book of ASTM Standards, Vol 06.04, volume information, refer to the standard's Document Summary page on the ASTM website.

**TABLE 2.1 Relative Summative Response Factors (RRF) Based on Effective Carbon Numbers, Data (ECNmg/kg)**

Compound	ECN	MW	Calculated RRF vs n-octane
Compound	Repeatability	Reproducibility	Range Studied
n-octane	-8.0	114	1.000
Acetone	1.61 + 0.035*Acetone	0.47 + 1.10*Acetone	0.5 – 26
Cumene	-9.0	120	0.936
Cumene	9.0	7.88 + 0.19*Cumene	45 – 290
3-methyl-2-cyclopentene-1-one	-4.9	-96	1.375
n-Propylbenzene (NPB)	2.11 + 0.03*NPB	96	55 – 195
n-propylbenzene	1.84	3.65 + 0.58*Phenol	1 – 40
Phenol	1.84	3.65 + 0.58*Phenol	1 – 40
tert-Butylbenzene (TBB)	-9.0	120	3.936
tert-Butylbenzene (TBB)	9.0	3.63 + 0.087*TBB	150 – 650
tert-butylbenzene	1-0-0	134	0.940
sec-Butylbenzene (SBB)	4.23 + 0.019*SBB	21.60 + 0.25*SBB	200 – 765
sec-butylbenzene	1-0-0	134	0.940
m-Cymene	0.31 + 0.035*mCymene	2.34 + 0.35*mCymene	2 – 50
2-phenyl-2-butene (cis or trans)	-9.9	132	0.936
o-Cymene	1.63	8.00	29 – 31
Acetophenone	-7.0	120	1.203
p-Cymene	5.12	15.79	10 – 18
2-phenyl-1-butene	-9.9	132	0.936
cis-2-Phenyl-2-Butene (CPB)	0.17 + 0.030*CPB	5.39 + 0.11*CPB	50 – 225
AMS oxide	-8.0	134	1.175
trans-2-Phenyl-2-Butene (TPB)	1.57	0.54 + 0.20*TPB	19 – 70
AMS	-8.9	118	0.930
1-Phenyl-1-Butene (PB)	4.00 + 0.018*PB	0.17 + 0.19*PB	0.2 – 735
2-phen-2-propanol	-8.5	136	1.123
Acetophenone (AP)	8.5	1.17 + 0.63*AP	15 – 123

$$RRF_{(i)} = \frac{ECN_{(STD)}}{ECN_{(i)}} \times \frac{MW_{(i)}}{MW_{(STD)}}$$

where:

- 15
- $RRF_{para-tert-Butylcatechol}$  = 2.21 relative response factor of compound *i* compared to the standard, 17.82 10 – 19
- Butylcatechol (PTBC)
- $ECN_{2-}$  = 0.33 effective carbon number of the standard, 0.75 + 0.60\*MBF 1 – 2
- Methylbenzofuran
- (MBF)
- $ECN_{2-}$  = 0.59 effective carbon number of compound *i*, 0.29 + 0.23\*PPA 1.5 – 15
- Phenylpropion aldehyde (PBA) \*PPA
- Alpha-AMS  $ECN_{2-}$  = 4.61 molecular weight of the standard, and 6.51 + 0.23\*AMS 13 – 32
- MW ethylstyrene Oxide (AMS\*EO)
- $ECN_{2-}$  = 0.38 molecular weight of compound *i*, 0.55 + 2.03\*DMBA 0.1 – 1
- zyl Alcohol (DMBA)

2.2 *Other Document:*

OSHA Regulations, 29 CFR, paragraphs 1910.1000 and 1910.1200<sup>3</sup>

### 3. Terminology

3.1 See Terminology D 4790 for definition of terms used in this test method.

### 4. Summary of Test Method

~~4.1 A known amount of internal standard~~

~~4.1 AMS (alpha-methylstyrene) is added to a sample of AMS. The prepared sample is mixed and analyzed by a gas chromatograph (GC) equipped with a flame ionization detector (FID). The peak area A precisely repeatable volume of each impurity and the internal standard sample to be analyzed is measured and injected onto the amount of each impurity is calculated from the ratio of the gas chromatograph. The peak areas of the impurities are measured and converted to concentrations via an external standard versus the peak area of the impurity. methodology. Purity by GC (the AMS content) is calculated by subtracting the sum of the impurities from 100.00. Results are reported in weight percent.~~

### 5. Significance and Use

5.1 This test method is suitable for setting specifications on the materials referenced in 1.2 and for use as an internal quality control tool where AMS is produced or is used in a manufacturing process. It may also be used in development or research work involving AMS.

5.2 This test method is useful in determining the purity of AMS with normal impurities present. If extremely high boiling or unusual impurities are present in the AMS, this test method would not necessarily detect them and the purity calculation would be erroneous.

### 6. Apparatus

6.1 *Gas Chromatograph*—Any instrument having a flame ionization detector that can be operated at the conditions given in Table 2. The system should have sufficient sensitivity to obtain a minimum peak height response for 0.01% *n*-octane 10 mg/kg acetophenone of twice the height of the signal background noise.

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<sup>3</sup> Available from U.S. Government Printing Office Superintendent of ASTM Standards, Vol T4.02: Documents, 732 N. Capitol St., NW, Mail Stop: SDE, Washington, DC 20401.

**TABLE 2 Instrumental Parameters**

Detector	flame ionization
Injection Port	capillary splitter
Column A:	
Tubing	fused silica
Stationary phase	crosslinked methylsilicone
Film thickness, $\mu\text{m}$	1.0
Length, m	60
<del>Diameter, mm</del>	<del>0.25</del>
Diameter, mm	0.32
Temperatures:	
Injector, $^{\circ}\text{C}$	250
<del>Detector, <math>^{\circ}\text{C}</math></del>	<del>250</del>
Detector, $^{\circ}\text{C}$	300
<del>Oven, <math>^{\circ}\text{C}</math></del>	<del>125</del>
Oven, $^{\circ}\text{C}$	85 hold for 13 min Ramp 1 = 6 $^{\circ}\text{C}/\text{min}$ to 125 $^{\circ}\text{C}$ , hold for 2 min Ramp 2 = 30 $^{\circ}\text{C}/\text{min}$ to 250 $^{\circ}\text{C}$ , hold for 7.5 min
Carrier gas	helium
Flow rate, ml/min	2
Flow rate, ml/min	3
Split ratio	70:1
Split ratio	60:1
Sample size, $\mu\text{l}$	1.0
Sample size, $\div \mu\text{l}$	1.0

6.2 *Columns*—The choice of column is based on resolution requirements. Any column may be used that is capable of resolving all significant impurities from AMS and the internal standard, AMS. The column described in Table 1 2 has been used successfully. Unless the analyst can be sure of peak identity—(, for example by gas chromatography-mass spectrometry (GC-MS)), the use of the column in Table 1 2 is strongly recommended.

6.3 *Recorder*—Electronic integration is recommended.

6.4 *Injector*—The specimen must be precisely and repeatably injected into the gas chromatograph. An automatic sample injection device is highly recommended. Manual injection can be employed if the precision stated in Table 1 can be reliably and consistently satisfied.

## 7. Reagents and Materials

7.1 *Purity of Reagents*—Reagent grade chemical shall be used in all tests. Unless otherwise indicated, it is intended that all reagents shall conform to the specification of the Committee on Analytical Reagents of the American Chemical Society, where such specifications are available.<sup>4</sup> Other grades may be used, provided it is first ascertained that the reagent is of sufficiently high purity to permit its use without lessening the accuracy of the determination.

7.1.1 Normal octane is the recommended internal standard of choice. Other compounds may be found acceptable provided they are sufficiently resolved from any impurity and from the AMS peak.

7.2 *Carrier Gas*—Chromatographic grade helium is recommended.

7.3 *Compressed Air*—Chromatographic grade.

7.4 *Hydrogen*—High purity.

## 8. Hazards

8.1 Consult current OSHA regulation, suppliers' Material Safety Data Sheets, and local regulations for all materials used in this test method.

## 9. Sampling and Handling

9.1 Sample the material in accordance with Practice D 3437.

## 10. Preparation of Apparatus

10.1 Follow manufacturer's instructions for mounting and conditioning the column into the chromatograph and adjusting the instrument to the conditions described in Table 1 2 allowing sufficient time for the equipment to reach equilibrium. See Practice E 1510 for more information on column installation. See Practice E 355 for additional information on gas chromatography practices and terminology.

## 11. Procedure

### 11.1 ~~Into~~ Calibration

11.1 Prepare a 100-mL volumetric flask, add 100  $\mu$ L synthetic mixture of *n*-octane high purity AMS containing impurities at concentrations representative of those expected in the samples to 99.90 mL be analyzed in accordance with Practice D 4307. The weight of AMS. Mix well. Assuming each hydrocarbon impurity must be measured to the nearest 0.1 mg. Because the availability of stock AMS with a density purity higher than 99.80 % is problematic, the method of 0.704 standard additions may be required for *n*-octane impurities such as tert-butylbenzene and 0.910 *n*-propylbenzene, as well as for AMS, a number of the other impurities listed in ~~n-o~~ 1.2 that are commonly present. In addition, nearly all commercially available AMS is stabilized with 10 to 30 mg/kg of para-tertiary-butylcatechol, requiring a standard addition methodology for this component as well unless AMS can be 0.0774 weight % obtained safely and directly from the point of manufacture.

11.2 Inject the resulting solution from 11.1 into the gas chromatograph, an appropriate amount of sample as previously determined according to 6.1 collect and start process the analysis.

11.3 Obtain a chromatogram and peak integration report. data. A typical chromatogram is illustrated in Fig. 1 illustrates a typical analysis of AMS using Column A and based on the conditions listed in Table 2.

11.3 Determine the response factor for each impurity in the calibration mixture as follows:

$$Rf_i = C_i/A_i \quad (1)$$

<sup>4</sup>Available from

<sup>4</sup> Reagent Chemicals, American Chemical Society Specifications, American Chemical Society, Washington, DC. For suggestions on the Superintendent testing of Documents, U.S. Government Printing Office, Washington, DC 20402. reagents not listed by the American Chemical Society, see *Analar Standards for Laboratory Chemicals*, BDH Ltd., Poole, Dorset, U.K., and the *United States Pharmacopeia and National Formulary*, U.S. Pharmacopeial Convention, Inc. (USPC), Rockville, MD.

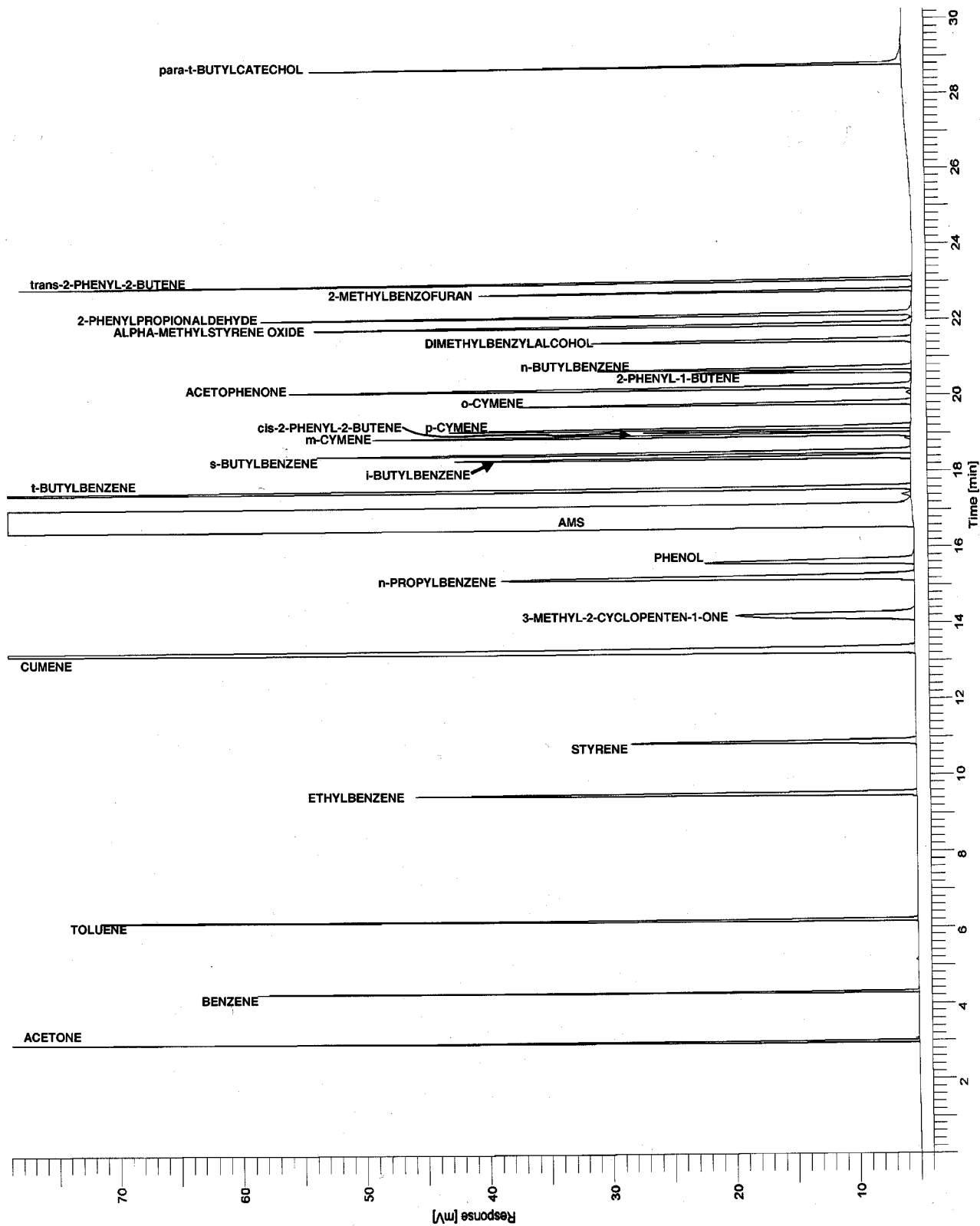


FIG. 1 Typical AMS Chromagram, AMS Method

where:

$Rfi$  = response factor for impurity  $i$ ,

$Ci$  = concentration of impurity  $i$  in the calibration mixture, and

$Ai$  = peak area of impurity  $i$ .

11.4 Initially analyze the calibration solution a minimum of three times and calculate an average  $Rfi$ . Subsequent calibrations may be a single analysis as long as the  $Rfis$  for all components of interest are within  $\pm 5\%$  of the initial validation  $Rfis$ . A “rolling” average as defined by most modern chromatographic software may also be used. The  $Rfi$  for tert-butylbenzene is used for the quantification of unknown impurities.

## 12. Procedure

12.1 Inject into the gas chromatograph an appropriate amount of sample as previously determined according to 6.1 and start the analysis.

12.2 Obtain a chromatogram and peak integration report. Fig. 1 illustrates a typical analysis of AMS using the column and conditions outlined in Table 2.

## 13. Calculations

12.1 Because some of

13.1 Of the impurities identified in AMS, only the butenylbenzene isomers are not available commercially, this test method is based on the use commercially. However, pure samples of theoretical response factors calculated with the use of effective carbon numbers (see Table 2):

12.2 For other impurities of known structure, the relative response factor these isomers can be calculated in a similar manner. For impurities of unknown structure, an average prepared, and testing has shown that all three isomers have the same response factor of 0.94 is suitable if the impurity is believed to consist of only carbon  $\pm 5\%$ , and hydrogen, and 1.2 that it is suitable if the impurity is believed equivalent to contain oxygen.

12.3 Calculate that for tert-butylbenzene  $\pm 5\%$ .<sup>5</sup>

13.2 Calculate the percent concentration of each impurity as follows:

$$C_i = \frac{(A_i)(RRF_i)(C_2)}{A_2} \quad (2)$$

$$C_i = (A_i)(Rfi)$$

$Rfi$

where:

$C_i$  = concentration of component  $i$ , weight percent, in mg/kg,

$A_i$  = peak area of component  $i$ , and

$RRFRfi$  = relative response factor for component  $i$ ;

$i$

$A_2$  = peak area of  $n$ -octane, and

$C_2$  = concentration of  $n$ -octane, weight percent.

12.3.43 Calculate the total concentration of all impurities in wt. % as follows:

$$C_t = \sum C_i$$

$$C_t = \sum C_i / 10000 \quad (3)$$

where:

$C_t$  = total concentration of all impurities.

12.3.54 Calculate the purity of AMS as follows:

$$\text{AMS, weight percent} = 100.00 - C_t \quad (4)$$

## 13. Report

13.1 Report the individual impurities to the nearest 0.01 %.

*Reagent Chemicals, American Chemical Society Specifications, American Chemical Society, Washington, DC. For suggestions on the testing of reagents not listed by the American Chemical Society, see *Analar Standards for Laboratory Chemicals*, BDH Ltd., Poole, Dorset, U.K., and the *United States Pharmacopeia and National Formulary*, U.S. Pharmacopeial Conventions, Inc. (USPC), Rockville, MD.*

<sup>5</sup> Supporting data are available from ASTM International Headquarters. Request RR:D16-1022.

~~13.2 Report the purity of AMS to the nearest 0.01 %.~~

#### **14. Report**

~~14.1 Report the individual impurities to the nearest 0.1 mg/kg.~~

~~14.2 Report the purity of AMS to the nearest 0.01 wt. %.~~

#### **15. Precision and Bias**

~~14.1 Precision—The following Bias<sup>5</sup>~~

~~15.1 The criteria in Table 1 should be used to judge the acceptability at the 95 % probability level of the results obtained by this test method. The criteria were derived from an interlaboratory study between six five laboratories (only one laboratory was used for repeatability data) on six samples run in triplicate by the same operator on the same day. The data results were obtained over one day derived using the same operators.~~

~~14.1.1 protocol set forth in Practice E 691.~~

~~15.2 Repeatability— Results in the same laboratory should not be considered suspect unless they differ by more than 0.03 %.~~

~~14.1.2 the amounts calculated from the appropriate equations in Table 1.~~

~~15.3 Reproducibility— Results submitted by two laboratories should not be considered suspect unless they differ by more than 0.09 %.~~

~~14.2 the amounts calculated from the appropriate equations in Table 1.~~

~~15.4 Bias— Since there is no accepted reference material suitable for determining the bias in this test method for measuring these impurities, bias has not been determined.~~

#### **156. Quality Guidelines**

~~16.1 Refer to Guide D 6809 for suggested QA/QC activities that can be used as a part of this method. It is recommended that the operator of this method select and perform relevant QA/QC activities like the ones in Guide D 6809 to help ensure the quality of data generated by this method.~~

#### **17. Keywords**

~~157.1 acetophenone; alpha methylstyrene; AMS; analysis by gas chromatography; benzene; butenylbenzene; butylbenzene; catechol; cumene; cymene; phenol; propylbenzene; toluene~~

### **SUMMARY OF CHANGES**

Committee D16 has identified the location of selected changes to this standard since the last issue (D 6144 - 97) that may impact the use of this standard.

- (1) Section 1 – Modified Scope and other sections to reflect external standard and broader scope of measured components
- (2) Section 2 – Practices D 4307 and E 691 were added as references.
- (3) Section 6 – Added 6.4 on injection technique.
- (4) Section 7 – Deleted 7.1.1 referencing n-octane as internal standard.
- (5) Section 11 – Modified extensively to reflect external standard methodology.
- (6) Section 13 – All reference to total carbon numbers removed, modified extensively to reflect external standard methodology.
- (7) Section 15 – Updated to reflect results of ILS and statistics generated by B.J. Lane following Practice E 691.
- (8) Section 16 – Converted to Quality Guidelines section.
- (9) Section 17 – Additional components added to keywords.

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