



Designation: D1091 – 00 (Reapproved 2005)

## Standard Test Methods for Phosphorus in Lubricating Oils and Additives<sup>1</sup>

This standard is issued under the fixed designation D1091; 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.

*This standard has been approved for use by agencies of the Department of Defense.*

### 1. Scope\*

1.1 These test methods cover the determination of phosphorus in unused lubricating oils and lubricating oil additives and their concentrates. The test methods are not restricted with respect to the type of phosphorus compounds that may be present—for example, trivalent or pentavalent phosphorus compounds, phosphines, phosphates, phosphonates, phosphorus sulfides, and so forth—since all are quantitatively converted to an aqueous solution of orthophosphate ion by oxidation of the sample during the course of analysis.

1.2 The values stated in SI units are to be regarded as the standard.

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.*

### 2. Referenced Documents

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

[D1193 Specification for Reagent Water](#)

[D4057 Practice for Manual Sampling of Petroleum and Petroleum Products](#)

[D4177 Practice for Automatic Sampling of Petroleum and Petroleum Products](#)

[D6299 Practice for Applying Statistical Quality Assurance and Control Charting Techniques to Evaluate Analytical Measurement System Performance](#)

### 3. Summary of Test Method

3.1 Organic material in the sample is removed and the phosphorus is converted to phosphate ion by oxidation with

<sup>1</sup> These test methods are under the jurisdiction of ASTM Committee D02 on Petroleum Products and Lubricants and are the responsibility of Subcommittee D02.03 on Elemental Analysis.

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<sup>2</sup> For referenced ASTM standards, visit the ASTM website, www.astm.org, or contact ASTM Customer Service at service@astm.org. For *Annual Book of ASTM Standards* volume information, refer to the standard's Document Summary page on the ASTM website.

sulfuric acid, nitric acid, and hydrogen peroxide. One of two procedures is then followed:

Photometric (Molybdivanado) Method  
Gravimetric Method

Sections  
**7-18**  
**7-11**  
and  
**19-25**

3.2 The photometric method is used where the phosphorus content is estimated to be under 2 %, and the gravimetric method is used for phosphorus contents of 2 % or over.

### 4. Significance and Use

4.1 Knowledge of the phosphorus content, and thus the phosphorus-containing additives, in a lubricating oil or additive can be used to predict performance characteristics. This test method is suitable for most applications requiring the determination of phosphorus.

### 5. Purity of Reagents

5.1 *Purity of Reagents*—Reagent grade chemicals shall be used in all tests. Unless otherwise indicated, it is intended that all reagents conform to the specifications of the Committee on Analytical Reagents of the American Chemical Society where such specifications are available.<sup>3</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.

5.2 *Purity of Water*—Unless otherwise indicated, references to water shall be understood to mean reagent water as defined by Type II or Type III of Specification [D1193](#).

### 6. Sampling

6.1 Obtain samples in accordance with the instructions in Practices [D4057](#) or [D4177](#).

6.2 Take care that the test specimen is thoroughly representative of the material to be tested and that the portion of the sample is thoroughly representative of the test unit.

<sup>3</sup> *Reagent Chemicals, American Chemical Society Specifications*, American Chemical Society, Washington, D.C. 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 Convention, Inc. (USPC), Rockville, MD.

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

**TABLE 1 Sample Size**

| Phosphorus Content, % | Approximate Weight of Sample, g | Precision of Weighing, plus or minus, g |
|-----------------------|---------------------------------|---|
|                       |                                 | Photometric (Molybdivanado) Method      |
| 0.002 to 0.2          | 2                               | 0.004                                   |
| 0.2 to 2              | 0.2                             | 0.0004                                  |
| Gravimetric Method    |                                 |   |
| 2 to 5                | 2                               | 0.004                                   |
| 5 to 10               | 1                               | 0.003                                   |
| 10 to 15              | 0.7                             | 0.002                                   |
| 15 to 25              | 0.4                             | 0.001                                   |

## OXIDATION OF THE SAMPLE

### 7. Scope

7.1 This test method covers a procedure for removal of organic material and subsequent conversion of phosphorus to phosphate ion in samples of unused lubricating oils, lubricating oil additives, and their concentrates.

### 8. Summary of Test Method

8.1 Organic material in the sample is destroyed and the phosphorus is converted to phosphate ion by oxidation with sulfuric acid, nitric acid, and hydrogen peroxide. The residual hydrogen peroxide is removed by diluting with water and evaporating several times to dense white fumes.

### 9. Apparatus

9.1 *Digestion Flasks*, Kjeldahl flasks, 300 mL, ground-glass stoppered.

9.2 *Digestion Rack*—A digestion rack constructed to hold one or more 300-mL Kjeldahl flasks at an angle of approximately 45° in such a fashion that direct heat is applied only to the bottom of the flask and such that the body and neck of the flask are insulated from the source of heat. Approximately three-fourths of the neck of the flask should be cooled by air at atmospheric temperature, preferably by directing an air stream against the neck of the flask. A Bunsen flame or high capacity electric heater are suitable heat sources.

### 10. Reagents

10.1 *Hydrogen Peroxide (30 %)*, concentrated hydrogen peroxide (H<sub>2</sub>O<sub>2</sub>) (**Warning**—Concentrated solutions are highly toxic and strong oxidants.) containing no more than 0.0002 % phosphorus.

10.2 *Nitric Acid (sp gr 1.42)*, concentrated nitric acid (HNO<sub>3</sub>).

10.3 *Sulfuric Acid (sp gr 1.84)*, concentrated sulfuric acid (H<sub>2</sub>SO<sub>4</sub>).

10.4 *White Oil*, phosphorus-free.

10.5 *Quality Control (QC) Samples*, preferably, portions of one or more liquid petroleum materials that are stable and representative of the samples of interest. These QC samples can be used to check the validity of the testing process, as described in Section 26.

### 11. Procedure

11.1 Weigh out a portion of the material to be analyzed, in accordance with Table 1, into a 300-mL Kjeldahl flask. Any convenient method of transferring the test specimen may be

used as long as care is taken to avoid getting the test specimen on the neck of the flask (see Note 1). Add H<sub>2</sub>SO<sub>4</sub> (3 mL for the photometric procedure, or 10 mL for the gravimetric procedure) and a 6-mm glass bead (see Note 2), and swirl the flask to mix the contents.

11.2 To obtain satisfactory accuracy with the small amounts of phosphorus involved, it is necessary to take extensive precautions in handling. The usual precautions of cleanliness, careful manipulation, and avoidance of contamination should be scrupulously observed; also, all glassware should be cleaned before use, with cleaning acid or by some procedure that does not involve use of commercial detergents. These compounds often contain alkali phosphates, which are strongly absorbed by glass surfaces and are not removed by ordinary rinsing. It is desirable to segregate a special stock of glassware for use only in the determination of phosphorus.

NOTE 1—The volume occupied by the glass bead (0.1 mL) can be ignored for ordinary work. Excessive bumping is encountered occasionally in the digestion of some organic phosphorus compounds. This bumping can be minimized by using a glass bead. Some difficulty can be experienced when using commercial boiling aids in obtaining a solution clear enough for photometric measurement of phosphorus (see Sections 12-18) even after centrifuging, due to the attrition of these boiling aids under the vigorous digestion procedure.

11.3 Make a blank determination following the same procedure and using the same amounts of all reagents and a similar size sample of phosphorus-free white oil. This blank is for use in the photometric method (see Sections 12-18).

11.4 Place the flask on the digestion rack under a hood and warm gently with a micro burner until the test specimen is charred, while cooling the neck of the flask, preferably by use of an air stream (see Note 2). Continue heating until dense white fumes appear (see Note 3). While boiling, continuously add 1 mL of HNO<sub>3</sub> dropwise (see Note 4) to oxidize the organic material. When the HNO<sub>3</sub> has boiled off and dense white fumes reappear, repeat the treatment with an additional 1 mL of HNO<sub>3</sub> (see Note 5). Continue the addition of HNO<sub>3</sub> in 1-mL increments until the digestion mixture is no darker than a straw color, indicating that almost all the organic matter has been oxidized.

NOTE 2—The amount of air used to cool the neck of the flask will at times have to be reduced or even shut off to allow vapors and fumes to leave the flask and to allow sample to come to dense white fumes. However, this should not be done until the test specimen is in a well-decomposed state; the air stream should be turned on again each time before the addition of the HNO<sub>3</sub> or H<sub>2</sub>O<sub>2</sub> (see 11.4).

NOTE 3—Excessive evaporation of H<sub>2</sub>SO<sub>4</sub> should be avoided to minimize any loss of phosphorus that may occur. Care should be exercised to avoid heating above the liquid level. Since there is some indication that with test specimens containing inorganic compounds (that is, barium or lead salts) there can be losses of phosphorus due to sintering or fusion of the phosphate and sulfate to the glass, it is well to examine the dried vessel after use to detect any opaque film of fused material.

NOTE 4—Unless the HNO<sub>3</sub> is added dropwise, it can force excessive amounts of vapor from the flask and lead to loss of phosphorus containing fumes.

NOTE 5—To minimize the loss of H<sub>2</sub>SO<sub>4</sub> in the digestion process, it is advisable not to prolong the dense white fumes stage between addition of HNO<sub>3</sub>.

11.5 Cool the flask slightly and add 10 drops (0.5 mL) of H<sub>2</sub>O<sub>2</sub>. Heat until dense white fumes appear, and while boiling,

cautiously add 1 mL of HNO<sub>3</sub> dropwise. When the HNO<sub>3</sub> has boiled off and dense white fumes reappear, repeat the treatment with H<sub>2</sub>O<sub>2</sub> and HNO<sub>3</sub> until the digestion mixture is colorless, at which time the organic material will be completely oxidized. Four treatments will usually suffice. The total amount of H<sub>2</sub>O<sub>2</sub> used should be noted, and the same amount used for each test specimen and the blank.

11.6 When oxidation is complete, allow the flask to cool, wash down the mouth and neck with a minimum amount of water (5 mL), and mix the contents. Return the flask to the digestion rack and continue heating to the appearance of dense white fumes. Repeat the process of the addition of water and heating to dense fumes several times. This will remove all traces of H<sub>2</sub>O<sub>2</sub>. (**Warning**—Use extreme care in fuming, in accordance with 11.5, to remove all traces of H<sub>2</sub>O<sub>2</sub> so that no color interference will be experienced when phosphorus is to be determined photometrically, as described in Sections 12-18.)

## PHOTOMETRIC (MOLYBDIVANADO) METHOD

### 12. Scope

12.1 This test method covers determination of total phosphorus in concentrations of less than two mass % (see Note 6), calculated on the basis of the original test specimen, in samples treated by the acid-oxidation procedure described in Sections 7-11.

NOTE 6—For phosphorus concentrations greater than or equal to two mass %, see Sections 19-25.

### 13. Summary of Test Method

13.1 After oxidation of organic material in the test specimen and quantitative conversion of the phosphorus to phosphate ion, the acidity of the digestion mixture is adjusted and the mixture diluted to suitable volume. Solutions of ammonium vanadate and ammonium molybdate are added in the order named. The addition of the molybdate solution to the acid vanadate-phosphate mixture results in the formation of a heteropoly acid, molybdivanadophosphoric acid, which is yellow in color. Although the exact composition of molybdivanadophosphoric acid is uncertain, solutions of this compound, when formed in accordance with carefully prescribed conditions, have been found to conform to the Beer-Lambert law for optical transmittance measurements made at 420 to 470 nm as a function of phosphorus content.

### 14. Apparatus

14.1 *Photoelectric Photometer*—A spectrophotometer capable of isolating a 5-nm spectral band at 430 and 460 nm is a suitable instrument for use in this determination. The instrument should be equipped with auxiliary facilities for handling 1, 2, and 5-cm cells, and a supply of these should be available. Other instruments such as photoelectric filter photometers may also be used.

NOTE 7—While not as desirable as photometers, visual color comparators can also be used, if necessary.

### 15. Reagents

15.1 *Ammonium Molybdate Solution*—Dissolve 50 g of ammonium molybdate (NH<sub>4</sub>)<sub>6</sub>Mo<sub>7</sub>O<sub>24</sub>·4H<sub>2</sub>O in warm water and dilute to 1 L. Filter before using.

15.2 *Ammonium Vanadate Solution*—Dissolve 2.5 g of ammonium vanadate (NH<sub>4</sub>VO<sub>3</sub>) in 500 mL of hot water, add 20 mL of concentrated nitric acid (HNO<sub>3</sub> relative density 1.42), and dilute to 1 L.

15.3 *Phosphate, Standard Solution (1 mL = 0.1 mg P)*—Dissolve 0.4393 g of potassium dihydrogen phosphate (KH<sub>2</sub>PO<sub>4</sub>) in water and dilute to 1 L. For best work, the salt should be twice recrystallized and vacuum-dried before use.

15.4 *Sulfuric Acid (relative density 1.84)*, concentrated sulfuric acid (H<sub>2</sub>SO<sub>4</sub>).

15.5 *QC Samples*, preferably, portions of one or more liquid petroleum materials that are stable and representative of the samples of interest. These QC samples can be used to check the validity of the testing process, as described in Section 26.

### 16. Calibration and Standardization

16.1 Introduce 0, 0.4, 0.8, 1.6, 2.4, 4.0, 4.8, 8.0, 16, 24, and 32 mL of standard phosphate solution into 100-mL ground-glass-stoppered volumetric flasks. Add sufficient H<sub>2</sub>SO<sub>4</sub> of any convenient concentration such that the final acid concentration after dilution to 100 mL will be 0.5 N. Dilute to 55 to 60 mL, and add 10 mL of ammonium vanadate solution and ammonium molybdate solution, in the order named, with adequate mixing between additions. Dilute to 100 mL, close with a ground-glass stopper, and mix thoroughly. Allow to stand at least 45 min but no longer than 60 min to develop the color.

16.2 Using the 1-cm cell and with the wave length set at 460 nm, adjust the photometer to read 100.0 % transmittance with the zero phosphate (reagent blank) standard. Although absorption cells are usually very closely matched, for best work it is recommended that two cells be used and that one be reserved for the blank and the other for the standard or sample solutions. Obtain transmittance measurements on solutions containing 0.4, 0.8, 1.6, 2.4, and 3.2 mg of phosphorus. These standards should give measurements falling between 90 and 20 % respectively. After making a measurement, return to the reagent blank cell. This should check the 100.0 % setting within 0.2 %. Repeat the reading of the standard and return to the blank. Obtain three readings in all of each standard solution. Using semilog graph paper, plot the average transmittance as a function of phosphorus content. The resultant curve should be a straight line.

16.3 In a similar manner, prepare calibration curves at 460 nm for the 2 and 5-cm cells, selecting concentrations from the series of standards that give readings between 20 and 90 %.

16.4 Finally, prepare a calibration curve for the 5-cm cell, using a wavelength setting of 430 nm. At this wavelength the molybdivanadophosphoric acid has a higher optical density, and the curve obtained will have a steeper slope. The advantages of having this calibration at the second wavelength are two-fold: (1) it provides increased sensitivity in the region of

low concentrations, and (2) it provides an independent confirmation of measurements made at 460 nm. Agreement between measurements at both wavelengths is a criterion of the absence of interference.

## 17. Procedure

17.1 To the cooled, decomposed sample in the Kjeldahl flask (see 11.5), add by visual observation sufficient  $\text{H}_2\text{SO}_4$  to bring the acidity to approximately one half the concentration present at the beginning of the acid-oxidation procedure (see 11.1). This step may not always be necessary (see Note 8). Cool the flask and contents and transfer to a 100-mL volumetric flask, using approximately 50 mL of water. (**Warning**—Extreme care should be exercised when adding water to  $\text{H}_2\text{SO}_4$ . It is advisable to add the water slowly, a small amount at a time, allowing it to run down the side of the flask, which is adequately cooled.)

NOTE 8—The acidity of the solution after acid oxidation is critical, since interference occurs from the appearance of an orange-yellow color, which forms in a neutral or too acid solution. The acidity for proper development of the desired color should be in the range from 0.4 *N* to 0.6 *N* in  $\text{H}_2\text{SO}_4$ . Adjustment of acidity can not be required when the losses of  $\text{H}_2\text{SO}_4$  have been kept to a minimum in the fuming steps of the acid-oxidation procedure; however, it may be necessary to further evaporate  $\text{H}_2\text{SO}_4$  in order to bring the acidity of the solution to approximately optimum normality.

17.2 Add 10 mL each of ammonium vanadate solution (see Note 9) and ammonium molybdate solution. It is important that these solutions be added in the order named, with adequate mixing between additions, to ensure the reproducible composition of the complex. Dilute to 100 mL, stopper with a ground-glass stopper, and mix thoroughly. Allow to stand at least 45 min but no longer than 60 min to develop the color. Maintain the temperature of this solution within 5°C of the temperature at which the calibration was performed.

NOTE 9—Remove the last trace of hydrogen peroxide since very little hydrogen peroxide is required to develop the maximum color of the vanadium-hydrogen peroxide complex. Any trace of  $\text{H}_2\text{O}_2$  will be evident by the reddish brown color obtained upon the addition of the vanadium reagent. When such is the case, the sample must be discarded and the acid-oxidation step will have to be repeated on a new test specimen.

17.3 When any insoluble matter is present, transfer a portion of the solution to a centrifuge tube, centrifuge at 1200 rpm for 5 min, and decant the clear supernatant liquid into the absorption cell. If desired, a portion of the solution may be drawn off by means of a filter stick. Avoid the use of filter paper as the colored complex may be adsorbed on it.

17.4 When the approximate phosphorus content is known, the path length of the absorption cell should be chosen to give a transmittance between 25 and 50 %. It is desirable to employ conditions such that readings fall within this range to reduce the error in the photometric measurement. If the phosphorus content is unknown, the analyst, with experience, will be able to select the best cell to use by visual observation. In the case of test specimens that prove to be too highly colored to be read directly, transfer an appropriate aliquot to another volumetric

flask and dilute with the reagent blank solution in order to maintain all the reagent concentrations at the proper level. Make all measurements at 460 nm, except for the extremely low concentrations (below 0.25 mg P/100 mL), which shall be made at 430 nm. In making the readings, adjust the galvanometer to 100.0 % with the reagent blank solution in the light path. Insert the sample in the light path, read the percentage transmittance to 0.1 %, and return to the reagent blank, which should check the original setting within 0.2 %. Readjust to 100.0 if necessary and repeat, obtaining at least three readings on the samples. These should agree within 0.2 %. Use the average of these readings to obtain the phosphorus content from the calibration curves.

17.5 *Overall Blank*—Although a reagent blank solution is used in preparing the calibration curves, an overall blank determination should be carried through on a sample of phosphorus-free white oil. No phosphorus should be detectable in such a blank.

## 18. Calculation

18.1 Calculate the percentage of phosphorus as follows:

$$\text{Phosphorus, mass \%} = ((P_s - P_b) \times D \times 100) / (1000 \times S) \quad (1)$$

where:

$P_s$  = milligrams of phosphorus in test specimen read from standard curve,

$P_b$  = milligrams of phosphorus in overall blank read from standard curve,

$D$  = dilution factor, if an aliquot is used (see 17.4), and

$S$  = mass of test specimen.

## GRAVIMETRIC METHOD<sup>2</sup>

## 19. Scope

19.1 This test method covers the determination of total phosphorus in concentrations of 2 mass % or more, (see Note 10), calculated on the basis of the original sample, in samples treated by the acid-oxidation procedure described in Sections 7-11.

NOTE 10—For phosphorus concentrations less than 2 mass %, see Sections 12-18.

## 20. Summary of Test Method

20.1 After oxidation of organic material in the test specimen and quantitative conversion of the phosphorus to phosphate ion, the phosphate ion is separated from interfering metals by precipitation as ammonium molybdiphosphate in nitric acid solution. After an ammoniacal solution of the phosphate ion is obtained, the phosphorus is precipitated as magnesium ammonium phosphate, ignited, and weighed as magnesium pyrophosphate.

## 21. Apparatus

21.1 *Electric Muffle Furnace*, capable of operating over a variable temperature range from 200 to 1100°C and of maintaining a temperature of  $1050 \pm 50^\circ\text{C}$ .